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September 12, 2016

Jeffrey Dyber, P.E. New York State Department of Environmental Conservation 625 Broadway Albany, New York, 12233-7015

Re: Pilot Test Work Plan for BCP # C224217

1815-1825 Ocean Avenue Brooklyn, New York Block: 7656 Lots: 55 & 58

Dear Mr. Dyber,

This Pilot Test Work Plan was prepared on behalf of Ocean Units LLC for the property known as the Former Tomat Service Station, located at 1815-1825 Ocean Avenue, Brooklyn, New York. The purpose of performing the pilot test is to obtain the field data needed to finalize the design of an Air Sparge / Soil Vapor Extraction (AS/SVE) system which is being proposed to remediated petroleum impacted soil and groundwater at the Site.

The results of sampling performed during the RI identified gasoline related VOCs in soil at the groundwater interface (approximately 20 to 25 feet below grade) above Restricted Residential Use SCOs in the vicinity of the two 4,000-gallon gasoline underground storage tanks located in front of the gasoline service station building on Lot 55. The contamination at the groundwater interface was found to extend to the southern area of the Site in the direction of groundwater flow.

Petroleum VOCs were detected within groundwater above NYSDEC Ambient Water Quality Standards (GQS) across the Site. The highest concentrations of gasoline-related VOCs in groundwater were detected within the groundwater samples collected closest to the tank pad and immediately down gradient of the two 4,000-gallon gasoline underground storage tanks (GW4, 15GW1 and 15GW2).

Conceptual Design AS /SVE System:

A conceptual layout of the system as shown in **Figure 1** includes nine sparging points and two SVE wells. This layout is based on an assumed radius of influence of 20 ft for the air sparging wells and 50 ft for the SVE wells which in our experience is typically achieved using standard design variables. These include constructing the sparge points using 2.5 feet of 0.020 slot 1-inch diameter PVC well-screen installed 10 feet below the static water table and connecting them to a blower capable of overcoming the breakout pressure plus 10 to 15 percent.

The SVE wells will be constructed of 2-inch diameter PVC with 10 feet of 0.020 inch slot screen set approximately 10 to 20 feet below grade; this will be connected to a 3-inch (or greater) diameter vent line to reduce friction loss. Given the variability in depth-to-groundwater measurements, the depth to groundwater will be locally verified before installing the SVE wells. The exact depth of the SVE wells will be approximately 1.5 feet above groundwater.

Air Sparge / Soil Vapor Extraction System Pilot Test:

The pilot test will consist of three parts: Soil Vapor Extraction-only, Air Sparging-only, and a combined AS/SVE test. For testing purposes, a single air sparging point and a single SVE well will be installed on site (**Figure 2**). One 2-inch diameter SVE well will be installed with GeoprobeTM equipment and constructed of 10 feet of 0.020 inch-PVC well screen; the exact depth of the well will be determined based on the measured depth-to-groundwater (estimated to be 20ft). The sparge point will be installed to a total depth of 38 feet below grade with GeoprobeTM direct push equipment and will be constructed of 1-inch diameter PVC casing with a 2.5-foot, 0.20-inch screened section. This will place the top of the well screen approximately 10 feet below the water table (estimated 35 feet). The screen will be gravel-packed using a Morie #1 filter media to 1 foot above the base of the casing, followed by one foot of bentonite pellets to prevent grout infiltration. The annular space will then be grouted with a bentonite-cement slurry to three feet below grade and backfilled with native material to grade. Six existing monitoring wells will be used as observation points.

SVE-Only Portion

The SVE portion of the test will be performed first and will consist of attaching a regenerative blower to the monitoring well and then recording negative pressure in the nearby observation wells to establish the effective radius of influence. For the purposes of this study, a reading of -0.1 inch H₂O will be regarded as the limit of influence.

Vacuum readings will be taken at the blower and wellhead with a diaphragm gauge, and at the observation points with a digital manometer. Contaminant concentration will be measured with a photoionization detector (PID) calibrated to provide a reading of total volatiles. VOCs will also be sampled via the T0-15 method; the collected data can determine the extent of effluent treatment by completing the discharge calculations as outlined in the DAR-1 policy document. Flow rates will be determined utilizing a vacuum/cfm chart supplied by the manufacturer. Previous to discharge into the atmosphere, an activated carbon system (1 55-gallon General Carbon Corp drum) will function as a filter for the untreated SVE effluent.

Three SVE-only tests will be run: one at full vacuum, one at 33% full vacuum and one at 66% full vacuum. The tests will be run from lowest applied vacuum (33%) to greatest (full). Each test will be run for approximately 15 minutes to allow for stabilization.

Sparge-Only Portion

During the sparge-only portion of the test, an air compressor capable of delivering 10 cfm at a maximum pressure of 10 psi will be attached to the sparge point. The compressor will be fitted with an oil-removing coalescing filter, pressure gauge, regulator and volumetric air flow meter.

The test will be run at two different pressure/flow levels, including pressures 10%, and 20%, over that required to displace the water column in the well (break-out pressure). At each range tested, water level and pressure readings will be recorded at the observation wells to determine the radius of influence of the sparge point.

Water level readings will be recorded prior to the start of the test and again at the test conclusion with an electronic tape. Additionally, dissolved oxygen (DO) levels will be recorded before and during the test to quantify changes during the process. A digital manometer will be used to monitor positive pressure levels at the observation points. A rise in water level of 0.02 feet or more, and/or a pressure reading of 0.02 inches of water were taken as the limit of influence.

SVE/Air Sparging Combined Portion

The final test will combine both SVE and Air Sparging to establish initial recovery rates and the overall pressure gradient. During this test, a vacuum will be applied to the SVE well with the sparge point running simultaneously at a wellhead pressure 10% above the calculated break-out pressure. The DO levels and VOC concentrations will be measured through the same methods mentioned in the AS-only and SVE-only test portions. Total volatile readings will be taken at the effluent stack for a period of thirty minutes with the PID. The effluent filter system (activated carbon drum) will be utilized in the combined test that was used in the SVE-only test.

Health and Safety Plan (HASP) & Community Air Monitoring Plan (CAMP):

The HASP and CAMP for the 1815-1825 Ocean Avenue Pilot Test Work Plan are included in the appendix.

The Remedial Action Work Plan (RAWP):

Once the pilot test is completed, the results will be included in the Site Remedial Action Work Plan (RAWP) or in a separate design document. The raw data will be included in an Excel chart. A graph will be generated based on the results of the SVE and AS tests, representing the pressure at each observation well as a function of its distance to the drawing well. In turn, these calculations can be used to determine the radius of influence, when also considering the soil composition and dissolved oxygen levels in the AS system. The total number of wells will be determined in the RAWP based on the ROI and room dimensions.

The equipment and treatment in the system will also be selected based on the pilot test results. The RAWP will present these design decisions and calculations. The blower will be selected based on the characteristic curve after determining the pressure and air flow in the needed in the system. DAR-1 calculations will be completed based on the VOC measurements collected via T0-15 methods; these calculations determine the necessity and extent of effluent treatment. The RAWP will include the T0-15 results and calculations. The effluent concentrations will be compared to the NYSDOH Air Quality standards as well. If activated carbon is considered an

appropriate treatment, the size of the system will be calculated based on the estimated concentration, airflow, and well diameters.

All final design parameters will be explained and outlined in the RAWP. Any choices that deviate from the original plan (included in the RIWP) will be justified.

Certification

I, <u>Ariel Czemerinski</u>, certify that I am currently a NYS registered professional engineer and that this Soil Vapor Intrusion Test 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)

Please feel free to give me a call with any questions.

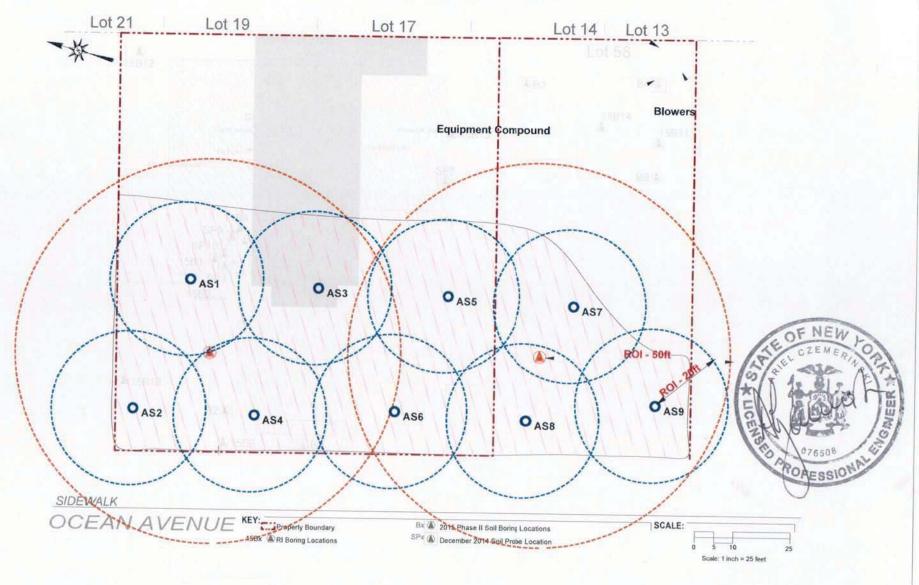
Regards,



Ariel Czemerinski

Figure 1: Conceptual AS/SVE Layout

GAC Vapor Phase Carbon Canisters





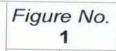
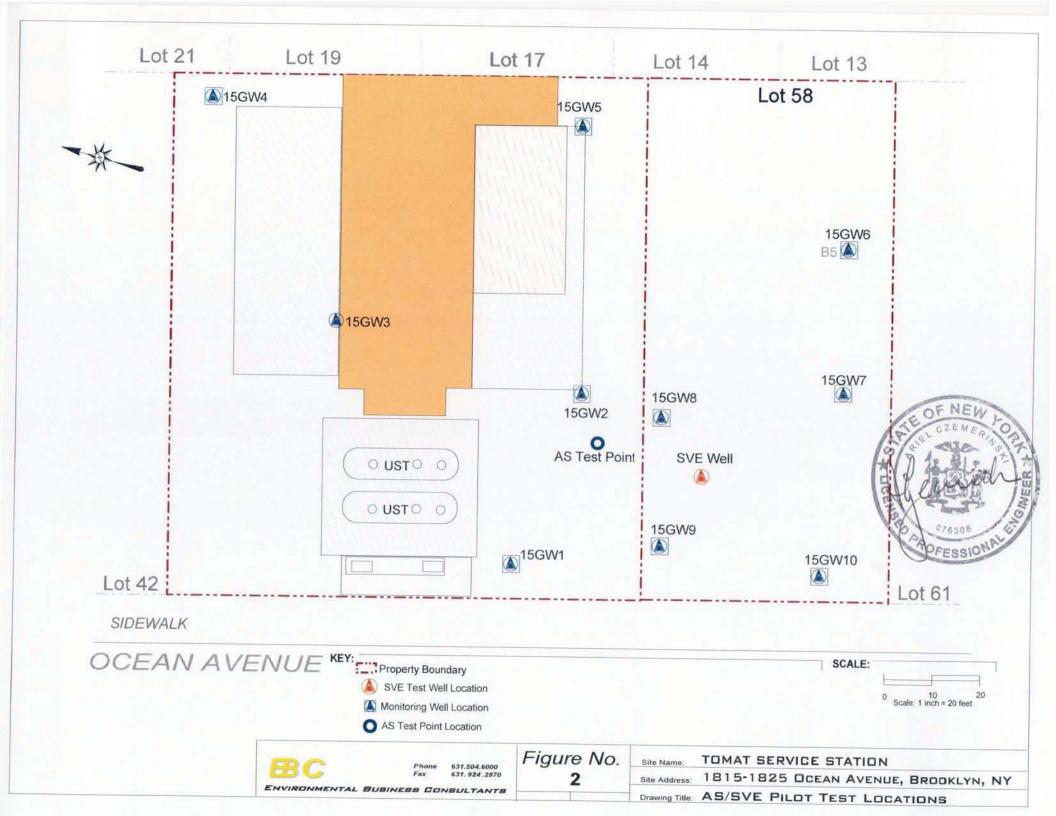


Figure 2: AS/SVE Pilot Test Locations



SVE/AS Pilot Test Work Plan: Construction Health and Safety Plan (CHASP)

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STATEMENT OF COMMITMENT

This Construction Health and Safety Plan (CHASP) has been prepared to ensure that workers are not exposed to risks from hazardous materials during the SVE/AS Pilot Test at the Tomat Service Station Site, 1815-1825 Ocean Avenue, Brooklyn, NY

This CHASP, which applies to persons present at the site actually or potentially exposed to hazardous materials, describes emergency response procedures for actual and potential chemical hazards. This CHASP is also intended to inform and guide personnel entering the work area or exclusion zone. Persons are to acknowledge that they understand the potential hazards and the contents of this Health and Safety policy by signing off on receipt of their individual copy of the document. Contractors and suppliers are retained as independent contractors and are responsible for ensuring the health and safety of their own employees.

1.0 INTRODUCTION AND SITE ENTRY REQUIREMENTS

This document describes the health and safety guidelines developed by Environmental Business Consultants (EBC) for the planned SVE/AS Pilot Test at the Tomat Service Station Site, 1815-1825 Ocean Avenue, Brooklyn, New York to protect on-site personnel, visitors, and the public from physical harm and exposure to hazardous materials or wastes during remedial activities. In accordance with the Occupational Safety and Health Administration (OSHA) 29 CFR Part 1910.120 Hazardous Waste Operations and Emergency Response Final rule, this CHASP, including the attachments, addresses safety and health hazards related to excavation, tank removal, loading and other soil disturbance activities and is based on the best information available. The CHASP may be revised by EBC at the request of Ocean Unites LLC and/or a regulatory agency upon receipt of new information regarding site conditions. Changes will be documented by written amendments signed by EBC's project manager, site safety officer and/or the EBC health and safety consultant.

Work performed under the remedial action will not involve confined space entry since excavation is minimal and limited to Geoprobe drilling.

1.1 **Training Requirements**

Personnel entering the exclusion zone or decontamination zone are required to be certified in health and safety practices for hazardous waste site operations as specified in the Federal OSHA Regulations CFR 1910.120e (revised 3/6/90).

Paragraph (e - 3) of the above referenced regulations requires that all on-site management personnel directly responsible for or who supervise employees engaged in hazardous waste operations, must initially receive 8 hours of supervisor training related to managing hazardous waste work.

Paragraph (e - 8) of the above referenced regulations requires that workers and supervisors receive 8 hours of refresher training annually on the items specified in Paragraph (e-1) and/or (e-3).

Additionally all on-site personnel must receive adequate site-specific training in the form of an on-site Health and Safety briefing prior to participating in field work with emphasis on the following:

- Protection of the adjacent community from hazardous vapors and / or dust which may be released during intrusive activities.
- Identification of chemicals known or suspected to be present on-site and the health effects and hazards of those substances.
- The need for vigilance in personnel protection, and the importance of attention to proper use, fit and care of personnel protective equipment.
- Decontamination procedures.
- Site control including work zones, access and security.
- Hazards and protection against heat or cold.
- The proper observance of daily health and safety practices, such as entry and exit of work



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zones and site. Proper hygiene during lunch, break, etc.

Emergency procedures to be followed in case of fire, explosion and sudden release of hazardous gases.

Health and Safety meetings will be conducted on a daily basis and will cover protective clothing and other equipment to be used that day, potential and chemical and physical hazards, emergency procedures, and conditions and activities from the previous day.

1.2 **Medical Monitoring Requirements**

Field personnel and visitors entering the exclusion zone or decontamination zone must have completed appropriate medical monitoring required under OSHA 29 CFR 1910.120(f) if respirators or other breathing related PPE is needed. Medical monitoring enables a physician to monitor each employee's health, physical condition, and his fitness to wear respiratory protective equipment and carry out on-site tasks.

1.3 Site Safety Plan Acceptance, Acknowledgment and Amendments

The project superintendent and the site safety officer are responsible for informing personnel (EBC employees and/or owner or owners representatives) entering the work area of the contents of this plan and ensuring that each person signs the safety plan acknowledging the on-site hazards and procedures required to minimize exposure to adverse effects of these hazards. A copy of the Acknowledgement Form is included in **Appendix A**.

Site conditions may warrant an amendment to the CHASP. Amendments to the CHASP are acknowledged by completing forms included in **Appendix B**.

1.4 **Key Personnel - Roles and Responsibilities**

Personnel responsible for implementing this Health and Safety Plan are:

Name	Title	Address	Contact Numbers
Mrs. Chawinie Reilly	EBC- Project Manager	1808 Middle Country Rd Ridge, NY 11961	(631) 504-6000
Mr. Kevin Waters	Health and Safety Officer	1808 Middle Country Rd Ridge, NY 11961	(631) 504-6000

The project manager is responsible for overall project administration and, with guidance from the site safety officer, for supervising the implementation of this CHASP. The site safety officer will conduct daily (tail gate or tool box) safety meetings at the project site and oversee daily safety issues. Each subcontractor and supplier (defined as an OSHA employer) is also responsible for the health and safety of its employees. If there is any dispute about health and safety or project activities, on-site personnel will attempt to resolve the issue. If the issue cannot be resolved at the site, then the project manager will be consulted.

The site safety officer is also responsible for coordinating health and safety activities related to hazardous material exposure on-site. The site safety officer is responsible for the following:

Ridge, NY 11961

- 1. Educating personnel about information in this CHASP and other safety requirements to be observed during site operations, including, but not limited to, decontamination procedures, designation of work zones and levels of protection, air monitoring, fit testing, and emergency procedures dealing with fire and first aid.
- 2. Coordinating site safety decisions with the project manager.
- 3. Designating exclusion, decontamination and support zones on a daily basis.
- 4. Monitoring the condition and status of known on-site hazards and maintaining and implementing the air quality monitoring program specified in this CHASP.
- 5. Maintaining the work zone entry/exit log and site entry/exit log.
- 6. Maintaining records of safety problems, corrective measures and documentation of chemical exposures or physical injuries (the site safety officer will document these conditions in a bound notebook and maintain a copy of the notebook on-site).

The person who observes safety concerns and potential hazards that have not been addressed in the daily safety meetings should immediately report their observations/concerns to the site safety officer or appropriate key personnel.

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2.0 SITE BACKGROUND AND SCOPE OF WORK

The street address for the Site is 1815-1825 Ocean Avenue, Brooklyn, NY. The Site is located in the Midwood section of Kings County and is comprised of two tax parcels totaling 16,555 square feet (0.38). The Site has approximately 150.5 ft of frontage along Ocean Avenue and is approximately 110 ft deep. Currently the property is developed with a one-story gasoline service station building (Lot 55) and a parking lot (Lot 58). The building has a footprint approximately 2,190 s.f. which, according to the NYC Department of Buildings, was constructed in 1931. These building are being demolished as part of the redevelopment of the Site.

The Site has a long history of use as a gas station since the 1930's.

2.1 **Previous Investigations**

2.1.1 December 30, 2013 –Subsurface Assessment Report (Hydrotech Environmental Corp) Hydro Tech Environmental, Corp. has performed a Subsurface Assessment at the property located at 1815 Ocean Avenue, Brooklyn, New York. This assessment was conducted on behalf of Tomat Service Center Inc. based upon their request to investigate the overall soil and groundwater quality.

The assessment consisted of the performance of the installation and sampling of a series of soil probes, groundwater probes, and monitoring wells. A Hydro Tech geologist screened all soil samples in the field for organic vapors utilizing a Photoionization Detector. Select soil, groundwater, and monitoring well samples were analyzed at a State-certified laboratory for volatile organic compounds (VOCs) and semi-volatile organic compounds (SVOCs). A select soil and groundwater sample was additionally analyzed for diesel range organic compounds and gasoline range organic compounds.

VOCs were detected in soil samples at the groundwater interface and in the groundwater in the vicinity of a fuel oil UST located to the north of the building at concentrations exceeding their respective regulatory standards. The petroleum compounds identified in the soil and groundwater samples was been classified as #2 fuel oil, as per the fingerprint analysis, obtained from the 23-25 foot soil sample from SP-9 and the groundwater sample from MW-1.

Dissolved VOCs (MW2 and MW3) were detected in groundwater beneath the southern portion of the Site, to the south of gasoline tanks and pump islands, at high concentrations exceeding their regulatory standards. Total VOCs in MW2 were noted to be 4,006.6 ug/L and at 6,471.7 ug/L in MW3. These findings indicated that groundwater had been impacted. Spill #1408472 is associated with the Site.

In addition, a ground penetrating radar survey (GPR) was conducted to clear sampling locations and discover any anomalies. One anomaly was identified on the north side of the interior of the site and is associated with a UST

2.1.2 Phase II Investigation Report – 1815 Ocean Avenue, Brooklyn NY. Environmental Business Consultants (EBC) dated February 25, 2015

The field work portion of the investigation was performed on February 7, 2015. The work



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consisted of the installation of four soil borings, two permanent monitoring wells, two temporary monitoring wells, and the collection and analysis of related samples. Four soil boring locations (B1 through B4) were selected. All of the borings were advanced with Geoprobe™ direct push equipment to a depth of 25 ft. Soil was characterized as a brown slit and sand with some historic fill material mixed in from surface grade to generally 1 foot below grade with the deepest locations at approximately 6-8 feet below grade followed by a brown silt and brown coarse fine sand to the termination depth. Groundwater was encountered and is expected at approximately 17 to 22 ft below grade.

PID readings of 130 ppm and petroleum odors were noted in B1 within the 21-25 foot interval. Petroleum odors were noted in the 21-25 foot interval of B2. PID readings of 250 ppm and petroleum odors were noted in B3 within the 21-25 foot interval. PID readings of 200 ppm were noted in B4 within the 21-25 foot interval. Soil samples were collected from the following intervals; 0-2 feet (B2 and B3) and 22-24 feet (B1-B4).

EBC collected four groundwater samples from each of the characterization borings (B1, B2, B3, B4). Permanent monitoring wells were installed for locations GW2 and GW4 by advancing the borehole to the water table (approx. 17 to 22 ft bgs) and installing a one-inch diameter PVC well 5-feet below the water table interface. Groundwater samples were collected in pre-cleaned, laboratory supplied glassware, stored in a cooler with ice and submitted to Phoenix Laboratories for analysis of VOCs by EPA Method 8260.

Deep soil and groundwater samples were analyzed for volatile organic compounds (VOCs) by USEPA method 8260. Shallow soil samples were analyzed for total lead.

Soil sample results were compared to the Unrestricted Use and Restricted Residential Use Soil Cleanup Objectives (SCOs) as presented in 6 NYCRR Part 375 Subparts 375-1 to 375-4 & 375-6. The following VOCs; 1,2,4-trimethylbenzene (maximum of 31,000 µg/Kg), 1,3,5trimethylbenzene (maximum of 9,900 µg/Kg), m&p-Xylenes (maximum of 2,300 µg/Kg), methylene chloride (maximum of 280 µg/Kg), naphthalene (at 14,000 µg/Kg) and o-xylene (maximum of 570 µg/Kg) were detected above Unrestricted Use SCOs in samples B1 and B2. VOCs including 2-isopropyltoluene (230 µg/Kg), acetone (maximum 34 µg/Kg), ethylbenzene (maximum 480 μg/Kg), n-butylbenzene (maximum 2,700 μg/Kg), n-propylbenzene (maximum 2,100 µg/Kg), p-isopropyltoluene (maximum 600 µg/Kg), sec-butylbenzene (maximum 860 μg/Kg) were detected at trace amounts in all soil samples. Lead was detected above Unrestricted Use SCOs in B2 and B3 shallow soil samples; at a maximum of 366 mg/kg.

Groundwater results were compared to the New York State Ambient Water Quality Standards and Guidance Values (6 NYCRR Part 703) as presented in the Technical & Operational Guidance Series (TOGS) 1.1.1 (1998). Several VOCs including 1,2,4-trimethylbenzene, 1,3,5trimethylbenzene, ethylbenzene, isopropylbenzene, m&p-Xylenes, naphthalene, n-butylbenzene, n-propylbenzene, o-xylene, p-isopropyltoluene and sec-butylbenzene were reported in all of the four monitoring wells above groundwater standards. 2-isopropyltoluene was reported in one groundwater well (GW4) above groundwater standards. Acetone was reported in two groundwater well (GW2 and GW3) above groundwater standards. Benzene was reported in one groundwater well (GW1) above groundwater standards. Toluene was reported in two groundwater well (GW1 and GW4) above groundwater standards.

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Soil and groundwater samples collected in the vicinity of the USTs indicate gasoline contamination. Several gasoline related VOCs were detected above Unrestricted Use SCOs in soil samples B1 and B2. Lead was detected above Unrestricted Use SCOs in B2 and B3 shallow soil samples. Gasoline related VOCs were noted above groundwater standards in all groundwater samples. These results indicate that further remedial action will be required.

2.1.3 Phase II Data Summary for 1825 Ocean Avenue, Brooklyn, NY. Environmental Business Consultants (EBC) dated April 29, 2015

The field work portion of the investigation was performed on April 22, 2015. The work consisted of the installation of five soil borings, three permanent monitoring wells, and the collection and analysis of related samples. Five soil boring locations (B1 through B5) were selected. Borings B1 and B2 were advanced with GeoprobeTM direct push equipment to a depth of 25 ft. Borings B3, B4 was advanced with GeoprobeTM direct push equipment to a depth of 15 ft. B5 was advanced with a GeoprobeTM to a depth of 20-23 feet. Soil was characterized as a brown medium fine sand with some historic fill material mixed in from surface grade to approximately 6-8 feet below grade followed by a brown sand to the termination depth. Groundwater was encountered and is expected at approximately 23 ft below grade.

Petroleum odors were noted in B1 and B2 within the 20-25 foot interval. Soil samples were collected from the following intervals; 0-2 feet (B1, B2 and B3) and 23-25 feet (B1 and B2).

EBC collected three groundwater samples from each of the characterization borings (B1, B2 and B5). Permanent monitoring wells were installed for locations B1, B2 and B5 by advancing the borehole to the water table (approx. 20 to 23 ft bgs) and installing a one-inch diameter PVC well 5-feet below the water table interface. Groundwater samples were collected in pre-cleaned, laboratory supplied glassware, stored in a cooler with ice and submitted to Phoenix Laboratories for analysis of VOCs by EPA Method 8260.

Deep soil and groundwater samples were analyzed for volatile organic compounds (VOCs) by USEPA method 8260. Shallow soil samples were analyzed for SVOCs (CP51), PCBs and TAL Metals. Deep soil samples were analyzed for (VOCs) by USEPA method 8260 and SVOCs (CP51).

Soil sample results were compared to the Unrestricted Use and Restricted Residential Use Soil Cleanup Objectives (SCOs) as presented in NYSDEC CP51 Soil Cleanup Guidance (10/21/10). The following VOCs; ethylbenzene (maximum of 22,000 µg/Kg), m&p-Xylenes (maximum of 45,000 μg/Kg), naphthalene (at 27,000 μg/Kg), n-Butylbenzene (at 17,000 μg/Kg) and n-Propylbenzene (at 35,000 µg/Kg) were detected above Unrestricted Use SCOs in samples B1 and B2. The following VOCs; 1,3,5-trimethylbenzene (maximum of 70,900 µg/Kg) and 1,2,4trimethylbenzene (maximum of 230,000 µg/Kg) were above RRSCOs in sample B2. No SVOCs and PCBs were detected above UUSCOs. The following metals; copper (at 56.1 mg/Kg), mercury (at 0.32 mg/Kg) and zinc (at 193 mg/Kg) were detected above UUSCOs in sample B3. Lead (at 1,860 mg/Kg) was detected above RRSCOs in sample B3.

Groundwater results were compared to the New York State Ambient Water Quality Standards and Guidance Values (6 NYCRR Part 703) as presented in the Technical & Operational

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Guidance Series (TOGS) 1.1.1 (1998). Several VOCs including 1,2,4-trimethylbenzene, 1,3,5trimethylbenzene, 2-Isopropyltoluene, benzene, ethylbenzene, isopropylbenzene, m&p-Xylenes, naphthalene, n-butylbenzene, n-propylbenzene, o-xylene, p-isopropyltoluene, sec-butylbenzene and toluene were reported in all of the three monitoring wells above groundwater standards. Soil and groundwater samples collected indicate gasoline contamination. Several gasoline related VOCs were detected above Unrestricted Use and Residential Restricted SCOs in soil samples B1 and B2. Lead was detected above Residential Restricted SCOs in the B3 shallow soil sample. Gasoline related VOCs were noted above groundwater standards in all groundwater samples. These results indicate that further remedial action will be required. A spill was called in for this site and assigned a SPILL # 1501018

2.1.4 Phase I Screening for 1815-1825 Ocean Avenue, Brooklyn, NY. Environmental Business Consultants (EBC) dated April 30, 2015.

Prior to the construction of the existing improvements (circa 1950), the Site was occupied by a gas station (lot 55) and 2-story dwelling (lot 58) in the 1930's. From 1895 to 1906, the site was vacant land (lot 55) and a 2-story dwelling (lot 58).

In the 1930 Sanborn map four gasoline tanks are depicted on the west side of the Site. In the 1950-2007 Sanborn maps, five gasoline tanks are depicted on the west side of the Site.

2.2 **Redevelopment Plans**

The redevelopment project consists of the demolition of the existing gas station and auto repair facility and the construction of a new 8-story residential building which will cover 65 percent of the Site. The Site will be developed with a full cellar. The cellar will occupy the entire footprint of the building down to about 10-12 feet below grade. The cellar will be utilized for bike storage, refuse area, elevator mechanical room, sprinkler room, water meter room, electrical meter room, plumbing room, recreation rooms and a gas meter room. The slab on grade area (44.3 x 110 foot area) on the north side of the Site will be excavated to a depth of 2 feet. The elevator pit will be excavated to approximately 15 feet below grade. The first floor of the Site will be developed with residential units, recreation space, and 45 parking spaces. The upper floors are developed with residential units. The maximum height of the building will be proposed at eighty feet tall with additional height provided for accessory spaces as permitted. The front yard will match the adjacent buildings on the street with a minimum of forty feet in the rear yard. The building will be proposed up to the side lot property lines. The building will be proposed with a maximum of ninety-three units with a mix of one, two and three bedroom units.

2.3 Description of SVE/Air Sparging Pilot Test Work Plan

This SVE/AS Pilot Test Work Plan addresses how to set-up the pilot test and the utility of the results. The pilot test will consist of three parts: Soil Vapor Extraction-only, Air Sparging-only, and a combined AS/SVE test. The results of each portion will be used to determine the design parameters of the full-scale system; the Site RAWP will include the results and final design decisions.

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3.0 HAZARD ASSESSMENT

This section identifies the hazards associated with the proposed scope of work, general physical hazards that can be expected at most sites; and presents a summary of documented or potential chemical hazards at the site. Every effort must be made to reduce or eliminate these hazards. Those that cannot be eliminated must be guarded against using engineering controls and/or personal protective equipment.

3.1 **Physical Hazards**

3.1.1 Tripping Hazards

An area of risk associated with on-site activities are presented by uneven ground, concrete, curbstones or equipment which may be present at the site thereby creating a potential tripping hazard. During intrusive work, care should be taken to mark or remove any obstacles within the exclusion zone.

3.1.2 Climbing Hazards

During site activities, workers may have to work on excavating equipment by climbing. The excavating contractor will conform with any applicable NIOSH and OSHA requirements or climbing activities.

3.1.3 Cuts and Lacerations

Field activities that involve excavating activities usually involve contact with various types of machinery. A first aid kit approved by the American Red Cross will be available during all intrusive activities.

3.1.4 Lifting Hazards

Improper lifting by workers is one of the leading causes of industrial injuries. Field workers in the excavation program may be required to lift heavy objects. Therefore, all members of the field crew should be trained in the proper methods of lifting heavy objects. All workers should be cautioned against lifting objects too heavy for one person.

3.1.5 Utility Hazards

Before conducting any excavation, the excavation contractor will be responsible for locating and verifying all existing utilities at each excavation.

3.1.6 Traffic Hazards

All traffic, vehicular and pedestrian, shall be maintained and protected at all times consistent with local, state and federal agency regulations regarding such traffic and in accordance with NYCDOT guidelines. The excavation contractor shall carry on his operations without undue interference or delays to traffic. The excavation contractor shall furnish all labor, materials, guards, barricades, signs, lights, and anything else necessary to maintain traffic and to protect his work and the public, during operations.

3.2 **Work in Extreme Temperatures**

Work under extremely hot or cold weather conditions requires special protocols to minimize the chance that employees will be affected by heat or cold stress.



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3.2.1 Heat Stress

The combination of high ambient temperature, high humidity, physical exertion, and personal protective apparel, which limits the dissipation of body heat and moisture, can cause heat stress.

The following prevention, recognition and treatment strategies will be implemented to protect personnel from heat stress. Personnel will be trained to recognize the symptoms of heat stress and to apply the appropriate treatment.

1. Prevention

- a. Provide plenty of fluids. Available in the support zone will be a 50% solution of fruit punch and water or plain water.
- b. Work in Pairs. Individuals should avoid undertaking any activity alone.
- c. Provide cooling devices. A spray hose and a source of water will be provided to reduce body temperature, cool protective clothing and/or act as a quick-drench shower in case of an exposure incident.
- d. Adjustment of the work schedule. As is practical, the most labor-intensive tasks should be carried out during the coolest part of the day.

2. Recognition and Treatment

Heat Rash (or prickly heat):

Continuous exposure to hot and humid air, aggravated by chafing Cause:

clothing.

Eruption of red pimples around sweat ducts accompanied by Symptoms:

intense itching and tingling.

Treatment: Remove source or irritation and cool skin with water or wet cloths.

b. Heat Cramps (or heat prostration)

Cause: Profuse perspiration accompanied by inadequate replenishment of

body water and electrolytes.

Muscular weakness, staggering gait, nausea, dizziness, shallow Symptoms:

breathing, pale and clammy skin, approximately normal body

temperature.

Treatment: Perform the following while making arrangement for transport to a

> medical facility. Remove the worker to a contamination reduction zone. Remove protective clothing. Lie worker down on back in a cool place and raise feet 6 to 12 inches. Keep warm, but loosen all clothing. If conscious, provide sips of salt-water solution, using one teaspoon of salt in 12 ounces of water. Transport to a medical

facility.

c. Heat Stroke

Cause: Same as heat exhaustion. This is also an extremely serious

condition.

Dry hot skin, dry mouth, dizziness, nausea, headache, rapid pulse. Symptoms:

Treatment: Cool worker immediately by immersing or spraying with cool

water or sponge bare skin after removing protective clothing.

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Transport to hospital.

3.2.2 Cold Exposure

Exposure to cold weather, wet conditions and extreme wind-chill factors may result in excessive loss of body heat (hypothermia) and /or frostbite. To guard against cold exposure and to prevent cold injuries, appropriate warm clothing should be worn, warm shelter must be readily available, rest periods should be adjusted as needed, and the physical conditions of on-site field personnel should be closely monitored. Personnel and supervisors working on-site will be made aware of the signs and symptoms of frost bite and hypothermia such as:

- Shivering;
- reduced blood pressure;
- reduced coordination;
- drowsiness;
- impaired judgment;
- fatigue;
- pupils dilated but reactive to light; and,
- numbing of the toes and fingers.

3.3 Chemical Hazards

The RI Investigation identified chlorinated and petroleum volatile organic compounds (VOCs) in soil, groundwater and soil vapor and free phase fuel oil and semi-volatile compounds in soil at the Site. In addition "Urban fill" materials which contain elevated levels of SVOCs and metals as well as other compounds were identified throughout the Site. These "contaminants" are not related to a chemical release occurring on the site, but are inherent in the reworked fill material in the area which contains ash and bits of tar and asphalt. Considering the previous sampling results and the past and present use of the site, the following compounds are considered for the site as potential contaminants: volatile organic compounds (VOCs), semi-volatile organic compounds (SVOCs), pesticides, polychlorinated biphenyl's (PCBs), and heavy metals such as arsenic, chromium, lead and mercury.

Based on the findings of the Remedial Investigation and the inherent properties of urban fill, the following compounds are considered for the site as potential contaminants: volatile organic compounds (VOCs), semi-volatile organic compounds (SVOCs), pesticides, and heavy metals.

Volatile organic compounds reported to be present in soil include the following:

1,2,4-trimethylbenzene	1,3,5-trimethylbenzene		ethylbenzene	isopropylbenzene
n-Butylbenzene	Toluene	napthalene		
n-propylbenzene	m&p-xylenes	o-xylenes		

Semi-Volatile organic compounds expected to be in the fill materials include the following:

Napthalene

Metals expected to be present in fill materials include the following:

Barium C	Chromium	Copper	Lead	Mercury	Nickel	Zinc
----------	----------	--------	------	---------	--------	------

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The primary routes of exposure to these contaminants are inhalation, ingestion and absorption.

Appendix C includes information sheets for suspected chemicals that may be encountered at the site. Also included under the appendix are procedures for handling and storing the chemical oxidant. These procedures will be followed to protect workers and the public.

3.3.1 Respirable Dust

Dust may be generated from vehicular traffic and/or excavation activities. If visible observation detects elevated levels of dust, a program of wetting will be employed by the site safety officer. If elevated dust levels persist, the site safety office will employ dust monitoring using a particulate monitor (Miniram or equivalent). If monitoring detects concentrations greater than $150 \, \mu \text{g/m}3$ over daily background, the site safety officer will take corrective actions as defined herein, including the use of water for dust suppression and if this is not effective, requiring workers to wear APRs with high efficiency particulate air (HEPA) cartridges.

Absorption pathways for dust and direct contact with soils or groundwater will be mitigated with the implementation of latex gloves, hand washing and decontamination exercises when necessary.

3.3.2 Dust Control and Monitoring During Earthwork

Dust generated during excavation activities or other earthwork may contain contaminants identified in soils at the site. Dust will be controlled by wetting the working surface with water. Calcium chloride may be used if the problem cannot be controlled with water. Air monitoring and dust control techniques are specified in a site specific Dust Control Plan (if applicable). Site workers will not be required to wear APR's unless dust concentrations are consistently over 150 $\mu g/m^3$ over site-specific background in the breathing zone as measured by a dust monitor unless the site safety officer directs workers to wear APRs. The site safety officer will use visible dust as an indicator to implement the dust control plan.

3.3.3 Organic Vapors

Elevated levels of chlorinated VOCs were detected in soil, soil gas and groundwater samples collected during previous investigations at the site. Therefore, excavation activities may cause the release of organic vapors to the atmosphere. The site safety officer will periodically monitor organic vapors with a Photoionization Detector (PID) during excavation activities to determine whether organic vapor concentrations exceed action levels shown in Section 5 and/or the Community Air Monitoring Plan.



4.0 PERSONAL PROTECTIVE EQUIPMENT

Personal protective equipment (PPE) shall be selected in accordance with the site air monitoring program, OSHA 29 CFR 1910.120(c), (g), and 1910.132. Protective equipment shall be NIOSH approved and respiratory protection shall conform to OSHA 29 CFR Part 1910.133 and 1910.134 specifications; head protection shall conform to 1910.135; eye and face protection shall conform to 1910.133; and foot protection shall conform to 1910.136. The only true difference among the levels of protection from D thru B is the addition of the type of respiratory protection. It is anticipated that work will be performed in Level D PPE.

4.1 Level D

Level D PPE shall be donned when the atmosphere contains no known hazards and work functions preclude splashes, immersion, or the potential for inhalation of, or contact with, hazardous concentrations of harmful chemicals. Level D PPE consists of:

- standard work uniform, coveralls, or tyvek, as needed;
- steel toe and steel shank work boots:
- hard hat;
- gloves, as needed;
- safety glasses;
- hearing protection;
- equipment replacements are available as needed.

4.2 Level C

Level C PPE shall be donned when the concentrations of measured total organic vapors in the breathing zone exceed background concentrations (using a portable OVA, or equivalent), but are less than 5 ppm. The specifications on the APR filters used must be appropriate for contaminants identified or expected to be encountered. Level C PPE shall be donned when the identified contaminants have adequate warning properties and criteria for using APR have been met. Level C PPE consists of:

- chemical resistant or coated tyvek coveralls;
- steel-toe and steel-shank workboots;
- chemical resistant overboots or disposable boot covers;
- disposable inner gloves (surgical gloves);
- disposable outer gloves;
- full face APR fitted with organic vapor/dust and mist filters or filters appropriate for the identified or expected contaminants;
- hard hat;
- splash shield, as needed; and,
- ankles/wrists taped with duct tape.

The site safety officer will verify if Level C is appropriate by checking organic vapor concentrations using compound and/or class-specific detector tubes.



The exact PPE ensemble is decided on a site-by-site basis by the Site Safety Officer with the intent to provide the most protective and efficient worker PPE.

4.3 Activity-Specific Levels of Personal Protection

The required level of PPE is activity-specific and is based on air monitoring results (Section 4.0) and properties of identified or expected contaminants. It is expected that site work will be performed in Level D. If air monitoring results indicate the necessity to upgrade the level of protection engineering controls (i.e. Facing equipment away from the wind and placing site personnel upwind of drilling locations, active venting, etc.) will be implemented before requiring the use of respiratory protection.



5.0 AIR MONITORING AND ACTION LEVELS

29 CFR 1910.120(h) specifies that monitoring shall be performed where there may be a question of employee exposure to hazardous concentrations of hazardous substances in order to assure proper selection of engineering controls, work practices and personal protective equipment so that employees are not exposed to levels which exceed permissible exposure limits, or published exposure levels if there are no permissible exposure limits, for hazardous substances.

5.1 **Air Monitoring Requirements**

If excavation work is performed, air will be monitored for VOCs with a portable ION Science 3000EX photoionization detector, or the equivalent. If necessary, Lower Explosive Limit (LEL) and oxygen will be monitored with a Combustible Gas Indicator (CGI). If appropriate, fugitive dust will be monitored using a MiniRam Model PDM-3 aerosol monitor. Air will be monitored when any of the following conditions apply:

- initial site entry;
- during any work where a potential IDLH condition or flammable atmosphere could develop;
- excavation work begins on another portion of the site;
- contaminants, other than those previously identified, have been discovered;
- each time a different task or activity is initiated;
- during trenching and/or excavation work.

The designated site safety officer will record air monitoring data and ensure that air monitoring instruments are calibrated and maintained in accordance with manufacturer's specifications. Instruments will be zeroed daily and checked for accuracy. Monitoring results will be recorded in a field notebook and will be transferred to instrument reading logs.

5.2 Work Stoppage Responses

The following responses will be initiated whenever one or more of the action levels necessitating a work stoppage are exceeded:

- 1 The SSO will be consulted immediately
- 2 All personnel (except as necessary for continued monitoring and contaminant migration, if applicable) will be cleared from the work area (eg from the exclusion zone).
- 3 Monitoring will be continued until intrusive work resumes.

Action Levels During Excavation Activities

Instrument readings will be taken in the breathing zone above the excavation pit unless otherwise noted. Each action level is independent of all other action levels in determining responses.

Organic Vapors (PID)	LEL %	Responses
0-1 ppm above background	0%	Continue excavating
		Level D protection
		• Continue monitoring every 10 minutes



1-5 ppm Above Background, Sustained Reading	1-10%	 Continue excavating Go to Level C protection or employ engineering controls Continue monitoring every 10 minutes
5-25 ppm Above Background, Sustained Reading	10-20%	 Discontinue excavating, unless PID is only action level exceeded. Level C protection or employ engineering controls Continue monitoring for organic vapors 200 ft downwind Continuous monitoring for LEL at excavation pit
>25 ppm Above Background, Sustained Reading	>20%	 Discontinue excavating Withdraw from area, shut off all engine ignition sources. Allow pit to vent Continuous monitoring for organic vapors 200 ft downwind.

Notes: Air monitoring will occur in the breathing zone 30 inches above the excavation pit. Readings may also be taken in the excavation pit but will not be used for action levels.

If action levels for any one of the monitoring parameters are exceeded, the appropriate responses listed in the right hand column should be taken. If instrument readings do not return to acceptable levels after the excavation pit has been vented for a period of greater than one-half hour, a decision will then be made whether or not to seal the pit with suppressant foam.

If, during excavation activities, downwind monitoring PID readings are greater than 5 ppm above background for more than one-half hour, excavation will stop until sustained levels are less then 5 ppm (see Community Air Monitoring Plan).

6.0 SITE CONTROL

6.1 Work Zones

The primary purpose of site controls is to establish the perimeter of a hazardous area, to reduce the migration of contaminants into clean areas, and to prevent access or exposure to hazardous materials by unauthorized persons. When operations are to take place involving hazardous materials, the site safety officer will establish an exclusion zone, a decontamination zone, and a support zone. These zones "float" (move around the site) depending on the tasks being performed on any given day. The site safety officer will outline these locations before work begins and when zones change. The site safety officer records this information in the site log book. If contamination is encountered then the Site Safety officer will establish the zones as follows:

Tasks requiring OSHA 40-hour Hazardous Waste Operations and Emergency Response Operations training are carried out in the exclusion zone. The exclusion zone is defined by the site safety officer but will typically be a 50-foot area around work activities. Gross decontamination (as determined by the site Health and Safety Officer) is conducted in the exclusion zone; all other decontamination is performed in the decontamination zone or trailer.

Protective equipment is removed in the decontamination zone. Disposable protective equipment is stored in receptacles staged in the decontamination zone, and non-disposable equipment is decontaminated. All personnel and equipment exit the exclusion zone through the decontamination zone. If a decontamination trailer is provided the first aid equipment, an eye wash unit, and drinking water are kept in the decontamination trailer.

The support zone is used for vehicle parking, daily safety meetings, and supply storage. Eating, drinking, and smoking are permitted only in the support zone. When a decontamination trailer is not provided, the eye wash unit, first aid equipment, and drinking water are kept at a central location designated by the site safety officer.

6.2 General Site Work

A general excavation contractor may complete the site excavation/grading as needed for the footing installation, or as deemed necessary by the Interim Remedial Measure Work Plan and/or Project Manager. All onsite employees must have obtained OSHA 24-hour Hazardous Waste Operations and Emergency Response Operations training prior to performing soil disturbing activities.

7.0 CONTINGENCY PLAN/EMERGENCY RESPONSE PLAN

Site personnel must be prepared in the event of an emergency. Emergencies can take many forms: illnesses, injuries, chemical exposure, fires, explosions, spills, leaks, releases of harmful contaminants, or sudden changes in the weather.

Emergency telephone numbers and a map to the hospital will be posted in the command post. Site personnel should be familiar with the emergency procedures, and the locations of site safety, first aid, and communication equipment.

7.1 **Emergency Equipment On-site**

Private telephones: Site personnel.

Two-way radios: Site personnel where necessary.

On-site vehicle horns*. Emergency Alarms:

First aid kits: On-site, in vehicles or office.

Fire extinguisher: On-site, in office or on equipment.

7.2 **Emergency Telephone Numbers**

General Emergencies	911
New York City Police	911
New York Community Hospital	1-718-692-5300
NYSDEC Spills Division	1-800-457-7362
NYSDEC Division of Env. Remediation	1-718-482-4900
NYCDEP	1-718-699-9811
NYC Department of Health	1-212-788-4711
NYC Fire Department	911
National Response Center	1-800-424-8802
Poison Control	1-212-340-4494
Site Safety Officer	1-631-504-6000
Alternate Site Safety Officer	1-631-504-6000

7.3 Personnel Responsibilities During an Emergency

The project manager is primarily responsible for responding to and correcting any emergency situations. However, in the absence of the project manager, the site safety officer shall act as the project manager's on-site designee and perform the following tasks:

Take appropriate measures to protect personnel including: withdrawal from the exclusion zone, evacuate and secure the site, or upgrade/downgrade the level of protective clothing and respiratory protection;

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^{*} Horns: Air horns will be supplied to personnel at the discretion of the project superintendent or site safety officer.

- Ensure that appropriate federal, state, and local agencies are informed and emergency response plans are coordinated. In the event of fire or explosion, the local fire department should be summoned immediately. If toxic materials are released to the air, the local authorities should be informed in order to assess the need for evacuation:
- Ensure appropriate decontamination, treatment, or testing for exposed or injured personnel;
- Determine the cause of incidents and make recommendations to prevent recurrence; and,
- Ensure that all required reports have been prepared.

The following key personnel are planned for this project:

Project Manager Mrs. Chawinie Reilly (631) 504-6000 • Site Safety Officer Mr. Kevin Waters (631) 504-6000

7.4 **Medical Emergencies**

A person who becomes ill or injured in the exclusion zone will be decontaminated to the maximum extent possible. If the injury or illness is minor, full decontamination will be completed and first aid administered prior to transport. First aid will be administered while waiting for an ambulance or paramedics. A Field Accident Report (Appendix D) must be filled out for any injury.

A person transporting an injured/exposed person to a clinic or hospital for treatment will take the directions to the hospital (Appendix D) and information on the chemical(s) to which they may have been exposed (Appendix C).

7.5 Fire or Explosion

In the event of a fire or explosion, the local fire department will be summoned immediately. The site safety officer or his designated alternate will advise the fire commander of the location, nature and identification of the hazardous materials on-site. If it is safe to do so, site personnel may:

- use fire fighting equipment available on site; or,
- remove or isolate flammable or other hazardous materials that may contribute to the fire.

7.6 **Evacuation Routes**

Evacuation routes established by work area locations for each site will be reviewed prior to commencing site operations. As the work areas change, the evacuation routes will be altered accordingly, and the new route will be reviewed.

Under extreme emergency conditions, evacuation is to be immediate without regard for equipment. The evacuation signal will be a continuous blast of a vehicle horn, if possible, and/or by verbal/radio communication. When evacuating the site, personnel will follow these

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instructions:

- Keep upwind of smoke, vapors, or spill location.
- Exit through the decontamination corridor if possible.
- If evacuation through the decontamination corridor is not possible, personnel should remove contaminated clothing once they are in a safe location and leave it near the exclusion zone or in a safe place.
- The site safety officer will conduct a head count to ensure that all personnel have been evacuated safely. The head count will be correlated to the site and/or exclusion zone entry/exit log.
- If emergency site evacuation is necessary, all personnel are to escape the emergency situation and decontaminate to the maximum extent practical.

7.7 **Spill Control Procedures**

Spills associated with site activities may be attributed to project equipment and include gasoline, diesel and hydraulic oil. In the event of a leak or a release, site personnel will inform their supervisor immediately, locate the source of spillage and stop the flow if it can be done safely. A spill containment kit including absorbent pads, booms and/or granulated speedy dry absorbent material will be available to site personnel to facilitate the immediate recovery of the spilled material. Daily inspections of site equipment components including hydraulic lines, fuel tanks, etc. will be performed by their respective operators as a preventative measure for equipment leaks and to ensure equipment soundness. In the event of a spill, site personnel will immediately notify the NYSDEC (1-800-457-7362), and a spill number will be generated.

7.8 Vapor Release Plan

If work zone organic vapor (excluding methane) exceeds 5 ppm, then a downwind reading will be made either 200 feet from the work zone or at the property line, whichever is closer. If readings at this location exceed 5 ppm over background, the work will be stopped.

If 5 ppm of VOCs are recorded over background on a PID at the property line, then an off-site reading will be taken within 20 feet of the nearest residential or commercial property, whichever is closer. If efforts to mitigate the emission source are unsuccessful for 30 minutes, then the designated site safety officer will:

contact the local police;

Environmental Business Consultants

- continue to monitor air every 30 minutes, 20 feet from the closest off-site property. If two successive readings are below 5 ppm (non-methane), off-site air monitoring will be halted.
- All property line and off site air monitoring locations and results associated with vapor releases will be recorded in the site safety log book.



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APPENDIX A SITE SAFETY ACKNOWLEDGEMENT FORM

DAILY BREIFING SIGN-IN SHEET

Date: Per	rson Conducting Briefing:			
Project Name and Location:				
. AWARENESS (topics discussed, special safety concerns, recent incidents, etc):				
2. OTHER ISSUES (HASP changes, attendee com	ments, etc):			
3. ATTENDEES (Print Name):				
1.	11.			
2.	12.			
3.	13.			
4.	14.			
5.	15.			
6.	16.			
7.	17.			
8.	18.			
9.	19.			
10.	20.			

APPENDIX B SITE SAFETY PLAN AMENDMENTS

SITE SAFETY PLAN AMENDMENT FORM

Site Safety Plan Amendment #:		
Site Name:		
Reason for Amendment:		
Alternative Procedures:		
Required Changes in PPE:		
Project Superintendent (signature)	Date	
Health and Safety Consultant (signature)	Date	
	= 3335	
Site Safety Officer (signature)	Date	

APPENDIX C CHEMICAL HAZARDS

CHEMICAL HAZARDS

The attached International Chemical Safety Cards are provided for contaminants of concern that have been identified in soils and/or groundwater at the site.

1,2,4-TRIMETHYLBENZENE











 $\begin{array}{c} \text{Pseudocumene} \\ \text{C}_9 \text{H}_{12} \end{array}$

Molecular mass: 120,2

ICSC # 1433 CAS # 95-63-6 RTECS # DC3325000

UN # 1993

EC# 601-043-00-3

March 06, 2002 Peer reviewed



ICSC: 1433

TYPES OF HAZARD/ EXPOSURE	ACUTE HAZARDS/ SYMPTOMS	PREVENTION	FIRST AID/ FIRE FIGHTING
FIRE	Flammable.	NO open flames, NO sparks, and NO smoking.	Alcohol-resistant foam, dry powder, carbon dioxide.
EXPLOSION	Above 44°C explosive vapour/air mixtures may be formed.	Above 44°C use a closed system, ventilation, and explosion-proof electrical equipment. Prevent build-up of electrostatic charges (e.g., by grounding).	In case of fire: keep drums, etc., cool by spraying with water.
EXPOSURE		PREVENT GENERATION OF MISTS!	
•INHALATION	Confusion. Cough. Dizziness. Drowsiness. Headache. Sore throat. Vomiting.	Ventilation, local exhaust, or breathing protection.	Fresh air, rest. Refer for medical attention.
•SKIN	Redness. Dry skin.	Protective gloves.	Rinse skin with plenty of water or shower.
•EYES	Redness. Pain.	Safety spectacles.	First rinse with plenty of water for several minutes (remove contact lenses if easily possible), then take to a doctor.
•INGESTION	(See Inhalation).	Do not eat, drink, or smoke during work.	Rinse mouth. Do NOT induce vomiting. Refer for medical attention.

SPILLAGE DISPOSAL	STORAGE	PACKAGING & LABELLING
	Fireproof. Separated from strong oxidants.	
	Well closed. Keep in a well-ventilated room.	
remaining liquid in sand or inert absorbent	 	N symbol
and remove to safe place. Do NOT wash		R: 10-20-36/37/38-51/53
away into sewer. Do NOT let this chemical		S: 2-26-61
enter the environment. Personal protection:		UN Hazard Class: 3
filter respirator for organic gases and		UN Packing Group: III
vapours.		

SEE IMPORTANT INFORMATION ON BACK

ICSC: 1433

Prepared in the context of cooperation between the International Programme on Chemical Safety & the Commission of the European Communities (C) IPCS CEC 1994. No modifications to the International version have been made except to add the OSHA PELs, NIOSH RELs and NIOSH IDLH values.

1,2,4-TRIMETHYLBENZENE

I	PHYSICAL STATE; APPEARANCE: COLOURLESS LIQUID, WITH CHARACTERISTIC	ROUTES OF EXPOSURE: The substance can be absorbed into the body by		
M	ODOUR.	inhalation.		
P	PHYSICAL DANGERS:	INHALATION RISK: A harmful contamination of the air will be reached		
О	CHEMICAL DANGERS	rather slowly on evaporation of this substance at 20°C;		
R	CHEMICAL DANGERS: The substance decomposes on burning producing toxic	on spraying or dispersing, however, much faster.		
T	and irritating fumes Reacts violently with strong oxidants causing fire and explosion hazard.	EFFECTS OF SHORT-TERM EXPOSURE: The substance is irritating to the eyes the skin and the respiratory tract If this liquid is swallowed, aspiration		
A	OCCUPATIONAL EXPOSURE LIMITS: TLV: (as mixed isomers) 25 ppm as TWA (ACGIH	into the lungs may result in chemical pneumonitis. The substance may cause effects on the central nervous		
N	2004).	system		
Т	MAK: (as mixed isomers) 20 ppm 100 mg/m³ Peak limitation category: II(2) Pregnancy risk group: C (DFG 2004).	EFFECTS OF LONG-TERM OR REPEATED EXPOSURE:		
D	OSHA PEL±: none NIOSH REL: TWA 25 ppm (125 mg/m³)	The liquid defats the skin. Lungs may be affected by repeated or prolonged exposure, resulting in chronic		
A	NIOSH IDLH: N.D. See: <u>IDLH INDEX</u>	bronchitis The substance may have effects on the central nervous system blood See Notes.		
Т				
A				
PHYSICAL PROPERTIES	Boiling point: 169°C Melting point: -44°C Relative density (water = 1): 0.88 Solubility in water: very poor Relative vapour density (air = 1): 4.1	Relative density of the vapour/air-mixture at 20°C (air = 1): 1.01 Flash point: 44°C c.c. Auto-ignition temperature: 500°C Explosive limits, vol% in air: 0.9-6.4 Octanol/water partition coefficient as log Pow: 3.8		
ENVIDONMENTAL	The substance is toxic to aquatic organisms. Bioaccumulation of this chemical may occur in fish.			

ENVIRONMENTAL DATA



ICSC: 1433

NOTES

Use of alcoholic beverages enhances the harmful effect. Depending on the degree of exposure, periodic medical examination is suggested. See also ICSC 1155 1,3,5-Trimethylbenzene (Mesitylene), ICSC 1362 1,2,3-Trimethylbenzene (Hemimellitene), ICSC 1389 Trimethyl benzene (mixed isomers). 1,3,5-Trimethylbenzene (Mesitylene) is classified as a marine pollutant.

> Transport Emergency Card: TEC (R)-30GF1-III NFPA Code: H0; F2; R0;

ADDITIONAL INFORMATION

ICSC: 1433 1,2,4-TRIMETHYLBENZENE

(C) IPCS, CEC, 1994

IMPORTANT LEGAL NOTICE:

1,3,5-TRIMETHYLBENZENE











Molecular mass: 120.2

ICSC # 1155 CAS # 108-67-8 RTECS # <u>OX6825000</u>

UN # 2325

EC# 601-025-00-5

March 06, 2002 Peer reviewed



ICSC: 1155

TYPES OF HAZARD/ EXPOSURE	ACUTE HAZA	PREVENTION	FIRST AID/ FIRE FIGHTING
FIRE	Flammable.	NO open flames, NO sparks, and N smoking.	Alcohol-resistant foam, dry powder, carbon dioxide.
EXPLOSION	Above 50°C explosive v mixtures may be formed	Above 50°C use a closed system, ventilation, and explosion-proof electrical equipment. Prevent build-of electrostatic charges (e.g., by grounding).	In case of fire: keep drums, etc., cool by spraying with water.
EXPOSURE		PREVENT GENERATION OF MISTS!	
•INHALATION	Confusion. Cough. Dizz Drowsiness. Headache. Vomiting.	Ventilation, local exhaust, or breathing protection.	Fresh air, rest. Refer for medical attention.
•SKIN	Redness. Dry skin.	Protective gloves.	Remove contaminated clothes. Rinse skin with plenty of water or shower.
•EYES	Redness. Pain.	Safety spectacles.	First rinse with plenty of water for several minutes (remove contact lenses if easily possible), then take to a doctor.
•INGESTION	(See Inhalation).	Do not eat, drink, or smoke during work.	Rinse mouth. Do NOT induce vomiting. Refer for medical attention.

SPILLAGE DISPOSAL	STORAGE	PACKAGING & LABELLING
	Fireproof. Separated from strong oxidants.	
1	Well closed. Keep in a well-ventilated room.	Marine pollutant.
remaining liquid in sand or inert absorbent		Xi symbol
and remove to safe place. Do NOT wash		N symbol
away into sewer. Do NOT let this chemical		R: 10-37-51/53
enter the environment. (Extra personal		S: 2-61
protection: filter respirator for organic gases		UN Hazard Class: 3
and vapours.)		UN Packing Group: III

SEE IMPORTANT INFORMATION ON BACK

ICSC: 1155

Prepared in the context of cooperation between the International Programme on Chemical Safety & the Commission of the European Communities (C) IPCS CEC 1994. No modifications to the International version have been made except to add the OSHA PELs, NIOSH RELs and NIOSH IDLH values.

1,3,5-TRIMETHYLBENZENE

I	PHYSICAL STATE; APPEARANCE:	ROUTES OF EXPOSURE:
M	COLOURLESS LIQUID , WITH CHARACTERISTIC ODOUR.	The substance can be absorbed into the body by inhalation.
P	PHYSICAL DANGERS:	INHALATION RISK:
О		A harmful contamination of the air will be reached rather slowly on evaporation of this substance at 20°C;
R	CHEMICAL DANGERS: The substance decomposes on burning producing toxic	on spraying or dispersing, however, much faster.
T	and irritating fumes. Reacts violently with strong oxidants causing fire and explosion hazard.	EFFECTS OF SHORT-TERM EXPOSURE: The substance is irritating to the eyes the skin and the respiratory tract If this liquid is swallowed, aspiration
A	OCCUPATIONAL EXPOSURE LIMITS:	into the lungs may result in chemical pneumonitis. The
N	TLV (as mixed isomers): 25 ppm; (ACGIH 2001). MAK (all isomers): 20 ppm; 100 mg/m ³ ; class II 1 ©	substance may cause effects on the central nervous system.
Т	(2001) OSHA PEL±: none NIOSH REL: TWA 25 ppm (125 mg/m³)	EFFECTS OF LONG-TERM OR REPEATED EXPOSURE:
D	NIOSH IDLH: N.D. See: <u>IDLH INDEX</u>	The liquid defats the skin. Lungs may be affected by repeated or prolonged exposure, resulting in chronic bronchitis. The substance may have effects on the
A		central nervous system blood See Notes.
Т		
A		
PHYSICAL PROPERTIES	Boiling point: 165°C Melting point: -45°C Relative density (water = 1): 0.86 Solubility in water: very poor Vapour pressure, kPa at 20°C: 0.25	Relative vapour density (air = 1): 4.1 Relative density of the vapour/air-mixture at 20°C (air = 1): 1.01 Flash point: 50°C (c.c.) Auto-ignition temperature: 550°C Octanol/water partition coefficient as log Pow: 3.42
ENVIRONMENTAL	The substance is harmful to aquatic organisms. Bioaccum	nulation of this chemical may occur in fish.

NOTES

Use of alcoholic beverages enhances the harmful effect. Depending on the degree of exposure, periodic medical examination is indicated. See ICSC 1433 1,2,4-Trimethylbenzene (Pseudocumene), ICSC 1362 1,2,3-Trimethylbenzene (Hemimellitene), ICSC 1389 Trimethylbenzene (mixed isomers).

Transport Emergency Card: TEC (R)-30S2325 NFPA Code: H0; F2; R0

ICSC: 1155

ADDITIONAL INFORMATION

ICSC: 1155 1,3,5-TRIMETHYLBENZENE

(C) IPCS, CEC, 1994

IMPORTANT LEGAL NOTICE:

DATA

ETHYLBENZENE











Ethylbenzol Phenylethane EB C_8H_{10} / $C_6H_5C_2H_5$ Molecular mass: 106.2

ICSC # 0268 CAS # 100-41-4 RTECS # <u>DA0700000</u>

UN # 1175

EC # 601-023-00-4 March 13, 1995 Validated



ICSC: 0268

TYPES OF HAZARD/ EXPOSURE	ACUTE HAZARDS/ SYMPTOMS	PREVENTION	FIRST AID/ FIRE FIGHTING
FIRE	Highly flammable.	NO open flames, NO sparks, and NO smoking.	Powder, AFFF, foam, carbon dioxide.
EXPLOSION	Vapour/air mixtures are explosive.	Closed system, ventilation, explosion- proof electrical equipment and lighting. Do NOT use compressed air for filling, discharging, or handling.	In case of fire: keep drums, etc., cool by spraying with water.
EXPOSURE		PREVENT GENERATION OF MISTS!	
•INHALATION	Cough. Dizziness. Drowsiness. Headache.	Ventilation, local exhaust, or breathing protection.	Fresh air, rest. Refer for medical attention.
•SKIN	Dry skin. Redness.	Protective gloves.	Remove contaminated clothes. Rinse and then wash skin with water and soap.
•EYES	Redness. Pain. Blurred vision.	Face shield or eye protection in combination with breathing protection.	First rinse with plenty of water for several minutes (remove contact lenses if easily possible), then take to a doctor.
•INGESTION	(Further see Inhalation).	Do not eat, drink, or smoke during work.	Rinse mouth. Give a slurry of activated charcoal in water to drink. Refer for medical attention.

SPILLAGE DISPOSAL	STORAGE	PACKAGING & LABELLING
Ventilation. Collect leaking liquid in covered containers. Absorb remaining liquid in sand or inert absorbent and remove to safe place. Do NOT wash away into sewer. Personal protection: A filter respirator for organic gases and vapours.		F symbol Xn symbol R: 11-20 S: 2-16-24/25-29 UN Hazard Class: 3 UN Packing Group: II

SEE IMPORTANT INFORMATION ON BACK

ICSC: 0268

Prepared in the context of cooperation between the International Programme on Chemical Safety & the Commission of the European Communities (C) IPCS CEC 1994. No modifications to the International version have been made except to add the OSHA PELs, NIOSH RELs and NIOSH IDLH values.

ETHYLBENZENE ICSC: 0268

COLOURLESS LIQUID , WITH AROMATIC ODOUR. PHYSICAL DANGERS: The vapour mixes well with air, explosive mixtures are easily formed. R CHEMICAL DANGERS: Reacts with strong oxidants. Attacks plastic and rubber. TLV: 100 ppm as TWA 125 ppm as STEL A3 (confirmed animal carcinogen with unknown relevance to humans); BEI issued (ACGIH 2005). MAK: skin absorption (H); Carcinogen category: 3A; (DFG 2004). OSHA PEL:: TWA 100 ppm (435 mg/m³) SI 125 ppm (545 mg/m³) NIOSH IBLH: 800 ppm 10% LEL See: 100414 PHYSICAL PROPERTIES Boiling point: 136°C Melting point: 95°C Melting point: 95°C Melting point: 95°C Melting point: 95°C Solubility in water, g/100 ml at 20°C: 0.015 Vapour pressure, kPa at 20°C: 0.9 Solubility in water g/100 ml at 20°C: 0.015 Vapour pressure, kPa at 20°C: 0.9 Solubility in water g/100 ml at 20°C: 0.015 Vapour pressure, kPa at 20°C: 0.9 Relative density of the vapour/air-mixture at 20°C (air = 1): 1.02 ENVIRONMENTAL DATA The substance is harmful to aquatic organisms. The substance can be absorbed into the body by inhalation of its vapour, through the skin and by ingestion. The substance are absoruer, through the skin and by ingestion. INHALATION RISK: A harmful contamination of the air will be reached rather slowly on evaporation of this substance at 20°C. EFFECTS OF SHORT-TERM EXPOSURE: The substance is irritating to the eyes the skin and the respiratory tract Swallowing the liquid may cause aspiration into thugs with the risk of chemical pneumonitis. The substance is irritating to the eyes the skin and the respiratory tract Swallowing the liquid may cause aspiration into thugs with the risk of chemical pneumonitis. The substance at 20°C (air = 1): 1.02 EFFECTS OF SHORT-TERM EXPOSURE: The substance at 20°C of C. Relative density of the vapour/air-mixture at 20°C (air = 1): 1.02 Explosive limits, vol% in air: 1.0-6.7 Octanol/water partition coefficient as log Pow: 3.2 ENVIRONMENTAL DATA The substance is power and the respirator into the eyes the skin and the respirator into the ups with the ris	I	PHYSICAL STATE; APPEARANCE:	ROUTES OF EXPOSURE:					
PHYSICAL DANGERS: The vapour mixes well with air, explosive mixtures are easily formed. R CHEMICAL DANGERS: Reacts with strong oxidants. Attacks plastic and rubber. T OCCUPATIONAL EXPOSURE LIMITS: TLV: 100 ppm as TWA 125 ppm as STEL A3 (confirmed animal carcinogen with unknown relevance to humans); BEI issued (ACGIH 2005). MAK: skin absorption (H); Carcinogen category: 3A; (DFG 2004). OSHA PEL½: TWA 100 ppm (435 mg/m³) NIOSH REL: TWA 100 ppm (435 mg/m³) ST 125 ppm (545 mg/m³) NIOSH IDLH: 800 ppm 10%LEL See: 100414 PHYSICAL PROPERTIES Boiling point: 136°C Melting point: 95°C Relative density (water = 1): 0.9 Relative density (water = 1): 0.9 Relative vapour density (air = 1): 3.7 ENVIRONMENTAL DATA NOTES	M		inhalation of its vapour, through the skin and by					
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Reacts with strong oxidants. Attacks plastic and rubber. OCCUPATIONAL EXPOSURE LIMITS: TLV: 100 ppm as TWA 125 ppm as STEL A3 (confirmed animal carcinogen with unknown relevance to humans); BEI issued (ACGIH 2005). MAK: skin absorption (H); Carcinogen category: 3A; (DFG 2004). DSHA PEL¹: TWA 100 ppm (435 mg/m³) NIOSH REL: TWA 100 ppm (435 mg/m³) ST 125 ppm (545 mg/m³) NIOSH IDLH: 800 ppm 10%LEL See: 100414 PHYSICAL PROPERTIES Boiling point: 136°C Melting point: -95°C Relative density (water = 1): 0.9 Solubility in water, g/100 ml at 20°C: 0.015 Vapour pressure, kPa at 20°C: 0.9 Relative vapour density (wair = 1): 3.7 ENVIRONMENTAL DATA Reacts with strong oxidants. Attacks plastic and rubber. The substance is irritating to the eyes the skin and the respiratory tract Swallowing the liquid may cause aspiration into the lungs with the risk of chemical pneumonitis. The substance may cause effects on the central nervous system Exposure far above the OEL could cause lowering of consciousness. EFFECTS OF SHORT-TERM EXPOSURE: The substance is irritating to the eyes the skin and the respiratory tract Swallowing the liquid may cause aspiration into the lungs with the risk of chemical pneumonitis. The substance may cause effects on the central nervous system Exposure far above the OEL could cause lowering of consciousness. EFFECTS OF SHORT-TERM EXPOSURE: The substance is irritating to the eyes the skin and the respiratory tract Swallowing the liquid may cause aspiration into the lungs with the risk of chemical pneumonitis. The substance may cause effects on the central nervous system Exposure far above the OEL could cause lowering of consciousness. EFFECTS OF SHORT-TERM EXPOSURE: The substance is irritating to the eyes the skin and the respiration into the lungs with the risk of chemical pneumonitis. The substance is spiration into the lungs with the risk of central nervous system Exposure aspiration into the lungs with the risk of central nervous system Exposure far above the OEL could cause lowering of consciou			A harmful contamination of the air will be reached					
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To humans); BEI issued (ACGIH 2005). MAK: skin absorption (H); Carcinogen category: 3A; (DFG 2004). OSHA PEL†: TWA 100 ppm (435 mg/m³) NIOSH REL: TWA 100 ppm (435 mg/m³) ST 125 ppm (545 mg/m³) NIOSH IDLH: 800 ppm 10%LEL See: 100414 PHYSICAL PROPERTIES Boiling point: 136°C Melting point: -95°C Relative density (water = 1): 0.9 Solubility in water, g/100 ml at 20°C: 0.015 Vapour pressure, kPa at 20°C: 0.9 Relative vapour density (air = 1): 3.7 Carcinogen category: 3A; (DFG 2004). ENVIRONMENTAL DATA to humans); BEI issued (ACGIH 2005). MAK: skin absorption (H); Carcinogen category: 3A; (DFG 2004). EFFECTS OF LONG-TERM OR REPEATED EXPOSURE: Repeated or prolonged contact with skin may cause dermatitis. Repeated or prolonged contact with skin may cause dermatitis. Repeated or prolonged contact with skin may cause dermatitis. Repeated or prolonged contact with skin may cause dermatitis. Repeated or prolonged contact with skin may cause dermatitis. Repeated or prolonged contact with skin may cause dermatitis. Repeated or prolonged contact with skin may cause dermatitis. Repeated or prolonged contact with skin may cause dermatitis. Repeated or prolonged contact with skin may cause dermatitis. Repeated or prolonged contact with skin may cause dermatitis. Repeated or prolonged contact with skin may cause dermatitis. Repeated or prolonged contact with skin may cause dermatitis. Repeated or prolonged contact with skin may cause dermatitis. Repeated or prolonged contact with skin may cause dermatitis. PAUS A. 100 PERONGE A. 100 PER	A	TLV: 100 ppm as TWA 125 ppm as STEL A3	aspiration into the lungs with the risk of chemical					
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NIOSH REL: TWA 100 ppm (435 mg/m³) ST 125 ppm (545 mg/m³) NIOSH IDLH: 800 ppm 10%LEL See: 100414 T A PHYSICAL PROPERTIES Boiling point: 136°C Melting point: -95°C Relative density (water = 1): 0.9 Solubility in water, g/100 ml at 20°C: 0.015 Vapour pressure, kPa at 20°C: 0.9 Relative vapour density (air = 1): 3.7 ENVIRONMENTAL DATA Repeated or prolonged contact with skin may cause dermatitis. Repeated or prolonged contact with skin may cause dermatitis. Repeated or prolonged contact with skin may cause dermatitis. Repeated or prolonged contact with skin may cause dermatitis. Repeated or prolonged contact with skin may cause dermatitis. Repeated or prolonged contact with skin may cause dermatitis. Repeated or prolonged contact with skin may cause dermatitis. NOTES	T	Carcinogen category: 3A;	Ç					
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PHYSICAL PROPERTIES Boiling point: 136°C Melting point: -95°C Relative density (water = 1): 0.9 Solubility in water, g/100 ml at 20°C: 0.015 Vapour pressure, kPa at 20°C: 0.9 Relative vapour density (air = 1): 3.7 ENVIRONMENTAL DATA Boiling point: 136°C Melting point: -95°C Relative density of the vapour/air-mixture at 20°C (air = 1): 1.02 Flash point: 18°C c.c. Auto-ignition temperature: 432°C Explosive limits, vol% in air: 1.0-6.7 Octanol/water partition coefficient as log Pow: 3.2 NOTES	A		dermatitis.					
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NOTES		Melting point: -95°C Relative density (water = 1): 0.9 Solubility in water, g/100 ml at 20°C: 0.015 Vapour pressure, kPa at 20°C: 0.9	1): 1.02 Flash point: 18°C c.c. Auto-ignition temperature: 432°C Explosive limits, vol% in air: 1.0-6.7					
		The substance is harmful to aquatic organisms.						
The odour warning when the exposure limit value is exceeded is insufficient.		NOTES						
	The odour warning who	The odour warning when the exposure limit value is exceeded is insufficient.						

Transport Emergency Card: TEC (R)-30S1175 or 30GF1-I+II

NFPA Code: H2; F3; R0

ADDITIONAL INFORMATION

ICSC: 0268 ETHYLBENZENE

(C) IPCS, CEC, 1994

IMPORTANT LEGAL NOTICE:

CUMENE ICSC: 0170











(1-Methylethyl)benzene 2-Phenylpropane Isopropylbenzene C₉H₁₂ / C₆H₅CH(CH₃)₂ Molecular mass: 120.2

ICSC # 0170 CAS # 98-82-8 RTECS # <u>GR8575000</u>

UN # 1918

EC # 601-024-00-X April 13, 2000 Peer reviewed

chemical enter the environment. Personal

and vapours.

protection: filter respirator for organic gases



TYPES OF HAZARD/ EXPOSURE	ACUTE HAZ SYMPTO		PREVENTION		FIRST AID/ FIRE FIGHTING
FIRE	Flammable.		NO open flames, NO sparks, ar smoking.	nd NO	Powder, AFFF, foam, carbon dioxide.
EXPLOSION	Above 31°C explosive mixtures may be formed		Above 31°C use a closed system ventilation, and explosion-proof electrical equipment. Prevent but of electrostatic charges (e.g., by grounding).	f uild-up	In case of fire: keep drums, etc., cool by spraying with water.
EXPOSURE			PREVENT GENERATION OF MISTS!	7	
•INHALATION	Dizziness. Ataxia. Drov Headache. Unconscious		Ventilation, local exhaust, or breathing protection.		Fresh air, rest. Refer for medical attention.
•SKIN	Dry skin.		Protective gloves. Protective clothing.		Remove contaminated clothes. Rinse and then wash skin with water and soap.
•EYES	Redness. Pain.		Safety spectacles.		First rinse with plenty of water for several minutes (remove contact lenses if easily possible), then take to a doctor.
•INGESTION	(See Inhalation).	Do not eat, drink, or smoke during work.		ring	Rinse mouth. Do NOT induce vomiting. Refer for medical attention.
SPILLAGE DISPOSAL			STORAGE	PA	CKAGING & LABELLING
Collect leaking and spilled liquid in sealable containers as far as possible. Absorb remaining liquid in sand or inert absorbent and remove to safe place. Do NOT let this			parated from strong oxidants, keep in the dark. Store only if	Marine Note: (nbol

SEE IMPORTANT INFORMATION ON BACK

N symbol

R: 10-37-51/53-65 S: 2-24-37-61-62

UN Hazard Class: 3 UN Packing Group: III

ICSC: 0170

Prepared in the context of cooperation between the International Programme on Chemical Safety & the Commission of the European Communities (C) IPCS CEC 1994. No modifications to the International version have been made except to add the OSHA PELs, NIOSH RELs and NIOSH IDLH values.

CUMENE ICSC: 0170

I	PHYSICAL STATE; APPEARANCE: COLOURLESS LIQUID, WITH CHARACTERISTIC	ROUTES OF EXPOSURE: The substance can be absorbed into the body by		
M	ODOUR.	inhalation and through the skin.		
P	PHYSICAL DANGERS: As a result of flow, agitation, etc., electrostatic charges	INHALATION RISK: A harmful contamination of the air will be reached		
О	can be generated.	rather slowly on evaporation of this substance at 20°C.		
R	CHEMICAL DANGERS: Reacts violently with acids and strong oxidants causing	EFFECTS OF SHORT-TERM EXPOSURE: The substance is irritating to the eyes and the skin		
T .	fire and explosion hazard. The substance can form explosive peroxides.	Swallowing the liquid may cause aspiration into the lungs with the risk of chemical pneumonitis. The		
A N	OCCUPATIONAL EXPOSURE LIMITS: TLV: 50 ppm as TWA (ACGIH 2004).	substance may cause effects on the central nervous system Exposure far above the OEL may result in unconsciousness.		
T	MAK: 50 ppm 250 mg/m³ Peak limitation category: II(4);	EFFECTS OF LONG-TERM OR REPEATED		
D	skin absorption (H); Pregnancy risk group: C; (DFG 2004).	EXPOSURE: Repeated or prolonged contact with skin may cause dermatitis.		
A	OSHA PEL: TWA 50 ppm (245 mg/m ³) skin NIOSH REL: TWA 50 ppm (245 mg/m ³) skin			
Т	NIOSH IDLH: 900 ppm 10%LEL See: <u>98828</u>			
A				
PHYSICAL PROPERTIES	Boiling point: 152°C Melting point: -96°C Relative density (water = 1): 0.90 Solubility in water: none Vapour pressure, Pa at 20°C: 427 Relative vapour density (air = 1): 4.2	Relative density of the vapour/air-mixture at 20°C (air = 1): 1.01 Flash point: 31°C c.c. Auto-ignition temperature: 420°C Explosive limits, vol% in air: 0.9-6.5 Octanol/water partition coefficient as log Pow: 3.66		
ENVIRONMENTAL DATA	The substance is toxic to aquatic organisms.			
	NOTES			
Check for peroxides pri	Check for peroxides prior to distillation; eliminate if found. Transport Emergency Card: TEC (R)-30S1918 or 30GF1-III NFPA Code: H2; F3; R1			
	ADDITIONAL INFORMA	TION		
ICSC: 0170	(C) IPCS, CEC, 1994	CUMENE		

IMPORTANT LEGAL NOTICE:

NAPHTHALENE ICSC: 0667











Naphthene $C_{10}H_8$

Molecular mass: 128.18

ICSC # 0667 CAS # 91-20-3 RTECS # QJ0525000

UN # 1334 (solid); 2304 (molten)

EC # 601-052-00-2 April 21, 2005 Validated



TYPES OF HAZARD/ EXPOSURE	ACUTE HAZARDS/ SYMPTOMS	PREVENTION	FIRST AID/ FIRE FIGHTING
FIRE	Combustible.	NO open flames.	Powder, water spray, foam, carbon dioxide.
EAPLOSION	Above 80°C explosive vapour/air mixtures may be formed. Finely dispersed particles form explosive mixtures in air.	Prevent deposition of dust; closed system, dust explosion-proof electrical equipment and lighting.	
EXPOSURE		PREVENT DISPERSION OF DUST!	
•INHALATION	Headache. Weakness. Nausea. Vomiting. Sweating. Confusion. Jaundice. Dark urine.	Ventilation (not if powder), local exhaust, or breathing protection.	Fresh air, rest. Refer for medical attention.
•SKIN	MAY BE ABSORBED! (Further see Inhalation).	Protective gloves.	Rinse skin with plenty of water or shower.
•EYES		Safety spectacles.	First rinse with plenty of water for several minutes (remove contact lenses if easily possible), then take to a doctor.
•INGESTION	Abdominal pain. Diarrhoea. Convulsions. Unconsciousness. (Further see Inhalation).	Do not eat, drink, or smoke during work. Wash hands before eating.	Rest. Refer for medical attention.

SPILLAGE DISPOSAL	STORAGE	PACKAGING & LABELLING
organic gases and vapours. Do NOT let this	feedstuffs . Store in an area without drain or sewer access.	Do not transport with food and feedstuffs. Marine pollutant. Xn symbol N symbol R: 22-40-50/53 S: 2-36/37-46-60-61 UN Hazard Class: 4.1 UN Packing Group: III

SEE IMPORTANT INFORMATION ON BACK

ICSC: 0667

Prepared in the context of cooperation between the International Programme on Chemical Safety & the Commission of the European Communities (C) IPCS CEC 1994. No modifications to the International version have been made except to add the OSHA PELs, NIOSH RELs and NIOSH IDLH values.

NAPHTHALENE ICSC: 0667

I	PHYSICAL STATE; APPEARANCE:	ROUTES OF EXPOSURE:
M	WHITE SOLID IN VARIOUS FORMS , WITH CHARACTERISTIC ODOUR.	The substance can be absorbed into the body by inhalation, through the skin and by ingestion.
P	PHYSICAL DANGERS:	INHALATION RISK:
О	Dust explosion possible if in powder or granular form, mixed with air.	A harmful contamination of the air will be reached rather slowly on evaporation of this substance at 20°C. See Notes.
R	CHEMICAL DANGERS:	
Т	On combustion, forms irritating and toxic gases. Reacts with strong oxidants .	EFFECTS OF SHORT-TERM EXPOSURE: The substance may cause effects on the blood, resulting in lesions of blood cells (haemolysis). See Notes. The
A	OCCUPATIONAL EXPOSURE LIMITS: TLV: 10 ppm as TWA; 15 ppm as STEL; (skin); A4 (not	effects may be delayed. Exposure by ingestion may
N	classifiable as a human carcinogen); (ACGIH 2005).	
Т	MAK: skin absorption (H); Carcinogen category: 2; Germ cell mutagen group: 3B; (DFG 2004).	EFFECTS OF LONG-TERM OR REPEATED EXPOSURE: The substance may have effects on the blood, resulting
D	OSHA PEL <u>†</u> : TWA 10 ppm (50 mg/m ³) NIOSH REL: TWA 10 ppm (50 mg/m ³) ST 15 ppm (75	in chronic haemolytic anaemia. The substance may have effects on the eyes, resulting in the development of cataract. This substance is possibly carcinogenic to
A	mg/m ³) NIOSH IDLH: 250 ppm See: <u>91203</u>	humans.
T		
A		
PHYSICAL PROPERTIES	Boiling point: 218°C Sublimation slowly at room temperature Melting point: 80°C Density: 1.16 g/cm³ Solubility in water, g/100 ml at 25°C: none	Vapour pressure, Pa at 25°C: 11 Relative vapour density (air = 1): 4.42 Flash point: 80°C c.c. Auto-ignition temperature: 540°C Explosive limits, vol% in air: 0.9-5.9 Octanol/water partition coefficient as log Pow: 3.3
ENVIRONMENTAL DATA	The substance is very toxic to aquatic organisms. The sub aquatic environment.	stance may cause long-term effects in the
	NOTES	

Some individuals may be more sensitive to the effect of naphthalene on blood cells.

Transport Emergency Card: TEC (R)-41S1334 (solid); 41GF1-II+III (solid); 41S2304 (molten)

NFPA Code: H2; F2; R0;

ADDITIONAL INFORMATION

ICSC: 0667 NAPHTHALENE

(C) IPCS, CEC, 1994

IMPORTANT LEGAL NOTICE:

Material Safety Data Sheet

Normal-Butylbenzene, 99+%

ACC# 55434

Section 1 - Chemical Product and Company Identification

MSDS Name: Normal-Butylbenzene, 99+%

Catalog Numbers: AC107850000, AC107850050, AC107850250, AC107850500, AC107851000, AC107852500

AC107852500

For information in North America, call: 800-ACROS-01 For emergencies in the US, call CHEMTREC: 800-424-9300

Section 2 - Composition, Information on Ingredients

CAS#	Chemical Name	Percent	EINECS/ELINCS
104-51-8	n-Butylbenzene	>99	203-209-7

Section 3 - Hazards Identification

EMERGENCY OVERVIEW

Appearance: clear, colorless liquid. Flash Point: 59 deg C.

Warning! Flammable liquid and vapor. May cause eye and skin irritation. May cause respiratory and digestive tract irritation. The toxicological properties of this material have not been fully investigated.

Target Organs: Liver, nervous system.

Potential Health Effects

Eye: May cause eye irritation. The toxicological properties of this material have not been fully investigated. **Skin:** May cause skin irritation. The toxicological properties of this material have not been fully investigated. **Ingestion:** May cause gastrointestinal irritation with nausea, vomiting and diarrhea. The toxicological properties of this substance have not been fully investigated.

Inhalation: May cause respiratory tract irritation. The toxicological properties of this substance have not been fully investigated. Vapors may cause dizziness or suffocation.

Chronic: No information found.

Section 4 - First Aid Measures

Eyes: Flush eyes with plenty of water for at least 15 minutes, occasionally lifting the upper and lower eyelids. Get medical aid immediately.

Skin: Get medical aid. Flush skin with plenty of water for at least 15 minutes while removing contaminated clothing and shoes. Wash clothing before reuse.

Ingestion: Never give anything by mouth to an unconscious person. Get medical aid immediately. Do NOT induce vomiting. If conscious and alert, rinse mouth and drink 2-4 cupfuls of milk or water.

Inhalation: Remove from exposure and move to fresh air immediately. If not breathing, give artificial respiration. If breathing is difficult, give oxygen. Get medical aid.

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. Vapors may form an explosive mixture with air. During a fire, irritating and highly toxic gases may be generated by thermal decomposition or combustion. Will burn if involved in a fire. Use water spray to keep fire-exposed containers cool. Containers may explode in the heat of a fire. Flammable liquid and vapor. Vapors may be heavier than air. They can spread along the ground and collect in low or confined areas.

Extinguishing Media: For small fires, use dry chemical, carbon dioxide, water spray or alcohol-resistant foam. For large fires, use water spray, fog, or alcohol-resistant foam. Use water spray to cool fire-exposed containers. Water may be ineffective. Use agent most appropriate to extinguish fire. Do NOT use straight streams of water.

Flash Point: 59 deg C (138.20 deg F)

Autoignition Temperature: 412 deg C (773.60 deg F)

Explosion Limits, Lower: .80 vol %

Upper: 5.80 vol %

NFPA Rating: (estimated) Health: 1; Flammability: 2; 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. Clean up spills immediately, observing precautions in the Protective Equipment section. Remove all sources of ignition. Use a spark-proof tool. Provide ventilation. A vapor suppressing foam may be used to reduce vapors.

Section 7 - Handling and Storage

Handling: Wash thoroughly after handling. Remove contaminated clothing and wash before reuse. Use with adequate ventilation. 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. Keep container tightly closed. Keep away from heat, sparks and flame. Avoid ingestion and inhalation. Do not pressurize, cut, weld, braze, solder, drill, grind, or expose empty containers to heat, sparks or open flames.

Storage: Keep away from heat, sparks, and flame. Keep away from sources of ignition. Store in a tightly closed container. Store in a cool, dry, well-ventilated area away from incompatible substances. Flammables-area.

Section 8 - Exposure Controls, Personal Protection

Engineering Controls: Use adequate ventilation to keep airborne concentrations low. Use process enclosure, local exhaust ventilation, or other engineering controls to control airborne levels.

Exposure Limits

Chemical Name	ACGIH	NIOSH	OSHA - Final PELs
n-Butylbenzene	none listed	none listed	none listed

OSHA Vacated PELs: n-Butylbenzene: No OSHA Vacated PELs are listed for this chemical.

Personal Protective Equipment

Eyes: Wear appropriate protective eyeglasses or chemical safety goggles as described by OSHA's eye and face 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: Wear a NIOSH/MSHA or European Standard EN 149 approved full-facepiece airline respirator in the positive pressure mode with emergency escape provisions. 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: None reported. pH: Not available.

Vapor Pressure: 1.33 hPa @ 23 C

Vapor Density: 4.6

Evaporation Rate: Not available.

Viscosity: Not available.

Boiling Point: 183 deg C @ 760.00mm Hg **Freezing/Melting Point:**-88 deg C **Decomposition Temperature:**> 183 deg C

Solubility: insoluble

Specific Gravity/Density: .8600g/cm3

Molecular Formula:C10H14 Molecular Weight:134.22

Section 10 - Stability and Reactivity

Chemical Stability: Stable under normal temperatures and pressures.

Conditions to Avoid: Incompatible materials, ignition sources, excess heat, strong oxidants.

Incompatibilities with Other Materials: Oxidizing agents.

Hazardous Decomposition Products: Carbon monoxide, irritating and toxic fumes and gases, carbon dioxide.

Hazardous Polymerization: Has not been reported.

Section 11 - Toxicological Information

RTECS#:

CAS# 104-51-8: CY9070000

LD50/LC50: Not available.

Carcinogenicity:

CAS# 104-51-8: Not listed by ACGIH, IARC, NTP, or CA Prop 65.

Epidemiology: No information available. **Teratogenicity:** No information available.

Reproductive Effects: No information available.

Mutagenicity: No information available. **Neurotoxicity:** No information available.

Other Studies:

Section 12 - Ecological Information

Ecotoxicity: No data available. No information available.

Environmental: Rapidly volatilizes into the atmosphere where it is photochemically degraded by hydroxyl

radicals.

Physical: No information available. **Other:** No information available.

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: None listed.

Section 14 - Transport Information

	US DOT	Canada TDG
Shipping Name:	BUTYL BENZENES	No information available.
Hazard Class:	3	
UN Number:	UN2709	
Packing Group:	111	

Section 15 - Regulatory Information

US FEDERAL

TSCA

CAS# 104-51-8 is listed on the TSCA inventory.

Health & Safety Reporting List

CAS# 104-51-8: Effective 6/1/87, Sunset 12/19/95

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

None of the chemicals in this material have an RQ.

SARA Section 302 Extremely Hazardous Substances

None of the chemicals in this product have a TPQ.

SARA Codes

CAS # 104-51-8: immediate, fire.

Section 313 No chemicals are reportable under Section 313.

Clean Air Act:

This material does not contain any hazardous air pollutants.

This material does not contain any Class 1 Ozone depletors.

This material does not contain any Class 2 Ozone depletors.

Clean Water Act:

None of the chemicals in this product are listed as Hazardous Substances under the CWA.

None of the chemicals in this product are listed as Priority Pollutants under the CWA.

None of the chemicals in this product are listed as Toxic Pollutants under the CWA.

OSHA:

None of the chemicals in this product are considered highly hazardous by OSHA.

STATE

CAS# 104-51-8 can be found on the following state right to know lists: New Jersey, Pennsylvania, Massachusetts.

California Prop 65

California No Significant Risk Level: None of the chemicals in this product are listed.

European/International Regulations

European Labeling in Accordance with EC Directives

Hazard Symbols:

Not available.

Risk Phrases:

R 10 Flammable.

Safety Phrases:

S 16 Keep away from sources of ignition - No smoking.

S 24/25 Avoid contact with skin and eyes.

S 33 Take precautionary measures against static discharges.

S 37 Wear suitable gloves.

S 45 In case of accident or if you feel unwell, seek medical advice immediately (show the label where possible).

S 9 Keep container in a well-ventilated place.

S 28A After contact with skin, wash immediately with plenty of water

.

WGK (Water Danger/Protection)

CAS# 104-51-8: 1

Canada - DSL/NDSL

CAS# 104-51-8 is listed on Canada's DSL List.

Canada - WHMIS

This product has a WHMIS classification of B3, 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

Section 16 - Additional Information

MSDS Creation Date: 4/15/1998 Revision #4 Date: 3/16/2007

The information above is believed to be accurate and represents the best information currently available to us. However, we make no warranty of merchantability 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 Fisher 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 Fisher has been advised of the possibility of such damages.

Material Safety Data Sheet

Version 4.0 Revision Date 07/28/2010 Print Date 12/07/2011

1. PRODUCT AND COMPANY IDENTIFICATION

Product name : Propylbenzene

Product Number : P52407 Brand : Aldrich

Company : Sigma-Aldrich

3050 Spruce Street

SAINT LOUIS MO 63103

USA

Telephone : +1 800-325-5832 Fax : +1 800-325-5052 Emergency Phone # : (314) 776-6555

2. HAZARDS IDENTIFICATION

Emergency Overview

OSHA Hazards

Combustible Liquid

Target Organs

Lungs, Eyes, Kidney

GHS Label elements, including precautionary statements

Pictogram



Signal word Danger

Hazard statement(s)

H226 Flammable liquid and vapour.

H304 May be fatal if swallowed and enters airways.

H335 May cause respiratory irritation.

H401 Toxic to aquatic life.

Precautionary statement(s)

P261 Avoid breathing dust/ fume/ gas/ mist/ vapours/ spray.

P301 + P310 IF SWALLOWED: Immediately call a POISON CENTER or doctor/ physician.

P331 Do NOT induce vomiting.

HMIS Classification

Health hazard: 0
Chronic Health Hazard: *
Flammability: 2
Physical hazards: 0

NFPA Rating

Health hazard: 1
Fire: 2
Reactivity Hazard: 0

Potential Health Effects

Inhalation May be harmful if inhaled. May cause respiratory tract irritation.

Skin May be harmful if absorbed through skin. May cause skin irritation.

Eyes May cause eye irritation.

Ingestion

Aspiration hazard if swallowed - can enter lungs and cause damage. May be harmful if

swallowed.

3. COMPOSITION/INFORMATION ON INGREDIENTS

Synonyms : 1-Phenylpropane

Formula : C₉H₁₂

Molecular Weight : 120.19 g/mol

CAS-No. EC-No. Index-No. Concentration			
Propylbenzene			

4. FIRST AID MEASURES

General advice

Consult a physician. Show this safety data sheet to the doctor in attendance. Move out of dangerous area.

If inhaled

If breathed in, move person into fresh air. If not breathing give artificial respiration Consult a physician.

In case of skin contact

Wash off with soap and plenty of water. Consult a physician.

In case of eye contact

Rinse thoroughly with plenty of water for at least 15 minutes and consult a physician.

If swallowed

Do NOT induce vomiting. Never give anything by mouth to an unconscious person. Rinse mouth with water. Consult a physician.

5. FIRE-FIGHTING MEASURES

Suitable extinguishing media

For small (incipient) fires, use media such as "alcohol" foam, dry chemical, or carbon dioxide. For large fires, apply water from as far as possible. Use very large quantities (flooding) of water applied as a mist or spray; solid streams of water may be ineffective. Cool all affected containers with flooding quantities of water.

Special protective equipment for fire-fighters

Wear self contained breathing apparatus for fire fighting if necessary.

Further information

Use water spray to cool unopened containers.

6. ACCIDENTAL RELEASE MEASURES

Personal precautions

Use personal protective equipment. Avoid breathing vapors, mist or gas. Ensure adequate ventilation. Remove all sources of ignition. Beware of vapours accumulating to form explosive concentrations. Vapours can accumulate in low areas.

Environmental precautions

Prevent further leakage or spillage if safe to do so. Do not let product enter drains. Discharge into the environment must be avoided.

Methods and materials for containment and cleaning up

Contain spillage, and then collect with non-combustible absorbent material, (e.g. sand, earth, diatomaceous earth, vermiculite) and place in container for disposal according to local / national regulations (see section 13). Keep in suitable, closed containers for disposal.

7. HANDLING AND STORAGE

Precautions for safe handling

Avoid inhalation of vapour or mist.

Keep away from sources of ignition - No smoking. Take measures to prevent the build up of electrostatic charge.

Conditions for safe storage

Keep container tightly closed in a dry and well-ventilated place. Containers which are opened must be carefully resealed and kept upright to prevent leakage. Store in cool place.

8. EXPOSURE CONTROLS/PERSONAL PROTECTION

Contains no substances with occupational exposure limit values.

Personal protective equipment

Respiratory protection

Where risk assessment shows air-purifying respirators are appropriate use a full-face respirator with multi-purpose combination (US) or type ABEK (EN 14387) respirator cartridges as a backup to engineering controls. If the respirator is the sole means of protection, use a full-face supplied air respirator. Use respirators and components tested and approved under appropriate government standards such as NIOSH (US) or CEN (EU).

Hand protection

For prolonged or repeated contact use protective gloves.

Eye protection

Face shield and safety glasses

Skin and body protection

Choose body protection according to the amount and concentration of the dangerous substance at the work place.

Hygiene measures

Handle in accordance with good industrial hygiene and safety practice. Wash hands before breaks and at the end of workday.

9. PHYSICAL AND CHEMICAL PROPERTIES

Appearance

Form liquid, clear Colour colourless

Safety data

pH no data available

Melting point -99 °C (-146 °F) - lit.

Boiling point 159 °C (318 °F) - lit.

Flash point 42.0 °C (107.6 °F) - closed cup

Ignition temperature 450 °C (842 °F)

Lower explosion limit 0.8 %(V) Upper explosion limit 6 %(V)

Density 0.862 g/cm3 at 25 °C (77 °F)

Water solubility slightly soluble

10. STABILITY AND REACTIVITY

Chemical stability

Stable under recommended storage conditions.

Possibility of hazardous reactions

Vapours may form explosive mixture with air.

Conditions to avoid

Heat, flames and sparks.

Materials to avoid

Strong oxidizing agents

Hazardous decomposition products

Hazardous decomposition products formed under fire conditions. - Carbon oxides

11. TOXICOLOGICAL INFORMATION

Acute toxicity

LD50 Oral - rat - 6,040 mg/kg

Remarks: Behavioral:Somnolence (general depressed activity).

LC50 Inhalation - rat - 2 h - 65000 ppm

Skin corrosion/irritation

no data available

Serious eye damage/eye irritation

no data available

Respiratory or skin sensitization

no data available

Germ cell mutagenicity

no data available

Carcinogenicity

IARC: No component of this product present at levels greater than or equal to 0.1% is identified as probable,

possible or confirmed human carcinogen by IARC.

ACGIH: No component of this product present at levels greater than or equal to 0.1% is identified as a

carcinogen or potential carcinogen by ACGIH.

NTP: No component of this product present at levels greater than or equal to 0.1% is identified as a known or

anticipated carcinogen by NTP.

OSHA: No component of this product present at levels greater than or equal to 0.1% is identified as a

carcinogen or potential carcinogen by OSHA.

Reproductive toxicity

no data available

Specific target organ toxicity - single exposure (Globally Harmonized System)

May cause respiratory irritation.

Specific target organ toxicity - repeated exposure (Globally Harmonized System)

no data available

Aspiration hazard

May be fatal if swallowed and enters airways.

Potential health effects

Inhalation May be harmful if inhaled. May cause respiratory tract irritation.

Ingestion Aspiration hazard if swallowed - can enter lungs and cause damage. May be harmful if

swallowed.

Skin May be harmful if absorbed through skin. May cause skin irritation.

Eyes May cause eye irritation.

Signs and Symptoms of Exposure

Damage to the lungs., To the best of our knowledge, the chemical, physical, and toxicological properties have not been thoroughly investigated.

Additional Information

RTECS: DA8750000

12. ECOLOGICAL INFORMATION

Toxicity

Toxicity to fish LC50 - Oncorhynchus mykiss (rainbow trout) - 1.55 mg/l - 96.0 h

Aldrich - P52407 Page 4 of 6

Toxicity to daphnia and other aquatic

Immobilization EC50 - Daphnia magna (Water flea) - 2 mg/l - 24 h

Persistence and degradability

no data available

invertebrates.

Bioaccumulative potential

no data available

Mobility in soil

no data available

PBT and vPvB assessment

no data available

Other adverse effects

An environmental hazard cannot be excluded in the event of unprofessional handling or disposal.

Toxic to aquatic organisms, may cause long-term adverse effects in the aquatic environment.

Avoid release to the environment.

13. DISPOSAL CONSIDERATIONS

Product

This combustible material may be burned in a chemical incinerator equipped with an afterburner and scrubber. Observe all federal, state, and local environmental regulations. Contact a licensed professional waste disposal service to dispose of this material.

Contaminated packaging

Dispose of as unused product.

14. TRANSPORT INFORMATION

DOT (US)

UN-Number: 2364 Class: 3

Packing group: III

Proper shipping name: n-Propyl benzene

Marine pollutant: No

Poison Inhalation Hazard: No

IMDG

UN-Number: 2364 Class: 3

Packing group: III

EMS-No: F-E, S-D

Proper shipping name: PROPYLBENZENE

Marine pollutant: No

IATA

UN-Number: 2364 Class: 3

Packing group: III

Proper shipping name: n-Propylbenzene

15. REGULATORY INFORMATION

OSHA Hazards

Combustible Liquid

DSL Status

All components of this product are on the Canadian DSL list.

SARA 302 Components

SARA 302: No chemicals in this material are subject to the reporting requirements of SARA Title III, Section 302.

SARA 313 Components

SARA 313: This material does not contain any chemical components with known CAS numbers that exceed the threshold (De Minimis) reporting levels established by SARA Title III, Section 313.

SARA 311/312 Hazards

Fire Hazard

Massachusetts Right To Know Components

	CAS-No.	Revision Date
Propylbenzene	103-65-1	2007-03-01
Pennsylvania Right To Know Components		
	CAS-No.	Revision Date
Propylbenzene	103-65-1	2007-03-01
New Jersey Right To Know Components		
	CAS-No.	Revision Date
Propylbenzene	103-65-1	2007-03-01

California Prop. 65 Components

This product does not contain any chemicals known to State of California to cause cancer, birth defects, or any other reproductive harm.

16. OTHER INFORMATION

Further information

Copyright 2010 Sigma-Aldrich Co. License granted to make unlimited paper copies for internal use only. The above information is believed to be correct but does not purport to be all inclusive and shall be used only as a guide. The information in this document is based on the present state of our knowledge and is applicable to the product with regard to appropriate safety precautions. It does not represent any guarantee of the properties of the product. Sigma-Aldrich Co., shall not be held liable for any damage resulting from handling or from contact with the above product. See reverse side of invoice or packing slip for additional terms and conditions of sale.

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O-XYLENE ICSC: 0084











ortho-Xylene 1,2-Dimethylbenzene o-Xylol $C_6H_4(CH_3)_2/C_8H_{10}$ Molecular mass: 106.2

ICSC # 0084 CAS # 95-47-6 RTECS # <u>ZE2450000</u> UN # 1307

EC # 601-022-00-9 August 03, 2002 Validated



TYPES OF HAZARD/ EXPOSURE	ACUTE HAZ SYMPTO		PREVENTION		FIRST AID/ FIRE FIGHTING
FIRE	Flammable.		NO open flames, NO sparks, an smoking.	d NO	Powder, water spray, foam, carbon dioxide.
EXPLOSION	Above 32°C explosive mixtures may be formed		Above 32°C use a closed system ventilation, and explosion-proof electrical equipment. Prevent but of electrostatic charges (e.g., by grounding).	ild-up	In case of fire: keep drums, etc., cool by spraying with water.
EXPOSURE			STRICT HYGIENE! AVOID EXPOSURE OF (PREGNANT) WOMEN!)	
•INHALATION	Dizziness. Drowsiness. Nausea.	Headache.	Ventilation, local exhaust, or breathing protection.		Fresh air, rest. Refer for medical attention.
•SKIN	Dry skin. Redness.		Protective gloves.		Remove contaminated clothes. Rinse and then wash skin with water and soap.
•EYES	Redness. Pain.		Safety spectacles.		First rinse with plenty of water for several minutes (remove contact lenses if easily possible), then take to a doctor.
•INGESTION	Burning sensation. Abd (Further see Inhalation)		Do not eat, drink, or smoke duri work.	ng	Rinse mouth. Do NOT induce vomiting. Refer for medical attention.
SPILLAGE	E DISPOSAL		STORAGE	PA	CKAGING & LABELLING

SPILLAGE DISPOSAL	STORAGE	PACKAGING & LABELLING
		Note: C Xn symbol R: 10-20/21-38 S: 2-25 UN Hazard Class: 3 UN Packing Group: III
organic gases and vapours.)		

SEE IMPORTANT INFORMATION ON BACK

ICSC: 0084

Prepared in the context of cooperation between the International Programme on Chemical Safety & the Commission of the European Communities (C) IPCS CEC 1994. No modifications to the International version have been made except to add the OSHA PELs, NIOSH RELs and NIOSH IDLH values.

O-XYLENE ICSC: 0084

I	PHYSICAL STATE; APPEARANCE: COLOURLESS LIQUID, WITH CHARACTERISTIC ODOUR.	ROUTES OF EXPOSURE: The substance can be absorbed into the body by inhalation, through the skin and by ingestion.			
M P	PHYSICAL DANGERS: As a result of flow, agitation, etc., electrostatic charges can be generated.	INHALATION RISK: A harmful contamination of the air will be reached rather slowly on evaporation of this substance at 20°C.			
O R	CHEMICAL DANGERS: Reacts with strong acids strong oxidants	EFFECTS OF SHORT-TERM EXPOSURE: The substance is irritating to the eyes and the skin The substance may cause effects on the central nervous			
Т	OCCUPATIONAL EXPOSURE LIMITS: TLV: 100 ppm as TWA 150 ppm as STEL A4 (ACGIH	system If this liquid is swallowed, aspiration into the			
A	2001). BEI (ACGIH 2001). MAK: 100 ppm 440 mg/m³	EFFECTS OF LONG-TERM OR REPEATED			
N _	Peak limitation category: II(2) skin absorption (H); Pregnancy risk group: D	EXPOSURE: The liquid defats the skin. The substance may have effects on the central nervous system. Exposure to the			
T	(DFG 2005). EU OEL: 50 ppm as TWA 100 ppm as STEL	substance may enhance hearing damage caused by exposure to noise. Animal tests show that this substance			
D	(skin) (EU 2000).	possibly causes toxicity to human reproduction or development.			
A	OSHA PEL <u>†</u> : TWA 100 ppm (435 mg/m ³) NIOSH REL: TWA 100 ppm (435 mg/m ³) ST 150 ppm				
T A	(655 mg/m ³) NIOSH IDLH: 900 ppm See: <u>95476</u>				
A					
PHYSICAL PROPERTIES	Boiling point: 144°C Melting point: -25°C Relative density (water = 1): 0.88 Solubility in water: none Vapour pressure, kPa at 20°C: 0.7	Relative vapour density (air = 1): 3.7 Relative density of the vapour/air-mixture at 20°C (air = 1): 1.02 Flash point: 32°C c.c. Auto-ignition temperature: 463°C Explosive limits, vol% in air: 0.9-6.7 Octanol/water partition coefficient as log Pow: 3.12			
ENVIRONMENTAL DATA	The substance is toxic to aquatic organisms.				
	NOTES				

Depending on the degree of exposure, periodic medical examination is indicated. The recommendations on this Card also apply to technical xylene. See ICSC 0086 p-Xylene and 0085 m-Xylene.

Transport Emergency Card: TEC (R)-30S1307-III NFPA Code: H 2; F 3; R 0;

ADDITIONAL INFORMATION ICSC: 0084 (C) IPCS, CEC, 1994

IMPORTANT LEGAL NOTICE:

p-XYLENE ICSC: 0086











para-Xylene 1,4-Dimethylbenzene p-Xylol $C_6H_4(CH_3)_2/C_8H_{10}$ Molecular mass: 106.2

ICSC # 0086 CAS # 106-42-3 RTECS # <u>ZE2625000</u> UN # 1307

EC # 601-022-00-9 August 03, 2002 Validated



Above 27°C explosive vapour/air mixtures may be formed. Above 27°C use a closed system, ventilation, and explosion-proof electrical equipment. Prevent build-up of electrostatic charges (e.g., by grounding). STRICT HYGIENE! AVOID EXPOSURE OF (PREGNANT) WOMEN! -INHALATION Dizziness. Drowsiness. Headache. Nausea. Ventilation, local exhaust, or breathing protection. Fresh air, rest. Refer for medical attention. -SKIN Protective gloves. Remove contaminated clothes. R and then wash skin with water are soap. -EYES Redness. Pain. Safety spectacles. First rinse with plenty of water for several minutes (remove contact lenses if easily possible), then tail a doctor.	TYPES OF HAZARD/ EXPOSURE	ACUTE HAZ		PREVENTION	FIRST AID/ FIRE FIGHTING
EXPLOSION mixtures may be formed. ventilation, and explosion-proof electrical equipment. Prevent build-up of electrostatic charges (e.g., by grounding). by spraying with water. EXPOSURE STRICT HYGIENE! AVOID EXPOSURE OF (PREGNANT) WOMEN! Fresh air, rest. Refer for medical attention. •INHALATION Dizziness. Drowsiness. Headache. Nausea. Ventilation, local exhaust, or breathing protection. Fresh air, rest. Refer for medical attention. •SKIN Dry skin. Redness. Protective gloves. Remove contaminated clothes. R and then wash skin with water an soap. •EYES Redness. Pain. Safety spectacles. First rinse with plenty of water for several minutes (remove contact lenses if easily possible), then tal a doctor.	FIRE	Flammable.			
EXPOSURE EXPOSURE OF (PREGNANT) WOMEN! •INHALATION Dizziness. Drowsiness. Headache. Nausea. Ventilation, local exhaust, or breathing protection. Fresh air, rest. Refer for medical attention. •SKIN Dry skin. Redness. Protective gloves. Remove contaminated clothes. Rand then wash skin with water are soap. •EYES Redness. Pain. Safety spectacles. First rinse with plenty of water for several minutes (remove contact lenses if easily possible), then tall a doctor.	EXPLOSION			ventilation, and explosion-proof electrical equipment. Prevent build-u of electrostatic charges (e.g., by	
• Nausea. • Protective gloves. Remove contaminated clothes. Rand then wash skin with water an soap. • EYES Redness. Pain. Safety spectacles. First rinse with plenty of water for several minutes (remove contact lenses if easily possible), then tall a doctor. Rurning sensetion. Abdominal pain. Do not eat, drink, or smoke during. Rinse mouth. Do NOT induces.	EXPOSURE			EXPOSURE OF (PREGNANT)	
•SKIN Redness. Pain. Safety spectacles. First rinse with plenty of water for several minutes (remove contact lenses if easily possible), then tall a doctor. Burning sensetion Abdominal pain. Do not eat, drink, or smoke during. Rinse mouth, Do NOT induces.	•INHALATION		Headache.		Fresh air, rest. Refer for medical attention.
several minutes (remove contact lenses if easily possible), then tall a doctor. Rurning sensation Abdominal pain Do not eat drink or smoke during Rings mouth Do NOT induces.	•SKIN	Dry skin. Redness.		Protective gloves.	Remove contaminated clothes. Rinse and then wash skin with water and soap.
Burning sensation, Abdominal pain. Do not eat, drink, or smoke during Rinse mouth. Do NOT induce	•EYES	Redness. Pain.		Safety spectacles.	First rinse with plenty of water for several minutes (remove contact lenses if easily possible), then take to a doctor.
	•INGESTION			Do not eat, drink, or smoke during work.	Rinse mouth. Do NOT induce vomiting. Refer for medical attention.

SPILLAGE DISPOSAL	STORAGE	PACKAGING & LABELLING
		Note: C Xn symbol R: 10-20/21-38 S: 2-25 UN Hazard Class: 3 UN Packing Group: III

SEE IMPORTANT INFORMATION ON BACK

ICSC: 0086

Prepared in the context of cooperation between the International Programme on Chemical Safety & the Commission of the European Communities (C) IPCS CEC 1994. No modifications to the International version have been made except to add the OSHA PELs, NIOSH RELs and NIOSH IDLH values.

p-XYLENE ICSC: 0086

PILIBBIA					
I	PHYSICAL STATE; APPEARANCE: COLOURLESS LIQUID, WITH CHARACTERISTIC ODOUR.	ROUTES OF EXPOSURE: The substance can be absorbed into the body by inhalation, through the skin and by ingestion.			
M	ODOUR.	minatation, through the skin and by ingestion.			
P	PHYSICAL DANGERS: As a result of flow, agitation, etc., electrostatic charges can be generated.	INHALATION RISK: A harmful contamination of the air will be reached rather slowly on evaporation of this substance at 20°C.			
0					
R	CHEMICAL DANGERS: Reacts with strong acids strong oxidants	EFFECTS OF SHORT-TERM EXPOSURE: The substance is irritating to the eyes and the skin The substance may cause effects on the central nervous			
Т	OCCUPATIONAL EXPOSURE LIMITS: TLV: 100 ppm as TWA 150 ppm as STEL A4 (ACGIH	system If this liquid is swallowed, aspiration into the			
A	2001). BEI (ACGIH 2001). MAK: 100 ppm 440 mg/m ³	EFFECTS OF LONG-TERM OR REPEATED			
N	Peak limitation category: II(2)	EXPOSURE:			
Т	skin absorption (H); Pregnancy risk group: D (DFG 2005).	The liquid defats the skin. The substance may have effects on the central nervous system. Animal tests show that this substance possibly causes toxicity to human			
D	EU OEL: 50 ppm as TWA 100 ppm as STEL (skin) (EU 2000).				
A	OSHA PEL±: TWA 100 ppm (435 mg/m³) NIOSH REL: TWA 100 ppm (435 mg/m³) ST 150 ppm				
Т	(655 mg/m ³) NIOSH IDLH: 900 ppm See: <u>95476</u>				
A					
PHYSICAL PROPERTIES	Boiling point: 138°C Melting point: 13°C Relative density (water = 1): 0.86 Solubility in water: none Vapour pressure, kPa at 20°C: 0.9	Relative vapour density (air = 1): 3.7 Relative density of the vapour/air-mixture at 20°C (air = 1): 1.02 Flash point: 27°C c.c. Auto-ignition temperature: 528°C Explosive limits, vol% in air: 1.1-7.0 Octanol/water partition coefficient as log Pow: 3.15			
ENVIRONMENTAL DATA	The substance is toxic to aquatic organisms.				
NOTES					
Depending on the degree of exposure, periodic medical examination is indicated. The recommendations on this Card also apply to technical xylene. See ICSC 0084 o-Xylene and 0085 m-Xylene. Transport Emergency Card: TEC (R)-30S1307-III NFPA Code: H 2; F 3; R 0;					
ADDITIONAL INFORMATION					

ICSC: 0086 p-XYLENE

(C) IPCS, CEC, 1994

IMPORTANT LEGAL NOTICE:

m-XYLENE ICSC: 0085











meta-Xylene 1,3-Dimethylbenzene m-Xylol $C_6H_4(CH_3)_2/C_8H_{10}$ Molecular mass: 106.2

ICSC # 0085 CAS # 108-38-3 RTECS # <u>ZE2275000</u> UN # 1307

EC # 601-022-00-9 August 03, 2002 Validated



TYPES OF HAZARD/ EXPOSURE	ACUTE HAZAR SYMPTOMS		FIRST AID/ FIRE FIGHTING
FIRE	Flammable.	NO open flames, NO sparks, and smoking.	NO Powder, water spray, foam, carbon dioxide.
EXPLOSION	Above 27°C explosive vapo mixtures may be formed.	ur/air Above 27°C use a closed system, ventilation, and explosion-proof electrical equipment. Prevent buil of electrostatic charges (e.g., by grounding).	d-up In case of fire: keep drums, etc., cool by spraying with water.
EXPOSURE		STRICT HYGIENE!	
•INHALATION	Dizziness. Drowsiness. Head Nausea.	dache. Ventilation, local exhaust, or breathing protection.	Fresh air, rest. Refer for medical attention.
•SKIN	Dry skin. Redness.	Protective gloves.	Remove contaminated clothes. Rinse and then wash skin with water and soap.
•EYES	Redness. Pain.	Safety spectacles.	First rinse with plenty of water for several minutes (remove contact lenses if easily possible), then take to a doctor.
•INGESTION	Burning sensation. Abdomir (Further see Inhalation).	nal pain. Do not eat, drink, or smoke during work.	Rinse mouth. Do NOT induce vomiting. Refer for medical attention.
GDV - GD D GD - GD - GD - GD - GD - GD -			

SPILLAGE DISPOSAL	STORAGE	PACKAGING & LABELLING
		Note: C Xn symbol
remaining liquid in sand or inert absorbent and remove to safe place. Do NOT let this chemical enter the environment. (Extra personal protection: filter respirator for organic gases and vapours.)		R: 10-20/21-38 S: 2-25 UN Hazard Class: 3 UN Packing Group: III

SEE IMPORTANT INFORMATION ON BACK

ICSC: 0085

Prepared in the context of cooperation between the International Programme on Chemical Safety & the Commission of the European Communities (C) IPCS CEC 1994. No modifications to the International version have been made except to add the OSHA PELs, NIOSH RELs and NIOSH IDLH values.

m-XYLENE ICSC: 0085

[r					
I	PHYSICAL STATE; APPEARANCE: COLOURLESS LIQUID, WITH CHARACTERIS' ODOUR.	ROUTES OF EXPOSURE: TIC The substance can be absorbed into the body by inhalation, through the skin and by ingestion.			
M	ODOUK.	innaration, through the skin and by ingestion.			
P	PHYSICAL DANGERS: As a result of flow, agitation, etc., electrostatic charge can be generated.	ges A harmful contamination of the air will be reached rather slowly on evaporation of this substance at 20°C.			
0	eun de generateu.	rainer stowny on evaporation of this substance at 20°C.			
R	CHEMICAL DANGERS: Reacts with strong acids strong oxidants	EFFECTS OF SHORT-TERM EXPOSURE: The substance is irritating to the eyes and the skin The substance may cause effects on the central nervous			
Т	OCCUPATIONAL EXPOSURE LIMITS: TLV: 100 ppm as TWA 150 ppm as STEL A4 (ACC	system If this liquid is swallowed, aspiration into the			
A	2001). BEI (ACGIH 2001). MAK: 100 ppm 440 mg/m ³	EFFECTS OF LONG-TERM OR REPEATED			
N	Peak limitation category: II(2)	EXPOSURE:			
Т	skin absorption (H); Pregnancy risk group: D (DFG 2005).	The liquid defats the skin. The substance may have effects on the central nervous system Animal tests show that this substance possibly causes toxicity to human			
D	EU OEL: 50 ppm as TWA 100 ppm as STEL (skin) 2000).	(EU reproduction or development.			
A	OSHA PEL±: TWA 100 ppm (435 mg/m³) NIOSH REL: TWA 100 ppm (435 mg/m³) ST 150 j	ppm			
Т	(655 mg/m ³) NIOSH IDLH: 900 ppm See: <u>95476</u>				
A					
PHYSICAL PROPERTIES	Boiling point: 139°C Melting point: -48°C Relative density (water = 1): 0.86 Solubility in water: none Vapour pressure, kPa at 20°C: 0.8	Relative vapour density (air = 1): 3.7 Relative density of the vapour/air-mixture at 20°C (air = 1): 1.02 Flash point: 27°C c.c. Auto-ignition temperature: 527°C Explosive limits, vol% in air: 1.1-7.0 Octanol/water partition coefficient as log Pow: 3.20			
ENVIRONMENTAL DATA	The substance is toxic to aquatic organisms.				
NOTES					
Depending on the degree of exposure, periodic medical examination is indicated. The recommendations on this Card also apply to technical xylene. See ICSC 0084 o-Xylene and 0086 p-Xylene. NFPA Code: H 2; F 3; R 0; Transport Emergency Card: TEC (R)-30S1307-III					
ADDITIONAL INFORMATION					
iL					

ICSC: 0085 m-XYLENE

(C) IPCS, CEC, 1994

IMPORTANT LEGAL NOTICE:

TOLUENE ICSC: 0078











 $\begin{array}{c} \text{Methylbenzene} \\ \text{Toluol} \\ \text{Phenylmethane} \\ \text{C}_6\text{H}_5\text{CH}_3\,/\,\text{C}_7\text{H}_8 \end{array}$

Molecular mass: 92.1

ICSC # 0078 CAS # 108-88-3 RTECS # <u>XS5250000</u>

UN # 1294

EC# 601-021-00-3

October 10, 2002 Peer reviewed



TYPES OF HAZARD/ EXPOSURE	ACUTE HAZ SYMPTO		PREVENTION		FIRST AID/ FIRE FIGHTING
FIRE	Highly flammable.		NO open flames, NO sparks, ar smoking.	nd NO	Powder, AFFF, foam, carbon dioxide.
EXPLOSION	Vapour/air mixtures are	explosive. Closed system, ventilation, explosion-		In case of fire: keep drums, etc., cool by spraying with water.	
EXPOSURE			STRICT HYGIENE! AVOID EXPOSURE OF (PREGNANT WOMEN!	·)	
•INHALATION	Cough. Sore throat. Diz Drowsiness. Headache. Unconsciousness.		Ventilation, local exhaust, or breathing protection.		Fresh air, rest. Refer for medical attention.
•SKIN	Dry skin. Redness.		Protective gloves.		Remove contaminated clothes. Rinse and then wash skin with water and soap. Refer for medical attention.
•EYES	Redness. Pain.	Safety goggles.		First rinse with plenty of water for several minutes (remove contact lenses if easily possible), then take to a doctor.	
•INGESTION	Burning sensation. Abd (Further see Inhalation)		1 1 / /		Rinse mouth. Do NOT induce vomiting. Refer for medical attention.
SPILLAGE DISPOSAL		STORAGE	PA	CKAGING & LABELLING	
Evacuate danger area in large spill! Consult an expert in large spill! Remove all ignition sources. Ventilation. Collect leaking liquid in sealable containers. Absorb remaining liquid in sand or inert absorbent and remove to safe place. Do NOT wash away into sewer. Do NOT let this chemical enter the environment. Personal protection: self-contained breathing apparatus		Fireproof. Sep	parated from strong oxidants.	S: 2-36 UN Ha	

SEE IMPORTANT INFORMATION ON BACK

ICSC: 0078

Prepared in the context of cooperation between the International Programme on Chemical Safety & the Commission of the European Communities (C) IPCS CEC 1994. No modifications to the International version have been made except to add the OSHA PELs, NIOSH RELs and NIOSH IDLH values.

International Chemical Safety Cards

TOLUENE ICSC: 0078

I	PHYSICAL STATE; APPEARANCE: COLOURLESS LIQUID, WITH CHARACTERISTIC	ROUTES OF EXPOSURE: The substance can be absorbed into the body by
M	ODOUR.	inhalation, through the skin and by ingestion.
P	PHYSICAL DANGERS: The vapour mixes well with air, explosive mixtures are	INHALATION RISK: A harmful contamination of the air can be reached rather
О	formed easily. As a result of flow, agitation, etc., electrostatic charges can be generated.	quickly on evaporation of this substance at 20°C.
R	CHEMICAL DANGERS:	EFFECTS OF SHORT-TERM EXPOSURE: The substance is irritating to the eyes and the respiratory
Т	Reacts violently with strong oxidants causing fire and explosion hazard.	tract The substance may cause effects on the central nervous system If this liquid is swallowed, aspiration
A	OCCUPATIONAL EXPOSURE LIMITS:	into the lungs may result in chemical pneumonitis. Exposure at high levels may result in cardiac
N	TLV: 50 ppm as TWA (skin) A4 BEI issued (ACGIH 2004).	dysrhythmiaandunconsciousness.
Т	MAK: 50 ppm 190 mg/m³ H Peak limitation category: II(4) Pregnancy risk group: C (DFG 2004).	EFFECTS OF LONG-TERM OR REPEATED EXPOSURE: The liquid defats the skin. The substance may have
D	OSHA PEL±: TWA 200 ppm C 300 ppm 500 ppm (10-minute maximum peak)	effects on the central nervous system Exposure to the substance may enhance hearing damage caused by
A	NIOSH REL: TWA 100 ppm (375 mg/m ³) ST 150 ppm (560 mg/m ³)	exposure to noise. Animal tests show that this substance possibly causes toxicity to human reproduction or
Т	NIOSH IDLH: 500 ppm See: <u>108883</u>	development.
A		
PHYSICAL	Boiling point: 111°C Melting point: -95°C Relative density (water = 1): 0.87	Relative density of the vapour/air-mixture at 20°C (air = 1): 1.01 Flash point: 4°C c.c.
PROPERTIES	Solubility in water: none Vapour pressure, kPa at 25°C: 3.8	Auto-ignition temperature: 480°C Explosive limits, vol% in air: 1.1-7.1
	Relative vapour density (air = 1): 3.1	Octanol/water partition coefficient as log Pow: 2.69
ENVIRONMENTAL DATA	The substance is toxic to aquatic organisms.	

NOTES

Depending on the degree of exposure, periodic medical examination is suggested. Use of alcoholic beverages enhances the harmful effect.

Transport Emergency Card: TEC (R)-30S1294

NFPA Code: H 2; F 3; R 0;

ADDITIONAL INFORMATION

ICSC: 0078 TOLUENE

(C) IPCS, CEC, 1994

IMPORTANT LEGAL NOTICE:

NAPHTHALENE ICSC: 0667













Molecular mass: 128.18

ICSC # 0667 CAS # 91-20-3 RTECS # QJ0525000

UN # 1334 (solid); 2304 (molten)

EC # 601-052-00-2 April 21, 2005 Peer reviewed



TYPES OF HAZARD/ EXPOSURE	ACUTE HAZARDS/ SYMPTOMS	PREVENTION	FIRST AID/ FIRE FIGHTING
FIRE	Combustible.	NO open flames.	Powder, water spray, foam, carbon dioxide.
EXPLOSION	Above 80°C explosive vapour/air mixtures may be formed. Finely dispersed particles form explosive mixtures in air.	Prevent deposition of dust; closed system, dust explosion-proof electrical equipment and lighting.	
EXPOSURE		PREVENT DISPERSION OF DUST!	
•INHALATION	Headache. Weakness. Nausea. Vomiting. Sweating. Confusion. Jaundice. Dark urine.	Ventilation (not if powder), local exhaust, or breathing protection.	Fresh air, rest. Refer for medical attention.
•SKIN	MAY BE ABSORBED! (Further see Inhalation).	Protective gloves.	Rinse skin with plenty of water or shower.
•EYES		Safety spectacles.	First rinse with plenty of water for several minutes (remove contact lenses if easily possible), then take to a doctor.
•INGESTION	Abdominal pain. Diarrhoea. Convulsions. Unconsciousness. (Further see Inhalation).	Do not eat, drink, or smoke during work. Wash hands before eating.	Rest. Refer for medical attention.
SDILLAGE DISDOSAL STODAGE DACKACING & LADELLING			

SPILLAGE DISPUSAL	STURAGE	PACKAGING & LABELLING
Personal protection: filter respirator for	Separated from strong oxidants, food and	Do not transport with food and feedstuffs.
organic gases and vapours. Do NOT let this	feedstuffs Store in an area without drain or	Marine pollutant.
chemical enter the environment. Sweep	sewer access.	Xn symbol
spilled substance into covered containers; if		N symbol
appropriate, moisten first to prevent dusting.		R: 22-40-50/53
Carefully collect remainder, then remove to		S: 2-36/37-46-60-61
safe place.		UN Hazard Class: 4.1
		UN Packing Group: III

SEE IMPORTANT INFORMATION ON BACK

ICSC: 0667

Prepared in the context of cooperation between the International Programme on Chemical Safety & the Commission of the European Communities (C) IPCS CEC 1994. No modifications to the International version have been made except to add the OSHA PELs, NIOSH RELs and NIOSH IDLH values.

NAPHTHALENE ICSC: 0667

I M	PHYSICAL STATE; APPEARANCE: WHITE SOLID IN VARIOUS FORMS, WITH CHARACTERISTIC ODOUR.	ROUTES OF EXPOSURE: The substance can be absorbed into the body by inhalation, through the skin and by ingestion.				
P	PHYSICAL DANGERS: Dust explosion possible if in powder or granular form,	INHALATION RISK: A harmful contamination of the air will be reached				
О	mixed with air.	rather slowly on evaporation of this substance at 20°C. See Notes.				
R	CHEMICAL DANGERS: On combustion, forms irritating and toxic gases. Reacts	EFFECTS OF SHORT-TERM EXPOSURE:				
T	with strong oxidants	The substance may cause effects on the blood, resulting in lesions of blood cells (haemolysis) See Notes. The				
A	OCCUPATIONAL EXPOSURE LIMITS: TLV: 10 ppm as TWA 15 ppm as STEL (skin) A4 (not	effects may be delayed. Exposure by ingestion may result in death. Medical observation is indicated.				
N	classifiable as a human carcinogen); (ACGIH 2005).	EFFECTS OF LONG-TERM OR REPEATED				
Т	MAK: skin absorption (H); Carcinogen category: 2; Germ cell mutagen group: 3B; (DFG 2004).	EXPOSURE: The substance may have effects on the blood, resulting				
D	OSHA PEL±: TWA 10 ppm (50 mg/m³) NIOSH REL: TWA 10 ppm (50 mg/m³) ST 15 ppm (75	in chronic haemolytic anaemia. The substance may have effects on the eyes, resulting in the development of				
A	mg/m ³) NIOSH IDLH: 250 ppm See: <u>91203</u>	cataract. This substance is possibly carcinogenic to humans.				
T						
A						
PHYSICAL PROPERTIES	Boiling point: 218°C Sublimation slowly at room temperature Melting point: 80°C Density: 1.16 g/cm3 Solubility in water, g/100 ml at 25°C: none	Vapour pressure, Pa at 25°C: 11 Relative vapour density (air = 1): 4.42 Flash point: 80°C c.c. Auto-ignition temperature: 540°C Explosive limits, vol% in air: 0.9-5.9 Octanol/water partition coefficient as log Pow: 3.3				
ENVIRONMENTAL DATA	Hoguetic anymonment					
	NOTES					
Some individuals may be more sensitive to the effect of naphthalene on blood cells.						

Some individuals may be more sensitive to the effect of naphthalene on blood cells.

Transport Emergency Card: TEC (R)-41S1334 (solid); 41GF1-II+III (solid); 41S2304 (molten)

NFPA Code: H2; F2; R0;

ADDITIONAL INFORMATION

ICSC: 0667 NAPHTHALENE

(C) IPCS, CEC, 1994

IMPORTANT LEGAL NOTICE:

BARIUM SULFATE











ICSC: 0827

Barium sulphate Blanc fixe Artificial barite BaSO₄

Molecular mass: 233.43

ICSC # 0827 CAS # 7727-43-7 RTECS # <u>CR0600000</u>

October 20, 1999 Peer reviewed

TYPES OF HAZARD/	ACUTE HAZ		PREVENTION		FIRST AID/
EXPOSURE	SYMPTO	MS	INLVENTOR		FIRE FIGHTING
FIRE	Not combustible. Give irritating or toxic fume in a fire.				In case of fire in the surroundings: use appropriate extinguishing media.
EXPLOSION					
EXPOSURE			PREVENT DISPERSION OF DUST!	7	
•INHALATION			Local exhaust or breathing protection.		Fresh air, rest.
•SKIN			Protective gloves.		Remove contaminated clothes. Rinse skin with plenty of water or shower.
•EYES			Safety spectacles.		First rinse with plenty of water for several minutes (remove contact lenses if easily possible), then take to a doctor.
•INGESTION			Do not eat, drink, or smoke during work.		Rinse mouth.
SPILLAGE DISPOSAL			STORAGE	PAC	CKAGING & LABELLING
Sweep spilled substa appropriate, moisten dusting. Personal pro- respirator for inert p	otection: P1 filter			R: S:	

SEE IMPORTANT INFORMATION ON BACK

ICSC: 0827

Prepared in the context of cooperation between the International Programme on Chemical Safety & the Commission of the European Communities (C) IPCS CEC 1994. No modifications to the International version have been made except to add the OSHA PELs, NIOSH RELs and NIOSH IDLH values.

ICSC: 0827

BARIUM SULFATE

BARIUM SULFATE

I	PHYSICAL STATE; APPEARANCE: ODOURLESS TASTELESS, WHITE OR	ROUTES OF EXPOSURE: The substance can be absorbed into the body by				
M	YELLOWISH CRYSTALS OR POWDER.	inhalation of its aerosol.				
P	PHYSICAL DANGERS:	INHALATION RISK:				
О	CHEMICAL DANCEDC.	Evaporation at 20°C is negligible; a nuisance- causing concentration of airborne particles can,				
R	CHEMICAL DANGERS: Reacts violently with aluminium powder.	however, be reached quickly.				
Т	OCCUPATIONAL EXPOSURE LIMITS:	EFFECTS OF SHORT-TERM EXPOSURE:				
A	TLV: 10 mg/m³ as TWA; (ACGIH 2004). MAK: (Inhalable fraction) 4 mg/m³; (Respirable fraction) 1.5 mg/m³; (DFG 2004).	EFFECTS OF LONG-TERM OR REPEATED EXPOSURE:				
N	OSHA PEL±: TWA 15 mg/m³ (total) TWA 5	Lungs may be affected by repeated or prolonged exposure to dust particles, resulting in baritosis (a				
Т	mg/m³ (resp) NIOSH REL: TWA 10 mg/m³ (total) TWA 5	form of benign pneumoconiosis).				
D	mg/m ³ (resp) NIOSH IDLH: N.D. See: <u>IDLH INDEX</u>					
A						
Т						
A						
PHYSICAL PROPERTIES	Melting point (decomposes): 1600°C Density: 4.5 g/cm ³	Solubility in water: none				
ENVIRONMENTAL DATA						
	NOTES					
Occurs in nature as the mineral barite; also as barytes, heavy spar. Card has been partly updated in October 2005. See section Occupational Exposure Limits.						
	ADDITIONAL INFORM	ATION				

IMPORTANT LEGAL NOTICE:

ICSC: 0827

Neither NIOSH, the CEC or the IPCS nor any person acting on behalf of NIOSH, the CEC or the IPCS is responsible for the use which might be made of this information. This card contains the collective views of the IPCS Peer Review Committee and may not reflect in all cases all the detailed requirements included in national legislation on the subject. The user should verify compliance of the cards with the relevant legislation in the country of use. The only modifications made to produce the U.S. version is inclusion of the OSHA PELs, NIOSH RELs and NIOSH IDLH values.

(C) IPCS, CEC, 1994

CHROMIUM ICSC: 0029











Chrome Cr Atomic mass: 52.0 (powder)

ICSC # 0029 CAS # 7440-47-3 RTECS # <u>GB4200000</u>

ICSC: 0029

P

October 27, 2004 Peer reviewed

		PREVENTION		FIRST AID/ FIRE FIGHTING
Combustible under specif	fic conditions.	ic conditions. No open flames if in powder form.		In case of fire in the surroundings: use appropriate extinguishing media.
		Prevent deposition of dust; closed system, dust explosion-proof electrical equipment and lighting.		
		PREVENT DISPERSION OF I	DUST!	
Cough.	Local exhaust or brea		ection.	Fresh air, rest.
		Protective gloves.		Remove contaminated clothes. Rinse skin with plenty of water or shower.
Redness.		Safety goggles.		First rinse with plenty of water for several minutes (remove contact lenses if easily possible), then take to a doctor.
	Do not eat, drink, or smoke during work.		Rinse mouth.	
SPILLAGE DISPOSAL		STORAGE	PA	ACKAGING & LABELLING
Sweep spilled substance into containers; if appropriate, moisten first to prevent dusting. Personal protection: P2 filter respirator for harmful particles.			R: S:	
SEE IMPORTANT INFORMATION ON BACK				
	Cough. Cough.	Cough. Redness. CDISPOSAL The into containers; if the into containers; if the into containers in the into conta	Combustible under specific conditions. No open flames if in powder for Prevent deposition of dust; clos system, dust explosion-proof elequipment and lighting. PREVENT DISPERSION OF I Local exhaust or breathing prot Protective gloves. Redness. Safety goggles. Do not eat, drink, or smoke dur work. CDISPOSAL STORAGE et into containers; if rest to prevent dusting. efilter respirator for	Combustible under specific conditions. No open flames if in powder form. Prevent deposition of dust; closed system, dust explosion-proof electrical equipment and lighting. PREVENT DISPERSION OF DUST! Cough. Local exhaust or breathing protection. Protective gloves. Redness. Safety goggles. Do not eat, drink, or smoke during work. CDISPOSAL e into containers; if set to prevent dusting. efilter respirator for Reinter respirator for SYMPTOMS No open flames if in powder form. Protective gloseit; closed system, dust explosion of dust; closed system, dust explosion-proof electrical equipment and lighting. Protective gloves. Safety goggles.

International Chemical Safety Cards

CHROMIUM ICSC: 0029

т	PHYSICAL STATE; APPEARANCE:	ROUTES OF EXPOSURE:
1	GREY POWDER	

NIOSH RELs and NIOSH IDLH values.

M PHYSICAL DANGERS:

Dust explosion possible if in powder or granular form, mixed with air.

INHALATION RISK:

Prepared in the context of cooperation between the International Programme on Chemical Safety & the Commission of the European

Communities (C) IPCS CEC 1994. No modifications to the International version have been made except to add the OSHA PELs,

A harmful concentration of airborne particles can be reached quickly when dispersed.

o							
R	CHEMICAL DANGERS: Chromium is a catalytic substance and may cause reaction	EFFECTS OF SHORT-TERM EXPOSURE: May cause mechanical irritation to the eyesandthe					
Т	in contact with many organic and inorganic substances, causing fire and explosion hazard.	respiratory tract. EFFECTS OF LONG-TERM OR REPEATED					
A	OCCUPATIONAL EXPOSURE LIMITS:	EXPOSURE:					
N	TLV: (as Cr metal, Cr(III) compounds) 0.5 mg/m³ as TWA A4 (ACGIH 2004). MAK not established.						
Т	OSHA PEL*: TWA 1 mg/m ³ See Appendix C *Note: The PEL also applies to insoluble chromium salts.						
D	NIOSH REL: TWA 0.5 mg/m ³ See Appendix C NIOSH IDLH: 250 mg/m ³ (as Cr) See: 7440473						
A							
Т							
A							
PHYSICAL PROPERTIES	Boiling point: 2642°C Melting point: 1900°C Density: 7.15 g/cm ³	Solubility in water: none					
ENVIRONMENTAL DATA							
	NOTES						
The surface of the chro	mium particles is oxidized to chromium(III)oxide in air. See	ICSC 1531 Chromium(III) oxide.					
ADDITIONAL INFORMATION							
ICSC: 0029		CHROMIUM					
(C) IPCS, CEC, 1994							

IMPORTANT LEGAL NOTICE:

COPPER ICSC: 0240











Cu (powder)

ICSC # 0240 CAS # 7440-50-8 RTECS # <u>GL5325000</u>

September 24, 1993 Validated

TYPES OF HAZARD/ EXPOSURE	ACUTE HAZ SYMPTO		PREVENTION		FIRST AID/ FIRE FIGHTING
FIRE	Combustible.		NO open flames.		Special powder, dry sand, NO other agents.
EXPLOSION					
EXPOSURE			PREVENT DISPERSION OF D	UST!	
•INHALATION	Cough. Headache. Short Sore throat.	ness of breath.	Local exhaust or breathing protect	ction.	Fresh air, rest. Refer for medical attention.
•SKIN	Redness.		Protective gloves.		Remove contaminated clothes. Rinse and then wash skin with water and soap.
•EYES	Redness. Pain.		Safety goggles.		First rinse with plenty of water for several minutes (remove contact lenses if easily possible), then take to a doctor.
•INGESTION	Abdominal pain. Nausea. Vomiting.		Do not eat, drink, or smoke during work.		Rinse mouth. Refer for medical attention.
SPILLAGE DISPOSAL		STORAGE	PA	ACKAGING & LABELLING	
Sweep spilled substance into containers. Carefully collect remainder. Then remove to safe place. (Extra personal protection: P2 filter respirator for harmful particles).		Separated from	m - See Chemical Dangers. R: S:		
	S	EE IMPORT <i>A</i>	ANT INFORMATION ON BAC	K	
ICSC: 0240	Com	ared in the context of munities (C) IPCS C	EC 1994. No modifications to the Internation	amme on nal versior	Chemical Safety & the Commission of the European have been made except to add the OSHA PELs,

International Chemical Safety Cards

NIOSH RELs and NIOSH IDLH values.

COPPER ICSC: 0240

ī	PHYSICAL STATE; APPEARANCE: RED POWDER, TURNS GREEN ON EXPOSURE TO MOIST AIR.	ROUTES OF EXPOSURE: The substance can be absorbed into the body by inhalation and by ingestion.
M	PHYSICAL DANGERS:	INHALATION RISK: Evaporation at 20°C is negligible; a harmful concentration
P	CHEMICAL DANGERS:	of airborne particles can, however, be reached quickly when dispersed.

1		
0	Shock-sensitive compounds are formed with acetylenic	
	compounds, ethylene oxides and azides. Reacts with strong	EFFECTS OF SHORT-TERM EXPOSURE:
R	oxidants like chlorates, bromates and iodates, causing	Inhalation of fumes may cause metal fume fever. See
_	explosion hazard.	Notes.
T		
	OCCUPATIONAL EXPOSURE LIMITS:	EFFECTS OF LONG-TERM OR REPEATED
A	TLV: 0.2 mg/m ³ fume (ACGIH 1992-1993).	EXPOSURE:
N	TLV (as Cu, dusts & mists): 1 mg/m ³ (ACGIH 1992-1993).	
14	Intended change 0.1 mg/m³ Inhal.,	sensitization.
T	A4 (not classifiable as a human carcinogen);	
-	MAK: 0.1 mg/m³ (Inhalable fraction)	
	Peak limitation category: II(2) Pregnancy risk group: D	
D	(DFG 2005).	
	OSHA PEL*: TWA 1 mg/m ³ *Note: The PEL also applies	
A	to other copper compounds (as Cu) except copper fume.	
	NIOSH REL*: TWA 1 mg/m ³ *Note: The REL also	
T	applies to other copper compounds (as Cu) except Copper	
	fume.	
A	NIOSH IDLH: 100 mg/m ³ (as Cu) See: 7440508	
	(43 Cu) Sec. 7440500	
	Boiling point: 2595°C	Solubility in water:
PHYSICAL	Melting point: 1083°C	none
PROPERTIES	Relative density (water = 1): 8.9	
ENVIRONMENTAL		
DATA		
	NOTES	
The symptoms of metal	I fume fever do not become manifest until several hours.	
	ADDITIONAL INFORMA	ΓΙΟΝ
ICSC: 0240		COPPER

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LEAD ICSC: 0052











Lead metal
Plumbum
Pb
Atomic mass: 207.2
(powder)

ICSC # 0052 CAS # 7439-92-1 RTECS # <u>OF7525000</u>

October 08, 2002 Peer reviewed

TYPES OF HAZARD/ EXPOSURE	ACUTE HAZ SYMPTO		PREVENTION		FIRST AID/ FIRE FIGHTING
FIRE	Not combustible. Gives or toxic fumes (or gases				In case of fire in the surroundings: use appropriate extinguishing media.
EXPLOSION	Finely dispersed particle explosive mixtures in ai		Prevent deposition of dust; close system, dust explosion-proof electrical equipment and lightin		
EXPOSURE	See EFFECTS OF LON REPEATED EXPOSUR		PREVENT DISPERSION OF I AVOID EXPOSURE OF (PREGNANT) WOMEN!	OUST!	
•INHALATION			Local exhaust or breathing proto	ection.	Fresh air, rest.
•SKIN			Protective gloves.		Remove contaminated clothes. Rinse and then wash skin with water and soap.
•EYES			Safety spectacles.		First rinse with plenty of water for several minutes (remove contact lenses if easily possible), then take to a doctor.
•INGESTION	Abdominal pain. Nause	a. Vomiting.			Rinse mouth. Give plenty of water to drink. Refer for medical attention.
SPILLAGE DISPOSAL		STORAGE	PA	CKAGING & LABELLING	
Sweep spilled substance into containers; if Separated from		n food and feedstuffs			

SPILLAGE DISPOSAL	STORAGE	PACKAGING & LABELLING
appropriate, moisten first to prevent dusting.	Domana	R: S:

SEE IMPORTANT INFORMATION ON BACK

ICSC: 0052

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International Chemical Safety Cards

LEAD ICSC: 0052

PHYSICAL STATE; APPEARANCE: **ROUTES OF EXPOSURE:** BLUISH-WHITE OR SILVERY-GREY SOLID IN The substance can be absorbed into the body by VARIOUS FORMS, TURNS TARNISHED ON inhalation and by ingestion. EXPOSURE TO AIR. Ι INHALATION RISK: PHYSICAL DANGERS: A harmful concentration of airborne particles can be M Dust explosion possible if in powder or granular form, reached quickly when dispersed, especially if powdered. mixed with air. P EFFECTS OF SHORT-TERM EXPOSURE: CHEMICAL DANGERS: O On heating, toxic fumes are formed. Reacts with oxidants. Reacts with hot concentrated nitric acid, EFFECTS OF LONG-TERM OR REPEATED R boiling concentrated hydrochloric acid and sulfuric acid. **EXPOSURE:** Attacked by pure water and by weak organic acids in the The substance may have effects on the blood bone T presence of oxygen. marrow central nervous system peripheral nervous system kidneys, resulting in anaemia, encephalopathy OCCUPATIONAL EXPOSURE LIMITS: (e.g., convulsions), peripheral nerve disease, abdominal TLV: 0.05 mg/m³ A3 (confirmed animal carcinogen cramps and kidney impairment. Causes toxicity to N with unknown relevance to humans); BEI issued human reproduction or development. (ACGIH 2004). \mathbf{T} MAK: Carcinogen category: 3B; Germ cell mutagen group: 3A; (DFG 2004). D EU OEL: as TWA 0.15 mg/m³ (EU 2002). OSHA PEL*: 1910.1025 TWA 0.050 mg/m³ See Appendix C *Note: The PEL also applies to other lead compounds (as Pb) -- see Appendix C. Т NIOSH REL*: TWA 0.050 mg/m³ See Appendix C *Note: The REL also applies to other lead compounds A (as Pb) -- see Appendix C. NIOSH IDLH: 100 mg/m³ (as Pb) See: 7439921 Boiling point: 1740°C Density: 11.34 g/cm3 **PHYSICAL** Solubility in water: none **PROPERTIES** Melting point: 327.5°C Bioaccumulation of this chemical may occur in plants and in mammals. It is strongly advised that this ENVIRONMENTAL substance does not enter the environment. DATA NOTES Depending on the degree of exposure, periodic medical examination is suggested. Do NOT take working clothes home. Transport Emergency Card: TEC (R)-51S1872 ADDITIONAL INFORMATION

IMPORTANT LEGAL NOTICE:

ICSC: 0052

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LEAD

ICSC: 0056 **MERCURY**











Quicksilver Liquid silver Hg Atomic mass: 200.6

ICSC# 0056

CAS# 7439-97-6 RTECS # OV4550000

UN# 2809

EC# 080-001-00-0 April 22, 2004 Peer reviewed









TYPES OF HAZARD/ EXPOSURE	ACUTE HAZA SYMPTON		PREVENTION		FIRST AID/ FIRE FIGHTING
FIRE	Not combustible. Gives of toxic fumes (or gases) in				In case of fire in the surroundings: use appropriate extinguishing media.
EXPLOSION	Risk of fire and explosion	1.			In case of fire: keep drums, etc., cool by spraying with water.
EXPOSURE			STRICT HYGIENE! AVOID EXPOSURE OF (PREGNANT) WOMEN! AVOID EXPOSURE ADOLESCENTS AND CHILDI	OF	IN ALL CASES CONSULT A DOCTOR!
•INHALATION	Abdominal pain. Cough. Shortness of breath. Vom or elevated body tempera	iting. Fever	Local exhaust or breathing protection	ction.	Fresh air, rest. Artificial respiration if indicated. Refer for medical attention.
•SKIN	MAY BE ABSORBED!	Redness.	Protective gloves. Protective clos	thing.	Remove contaminated clothes. Rinse and then wash skin with water and soap. Refer for medical attention.
•EYES			Face shield, or eye protection in combination with breathing prote	ection.	First rinse with plenty of water for several minutes (remove contact lenses if easily possible), then take to a doctor.
•INGESTION			Do not eat, drink, or smoke durin work. Wash hands before eating.		Refer for medical attention.
CDILLACI	EDICDOCAT	_	STODACE	DA	CKACING & LADELLING

SPILLAGE DISPOSAL **STORAGE** PACKAGING & LABELLING Evacuate danger area in case of a large spill! Provision to contain effluent from fire Special material. Do not transport with food Consult an expert! Ventilation. Collect leaking extinguishing. Separated from food and and feedstuffs. and spilled liquid in sealable non-metallic feedstuffs Well closed. T symbol containers as far as possible. Do NOT wash N symbol away into sewer. Do NOT let this chemical R: 23-33-50/53 enter the environment. Chemical protection S: 1/2-7-45-60-61 suit including self-contained breathing UN Hazard Class: 8 apparatus. UN Packing Group: III

SEE IMPORTANT INFORMATION ON BACK

Prepared in the context of cooperation between the International Programme on Chemical Safety & the Commission of the ICSC: 0056 European Communities (C) IPCS CEC 1994. No modifications to the International version have been made except to add the OSHA PELs, NIOSH RELs and NIOSH IDLH values

MERCURY ICSC: 0056

I	PHYSICAL STATE; APPEARANCE: ODOURLESS, HEAVY AND MOBILE SILVERY	ROUTES OF EXPOSURE: The substance can be absorbed into the body by inhalation
M	LIQUID METAL.	of its vapour and through the skin, also as a vapour!
P	PHYSICAL DANGERS:	INHALATION RISK:
О	CHEMICAL DANCEDS.	A harmful contamination of the air can be reached very quickly on evaporation of this substance at 20°C.
R	CHEMICAL DANGERS: Upon heating, toxic fumes are formed. Reacts violently	EFFECTS OF SHORT-TERM EXPOSURE:
	with ammonia and halogens causing fire and explosion	The substance is irritating to the skin. Inhalation of the
T	hazard. Attacks aluminium and many other metals	vapours may cause pneumonitis. The substance may cause
\mathbf{A}	forming amalgams.	effects on the central nervous systemandkidneys. The effects may be delayed. Medical observation is indicated.
	OCCUPATIONAL EXPOSURE LIMITS:	erroris may ob domy our risourour cosor random is marcarour
N	TLV: 0.025 mg/m ³ as TWA (skin) A4 BEI issued	EFFECTS OF LONG-TERM OR REPEATED
T	(ACGIH 2004). MAK: 0.1 mg/m³ Sh	EXPOSURE: The substance may have effects on the central nervous
•	Peak limitation category: II(8) Carcinogen category: 3B	system kidneys, resulting in irritability, emotional
	(DFG 2003).	instability, tremor, mental and memory disturbances,
D	OSHA PEL±: C 0.1 mg/m ³	speech disorders. Danger of cumulative effects. Animal
A	NIOSH REL: Hg Vapor: TWA 0.05 mg/m ³ skin	tests show that this substance possibly causes toxic effects upon human reproduction.
11	Other: C 0.1 mg/m ³ skin	upon numan reproduction.
T	NIOSH IDLH: 10 mg/m ³ (as Hg) See: <u>7439976</u>	
A		
	Boiling point: 357°C	Vapour pressure, Pa at 20°C: 0.26
PHYSICAL	Melting point: -39°C	Relative vapour density (air = 1): 6.93
PROPERTIES	Relative density (water = 1): 13.5 Solubility in water:	Relative density of the vapour/air-mixture at 20°C (air = 1): 1.009
	none	1). 1.007
ENVIRONMENTAL DATA	The substance is very toxic to aquatic organisms. In the fortakes place, specifically in fish.	od chain important to humans, bioaccumulation
	NOTES	
Depending on the degree NOT take working clot	ee of exposure, periodic medical examination is indicated. I	No odour warning if toxic concentrations are present. Do
THO I Take WOLKING CIOU	nes nome.	Transport Emergency Card: TEC (R)-80GC9-II+III
	ADDITIONAL INFORMA	ATION
ICSC: 0056		MERCURY

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NICKEL ICSC: 0062











Ni Atomic mass: 58.7 (powder)

ICSC # 0062 CAS # 7440-02-0 RTECS # <u>QR5950000</u> EC # 028-002-00-7

October 17, 2001 Peer reviewed

TYPES OF HAZARD/ EXPOSURE	ACUTE HAZARDS/ SYMPTOMS	PREVENTION	FIRST AID/ FIRE FIGHTING
FIRE	Flammable as dust. Toxic fumes may be released in a fire.		Dry sand. NO carbon dioxide. NO water.
EXPLOSION	Finely dispersed particles form explosive mixtures in air.	Prevent deposition of dust; closed system, dust explosion-proof electrical equipment and lighting.	
EXPOSURE		PREVENT DISPERSION OF DUST! AVOID ALL CONTACT!	
•INHALATION	Cough. Shortness of breath.	Local exhaust or breathing protection.	Fresh air, rest.
•SKIN		Protective gloves. Protective clothing.	Remove contaminated clothes. Rinse and then wash skin with water and soap.
•EYES		Safety spectacles, or eye protection in combination with breathing protection.	First rinse with plenty of water for several minutes (remove contact lenses if easily possible), then take to a doctor.
•INGESTION		Do not eat, drink, or smoke during work.	Rinse mouth.

SPILLAGE DISPOSAL	STORAGE	PACKAGING & LABELLING
Vacuum spilled material. Carefully collect	Separated from strong acids.	
remainder, then remove to safe place. Personal		Xn symbol
protection: P2 filter respirator for harmful		R: 40-43
particles.		S: 2-22-36

SEE IMPORTANT INFORMATION ON BACK

ICSC: 0062

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International Chemical Safety Cards

NICKEL ICSC: 0062

PHYSICAL STATE; APPEARANCE:

SILVERY METALLIC SOLID IN VARIOUS FORMS.

ROUTES OF EXPOSURE:

The substance can be absorbed into the body by inhalation of the dust.

T

PHYSICAL DANGERS:

M P O R T A N T D A T A	Dust explosion possible if in powder or granular form, mixed with air. CHEMICAL DANGERS: Reacts violently, in powder form, with titanium powder and potassium perchlorate, and oxidants such as ammonium nitrate, causing fire and explosion hazard. Reacts slowly with non-oxidizing acids and more rapidly with oxidizing acids. Toxic gases and vapours (such as nickel carbonyl) may be released in a fire involving nickel. OCCUPATIONAL EXPOSURE LIMITS: TLV: (Inhalable fraction) 1.5 mg/m³ as TWA A5 (not suspected as a human carcinogen); (ACGIH 2004). MAK: (Inhalable fraction) sensitization of respiratory tract and skin (Sah); Carcinogen category: 1; (DFG 2004). OSHA PEL*‡: TWA 1 mg/m³ *Note: The PEL does not apply to Nickel carbonyl. NIOSH REL*: Ca TWA 0.015 mg/m³ See Appendix A	INHALATION RISK: Evaporation at 20°C is negligible; a harmful concentration of airborne particles can, however, be reached quickly when dispersed. EFFECTS OF SHORT-TERM EXPOSURE: May cause mechanical irritation. Inhalation of fumes may cause pneumonitis. EFFECTS OF LONG-TERM OR REPEATED EXPOSURE: Repeated or prolonged contact may cause skin sensitization. Repeated or prolonged inhalation exposure may cause asthma. Lungs may be affected by repeated or prolonged exposure. This substance is possibly carcinogenic to humans.
A	*Note: The REL does not apply to Nickel carbonyl. NIOSH IDLH: Ca 10 mg/m³ (as Ni) See: 7440020	
PHYSICAL PROPERTIES	Boiling point: 2730°C Melting point: 1455°C Density: 8.9 g/cm3	Solubility in water: none
ENVIRONMENTAL DATA		
	NOTES	
symptoms of asthma oft	ickel oxide fumes will be formed. Depending on the degree of ten do not become manifest until a few hours have passed and re essential. Anyone who has shown symptoms of asthma due	d they are aggravated by physical effort. Rest and medical

substance.

ADDITIONAL INFORMATION ICSC: 0062 **NICKEL** (C) IPCS, CEC, 1994

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ZINC POWDER ICSC: 1205











Blue powder
Merrillite
Zn
Atomic mass: 65.4
(powder)

ICSC # 1205

CAS # 7440-66-6 RTECS # ZG8600000

UN # 1436 (zinc powder or dust)

EC# 030-001-00-1

October 24, 1994 Peer reviewed









TYPES OF HAZARD/ EXPOSURE	ACUTE HAZA SYMPTON		PREVENTION		FIRST AID/ FIRE FIGHTING
FIRE	Highly flammable. Many cause fire or explosion. C irritating or toxic fumes (fire.	Gives off	NO open flames, NO sparks, and smoking. NO contact with acid(s) (s) and incompatible substances (see Chemical Dangers).	, base	Special powder, dry sand, NO other agents. NO water.
EXPLOSION	Risk of fire and explosion with acid(s), base(s), wat incompatible substances.	er and	Closed system, ventilation, explos proof electrical equipment and lig Prevent build-up of electrostatic charges (e.g., by grounding). Prev deposition of dust.	hting.	In case of fire: cool drums, etc., by spraying with water but avoid contact of the substance with water.
EXPOSURE			PREVENT DISPERSION OF DU STRICT HYGIENE!	JST!	
•INHALATION	Metallic taste and metal f Symptoms may be delayed		Local exhaust.		Fresh air, rest. Refer for medical attention.
•SKIN	Dry skin.		Protective gloves.		Rinse and then wash skin with water and soap.
•EYES			Safety spectacles.		First rinse with plenty of water for several minutes (remove contact lenses if easily possible), then take to a doctor.
•INGESTION	Abdominal pain. Nausea.	. Vomiting.	Do not eat, drink, or smoke during work. Wash hands before eating.	g	Rinse mouth. Refer for medical attention.
SPILLAGI	E DISPOSAL.		STORAGE	PΔ	CKAGING & LABELLING

SPILLAGE DISPOSAL	STORAGE	PACKAGING & LABELLING
	Fireproof. Separated from acids, bases oxidants	<u> </u>
NOT wash away into sewer. Sweep spilled	Dry.	F symbol
substance into containers. then remove to safe		N symbol
place. Personal protection: self-contained		R: 15-17-50/53
breathing apparatus.		S: 2-7/8-43-46-60-61
		UN Hazard Class: 4.3
		UN Subsidiary Risks: 4.2

SEE IMPORTANT INFORMATION ON BACK

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ICSC: 1205 ZINC POWDER

ROUTES OF EXPOSURE:

and by ingestion.

The substance can be absorbed into the body by inhalation

PHYSICAL STATE; APPEARANCE:

PHYSICAL DANGERS:

ODOURLESS GREY TO BLUE POWDER.

P	Dust explosion possible if in powder or granular form,	INHALATION RISK:
0	mixed with air. If dry, it can be charged electrostatically l swirling, pneumatic transport, pouring, etc.	
R	CHEMICAL DANGERS:	
Т	Upon heating, toxic fumes are formed. The substance is a strong reducing agent and reacts violently with oxidants. Reacts with water and reacts violently with acids and bas	Inhalation of fumes may cause metal fume fever. The
A	forming flammable/explosive gas (hydrogen - see	
N	ICSC0001) Reacts violently with sulfur, halogenated hydrocarbons and many other substances causing fire and explosion hazard.	EFFECTS OF LONG-TERM OR REPEATED EXPOSURE: Repeated or prolonged contact with skin may cause
T	OCCUPATIONAL EXPOSURE LIMITS:	dermatitis.
D	TLV not established.	
A		
T		
A		
PHYSICAL PROPERTIES	Boiling point: 907°C Melting point: 419°C Relative density (water = 1): 7.14	Solubility in water: reaction Vapour pressure, kPa at 487°C: 0.1 Auto-ignition temperature: 460°C
ENVIRONMENTAL DATA		
	NOTES	
violently with fire extir	amounts of arsenic, when forming hydrogen, may also for a neguishing agents such as water, halons, foam and carbon discours later. Rinse contaminated clothes (fire hazard) with pl	oxide. The symptoms of metal fume fever do not become
	ADDITIONAL INFORM	ATION
	ADDITIONAL INFORM	ATION
ICSC: 1205	JI	ZINC POWDER

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APPENDIX D HOSPITAL INFORMATION AND MAP FIELD ACCIDENT REPORT

FIELD ACCIDENT REPORT

This report is to be filled out by the designated Site Safety Officer after EVERY accident.

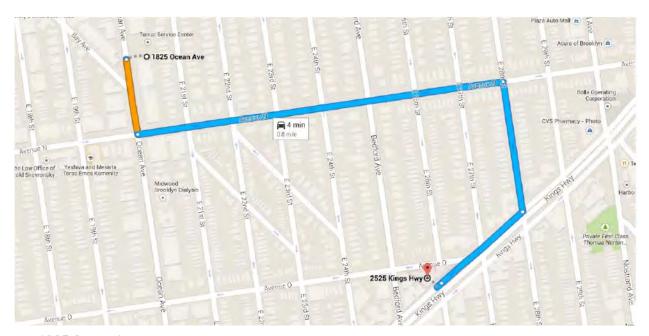
PROJECT NAME		PROJECT. NO		
Date of Accident	Time	Report By		
Type of Accident (Check One)	:			
() Vehicular	() Personal	() Property		
Name of Injured		DOB or Age		
How Long Employed				
Action Taken				
Did the Injured Lose Any Time	? How Much	(Days/Hrs.)?		
Was Safety Equipment in Us Shoes, etc.)?	e at the Time of the	Accident (Hard Hat, Safety Glasses,	Gloves,	Safety
(If not, it is the EMPLOYEE' Welfare Fund.)	S sole responsibility t	o process his/her claim through his/	<u> </u>	lth and
INDICATE STREET NAMES, I	DESCRIPTION OF VE	HICLES, AND NORTH ARROW		

HOSPITAL INFORMATION AND MAP

The hospital nearest the site is:

New York Community Hospital

2525 Kings Highway Brooklyn, New York 11229 0.8 miles – 4 min (718)-692-5300



o 1825 Ocean Ave

Brooklyn, NY 11230

Brooklyn, NY 11229

1. Head south on Ocean Ave toward Bay Ave

436 ft

2. Turn left onto Avenue N

3. Turn right onto E 28th St

4. Turn right onto Kings Hwy

5. Keep right to stay on Kings Hwy

5. Keep right to stay on Kings Hwy

5. Keep right to stay on Kings Hwy

6. Turn right onto Kings Hwy

7. Turn right onto Kings Hwy

8. Turn right onto Kings Hwy

9. Turn right onto Kings Hwy

9. Turn right to stay on Kings Hwy

9. Turn right to stay on Kings Hwy

9. Turn right to stay on Kings Hwy

SVE/AS Pilot Test Work Plan: Community Action Monitoring Plan (CAMP)

COMMUNITY AIR MONITORING PLAN

TOMAT SERVICE STATION 1815-1825 OCEAN AVENUE BROOKLYN, NY

August - 2016

COMMUNITY AIR MONITORING PLAN TABLE OF CONTENTS

1815-1825 Ocean Avenue, Brooklyn, NY

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APPENDICES

Appendix A Action Limit Report

1.0 INTRODUCTION

This Community Air Monitoring Plan (CAMP) has been prepared for the excavation and construction activities to be performed under the SVE/AS Pilot Test Work Plan at the Tomat Service Station located at 18165-1825 Ocean Avenue, Brooklyn, NY. The CAMP provides measures for protection for the downwind community (i.e., off-site receptors including residences, businesses, and on-site workers not directly involved in the investigation activities) from potential airborne contaminant releases resulting from excavation activities at the site.

Compliance with this CAMP is required during all activities associated with soil disturbance activities that have the potential to generate airborne particulate matter and volatile organic compounds (VOCs). These activities include excavation and loading of affected soil. This CAMP has been prepared to ensure that remedial activities do not adversely affect passersby, residents, or workers in the area immediately surrounding the Site and to preclude or minimize airborne migration of site-related contaminants to off-site areas.

1.1 Regulatory Requirements

This CAMP was established in accordance with the following requirements:

New York State Department of Health's (NYSDOH) Generic Community Air Monitoring Plan
as presented in DER-10 Technical Guidance for Site Investigation and Remediation (NYSDEC
May 3, 2010). This guidance specifies that a community air-monitoring program shall be
implemented to protect the surrounding community and to confirm that the work does not spread
contamination off-site through the air.

2.0 AIR MONITORING

Petroleum volatile organic compounds (VOCs) are the constituents of concern at the Site. The appropriate method to monitor air for these constituents during remediation activities is through real-time VOC and air particulate (dust) monitoring.

2.1 Meteorological Data

At a minimum, wind direction will be evaluated at the start of each workday, noon of each workday, and the end of each workday. These readings will be utilized to position the monitoring equipment in appropriate upwind and downwind locations.

2.2 Community Air Monitoring Requirements

To establish ambient air background concentrations, air will be monitored at several locations around the site perimeter before activities begin. These points will be monitored periodically in series during the site work. When the excavation area is within 20 feet of potentially exposed populations or occupied structures, the perimeter monitoring points will be located to represent the nearest potentially exposed individuals at the downwind location and will take into account the locations of ventilation system intakes of nearby structures.

Fugitive respirable dust will be monitored using a MiniRam Model PDM-3 aerosol monitor (or equivalent). Air will be monitored for VOCs with a portable Ionscience 3000 photoionization detector (PID), or equivalent. All air monitoring data will be documented in a site log book by the designated site safety officer. The site safety officer or delegate must ensure that air monitoring instruments are calibrated and maintained in accordance with manufacturer's specifications. All instruments will be zeroed daily and checked for accuracy. A daily log will be kept. If additional monitoring is required, the protocols will be developed and appended to this plan

When work area are within in 20 feet of the potentially exposed populations or occupied structures, the continuous monitoring locations for VOCs and particulates must reflect the nearest potentially exposed individuals and the location of ventilation system intakes for nearby structures. The use of engineering controls such base vapor / dust barriers, temporary negative pressure enclosures, or special ventilation devices should be considered to prevent exposures related to work activities and to control dust and odors. Consideration should be given to implementing the planned activities when potentially exposed populations are at a minimum, such as during the weekends or evening hours in non-residential settings.

- If total VOC concentrations opposite the walls of occupied structures or next to the intake
 vents exceed 1 ppm, monitoring should occur within the occupied structures (if access is
 granted by occupants). Background readings in the occupied spaces must be take prior to the
 commencement of the planned work. Any unusual background readings should be discussed
 with NYSDOH prior to the commencement of work.
- If total particulate concentrations opposite the walls of the occupied structures or next to the intake vents exceed 150 μ g/m³, work activities should be suspended until controls are implemented and are successful in reducing the total particulate concentration to 150 μ g/m³ or less at the monitoring point.

3.0 VOC MONITORING, RESPONSE LEVELS, AND ACTIONS

Volatile organic compounds (VOCs) will 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. 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.

- 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.
- 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.
- If the organic vapor level is above 25 ppm at the perimeter of the work area, activities must be shutdown. All 15-minute readings must be recorded and be available for State (DEC and DOH) personnel to review. Instantaneous readings, if any, used for decision purposes should also be recorded.

All readings will be recorded and made available for NYSDEC and NYSDOH personnel to review. If an exceedance of the Action Limits occurs, an Action Limit Report, as shown in Appendix A, will be completed.

3.1 Potential Corrective Measures and VOC Suppression Techniques

If the 15-minute integrated VOC level at the downwind location persists at a concentration that exceeds the upwind level by more than 5 ppm but less than 25 ppm during remediation activities, then vapor suppression techniques will be employed. The following techniques, or others, may be employed to mitigate the generation and migration of fugitive organic vapors:

- Collection of purge water in covered containers;
- storage of excess sample and drill cuttings in drums or covering with plastic



4.0 PARTICULATE MONITORING

Air monitoring for particulates (i.e., dust) will be performed continuously during excavation and loading activities using both air monitoring equipment and visual observation at upwind and downwind locations. Monitoring equipment capable of measuring particulate matter smaller than 10 microns (PM10) and capable of integrating (averaging) over periods of 15 minutes or less will be set up at upwind (i.e., background) and downwind locations, at heights approximately four to five feet above land surface (i.e., the breathing zone). Monitoring equipment will be MIE Data Ram monitors, or equivalent. The audible alarm on the particulate monitoring device will be set at 90 micrograms per cubic meter (μ g/m3). This setting will allow proactive evaluation of worksite conditions prior to reaching the action level of 100 μ g/m³ above background. The monitors will be calibrated at least once per day prior to work activities and recalibrated as needed thereafter. In addition, fugitive dust migration will be visually assessed during all intrusive work activities.

The following summarizes particulate action levels and the appropriate responses:

- If the downwind PM-10 particulate level is 100 μg/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 μg/m³ above the upwind level and provided that no visible dust is migrating from the work area.
- If, after implementation of dust suppression techniques, downwind PM-10 particulate levels are greater than 150 μg/m³ above the upwind level, work must be stopped and an evaluation of activities initiated. Work can resume provided that dust suppression measures (as described in Section 2.3.1 below) and other controls are successful in reducing the downwind PM-10 particulate concentration to within 150 μg/m³ of the upwind level and in preventing visible dust migration.

All readings will be recorded and be available for NYSDEC and NYSDOH personnel to review. If an exceedance of the Action Limits occurs, an Action Limit Report as shown in **Appendix A** will be completed.

4.1 Potential Particulate Suppression Techniques

If the integrated particulate level at the downwind location exceeds the upwind level by more than $100~\mu g/m^3$ at any time during remediation activities, then dust suppression techniques will be employed. The following techniques, or others, may be employed to mitigate the generation and migration of fugitive dusts:

- limiting the excavation size;
- backfilling the excavation;
- spraying water onto the excavation faces and equipment;



- covering soil stockpiles with 8-mil plastic sheeting;
- hauling waste materials in properly tarped containers; and/or
- limiting vehicle speeds onsite.

Work may continue with dust suppression techniques provided that downwind PM_{10} levels are not more than 150 $\mu g/m^3$ greater than the upwind levels.

There may also be situations where the dust is generated by remediation activities and migrates to downwind locations, but is not detected by the monitoring equipment at or above the action level. Therefore, if dust is observed leaving the working area, dust suppression techniques such as those listed above will be employed.

If dust suppression techniques do not lower particulates to below $150 \,\mu\text{g/m}^3$, or visible dust persists, work will be suspended until appropriate corrective measures are identified and implemented to remedy the situation.

All air monitoring readings will be recorded in the field logbook and will be available for the NYSDEC and NYSDOH personnel to review.

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5.0 DATA QUALITY ASSURANCE

5.1 Calibration

Instrument calibration shall be documented on instrument calibration and maintenance sheets or in the designated field logbook. All instruments shall be calibrated as required by the manufacturer. Calibration checks may be used during the day to confirm instrument accuracy. Duplicate readings may be taken to confirm individual instrument response.

5.2 Operations

All instruments shall be operated in accordance with the manufacturer's specifications. Manufacturers' literature, including an operations manual for each piece of monitoring equipment will be maintained on-site by the SSO for reference.

5.3 Data Review

The SSO will interpret all monitoring data based the established criteria and his/her professional judgment. The SSO shall review the data with the PM to evaluate the potential for worker exposure, upgrades/downgrades in level of protection, comparison to direct reading instrumentation and changes in the integrated monitoring strategy.

Monitoring and sampling data, along with all sample documentation will be periodically reviewed by the PM.

6.0 RECORDS AND REPORTING

All air readings must be recorded on daily air monitoring log sheets and made available for review by personnel from NYSDEC and NYSDOH.

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APPENDIX A ACTION LIMIT REPORT

CAMP ACTION LIMIT REPORT

Project Location:				
Date:	-	Time:		
Name:	-			
Contaminant:	PM-10:	VOC:		
Wind Speed:	_	Wind Direction:		
Temperature:	_	Barometric Pressure:		
DOWNWIND DATA Monitor ID #:	Location:	Level Reported:		
Monitor ID#:	Location:	_ Level Reported:		
UPWIND DATA Monitor ID #:	Location:	_ Level Reported:		
Monitor ID#:	Location:	_ Level Reported:		
BACKGROUND CORRECTED LEVELS				
Monitor ID #: Location:	_ Level Reported: Leve	el Reported:		
ACTIONS TAKEN				