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WORK PLAN FOR SOIL VAPOR EXTRACTION PILOT TEST

eder associates environmental scientists and engineers NATIONAL PRESTO INDUSTRIES. INC. EAU CLAIRE, WISCONSIN

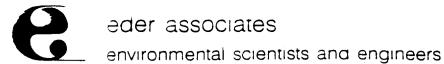
WORK PLAN FOR SOIL VAPOR EXTRACTION PILOT TEST

PROJECT #497-18 MARCH 1994

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Offices in New York, Wisconsin, Michigan, Georgia, Florida, and New Jersey

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March 17, 1994 File #497-18

Ms. Eugenia Chow Remedial Project Manager United States Environmental Protection Agency - Region 5 77 West Jackson Boulevard - HSRW6J Chicago, Illinois 60604-3590

> Re: National Presto Industries. Inc. Eau Claire. Wisconsin

Dear Ms. Chow:

Two copies of the Soil Vapor Extraction Pilot Test Work Plan are enclosed. The work plan has been revised in accordance with the comments contained in your March 8, 1994 letter to NPI. Please advise if you have any additional questions or comments. I am available to discuss the work plan at your convenience.

Very truly yours,

EDER ASSOCIATES

Nora M. Brew

Nora M. Brew Project Engineer

NB/cg Encl.

- cc: R. Riedl (2 copies)
 - S. Thon
 - O. Patel
 - D. Manning
 - R. Nauman
 - W. Warren
 - D. Gan

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VOCs to be Analyzed by Portable Gas Chromatograph

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1.0 INTRODUCTION

1.1 Background Information

The National Presto Industries, Inc. (NPI) site is an inactive manufacturing facility comprising approximately 320 acres in Eau Claire, Wisconsin. A map of the NPI site is presented on Figure 1. The site was originally owned by the United States Government and operated between 1940 and 1947 by its contractors, the U.S. Rubber Company (Uniroyal) as a small arms ammunition plant, and Western Electric (AT&T) as a radar tube manufacturing plant. NPI purchased the site from the government in 1947. Between 1947 and 1954, NPI used the facility to produce consumer products, projectile fuzes, and military aircraft parts. Since 1954, all manufacturing activities at the NPI site have been dedicated to defense work for the Department of Defense (DOD). Between 1954 and 1959, the facility produced military aircraft parts and between 1966 and 1980 the facility produced 8-inch and 105 mm projectiles for the Department of the Army (DOA). Active production ceased in February 1980.

Waste forge compound, which in its virgin state consists of approximately equal parts of asphalt, graphite and mineral oil, was the primary waste material generated by the projectile production operation. Volatile organic compounds (VOCs) became incorporated into the waste forge compound during the forging operations. Waste forge compound with a low solids content flowed by gravity and was pumped to Lagoon No. 1 with process wastewaters between 1966 and 1980. From 1966 to 1970, waste forge compound with a higher solids content was heated in a basement collection sump so that it could be pumped into drums. The drums were hauled to the Melby Road Disposal Site (MRDS) where the waste forge compound was disposed of. The waste disposal area was originally delineated based on aerial photographs from 1968, which showed a trench area at the MRDS. Disposal of waste forge compound at the MRDS ceased in 1970 after a reclamation and recycling program was developed by NPI. The MRDS area was regraded in 1970.

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The NPI site Remedial Investigation (RI) indicates the presence of VOCs and/or metals at the MRDS. The most commonly found VOCs were 1,1,1-trichloroethane (TCA), trichloroethylene (TCE), tetrachloroethylene (PCE), 1,1-dichloroethylene (1,1-DCE), and 1,1-dichloroethane (1,1-DCA).

As the Feasibility Study (FS) was being prepared for the NPI site, it became apparent that more information about the MRDS was needed to effectively evaluate potential remedial alternatives. Data gaps existed with respect to the vertical and areal extent of contamination and questions remained regarding the specific source of VOCs in groundwater. In addition, aerial photographs obtained from the Eau Claire County Health Department in early 1993 revealed that the waste disposal area was larger than originally delineated. Additional sampling and analysis was conducted at the MRDS in August 1993 to further define the nature and extent of waste disposal and soil contamination. The investigation was conducted in accord with the Pre-Design Pilot Studies Work Plan (Eder Associates, August 1993) approved by the United States Environmental Protection Agency (USEPA) and the Wisconsin Department of Natural Resources (WDNR).

The August 1993 investigation revealed that a variety of wastes were disposed of at the MRDS in a random manner. The wastes found included empty drums, drums containing waste forge compound, construction and demolition debris, and waste forge compound mixed with varying amounts of soil. Waste (primarily waste forge compound/soil mixture) and soil samples were collected from the test pits excavated and the soil borings drilled during the MRDS investigation and analyzed. The MRDS characterization results will be summarized in the RI report.

Soil vapor extraction (SVE) may be a component of the final remedy for the MRDS. An effective SVE system would act as a barrier to prevent the downward movement of vapors. A pilot test will be conducted to determine whether SVE would be effective in preventing VOC vapor transport through the vadose zone to groundwater at the MRDS. The pilot test will also provide the information required to determine design parameters (i.e. - radius of influence. air extraction rates, well spacing, screen intervals, and VOC concentrations in the extracted air) for a full-scale SVE system. This work plan describes the technical protocol for the SVE pilot test.

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1.2 Pilot Test Approach

The SVE pilot test will be conducted at three locations selected based on the waste and soil characteristics identified during the August 1993 investigation. The test locations are shown on Figure 2 and described in Section 2.0. The test will determine whether SVE would be effective in creating a vapor barrier.

The pilot test will be implemented in two phases. Each location will be tested independently to avoid interference between the results collected for each pilot system. An array of soil monitoring points will be used to measure subsurface pressures and vapor concentrations. Vapors will be extracted from each test well to establish subsurface pressures that are representative of steady-state or dynamic conditions during the first phase of the pilot test. This first phase of the test is expected to run for less than eight hours. The results will be used to determine the air permeability of the subsurface materials, the radius of influence of the vapor extraction wells, and the need for VOC emission control during longer term operation.

The second phase of the pilot test will provide extended system operating information to determine the effectiveness of SVE based on sustained negative pressures and decreases in VOC vapor concentrations. This second phase of the test is expected to run for about two weeks at each location.

2.0 TEST WELLS AND EQUIPMENT

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Three SVE test wells (VW-1 through VW-3) and nine nested monitoring points (MP-1 through MP-9) will be installed at the MRDS for the pilot test. The proposed locations of the vapor extraction wells and monitoring points are shown on Figure 2. Cross-sectional views of the vapor extraction wells and monitoring points are shown on Figures 3 and 4.

2.1 Soil Vapor Extraction Wells

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Vapor extraction well VW-1 will be located near the deep trench area and screened from 25 feet to 30 feet below grade. VW-2 will be located in the area where the highest VOC concentrations were found in waste and soil during the August 1993 investigation. VW-2 will have a 30-foot long screen beginning at 30 feet below grade. VW-3 will be located in the eastern portion of the MRDS where the wastes were relatively consistent and representative of conditions found at the eastern and western portions of the MRDS during the August 1993 work. VW-3 will be screened from about 5 feet to 10 feet below grade. All three vapor extraction wells will be screened in the unsaturated native soil beneath the waste.

The SVE wells will be constructed of four-inch diameter Schedule 40 PVC in boreholes drilled using a 6.25-inch inner diameter hollow stem auger. A temporary 12-inch steel outer casing will be installed to at least two feet below the waste at each drilling location. The 12-inch casing will be installed in a 15-inch borehole drilled using mud-rotary methods. Hollow stem augering will then proceed through the outer casing to the required depth. Each well will have a Schedule 40 PVC screen with 0.020-inch slots, and a filter pack of #30 silica sand terminating approximately two feet above the top of the screen. A hydrated bentonite seal will be placed on top of a fine sand filter pack seal. Water will be added to hydrate the bentonite at five foot intervals. A native soil surface seal will be placed on top of the hydrated bentonite seal. The wells will terminate above grade with a tee fitting for connection to the vacuum blower. A detail of the

vapor extraction wells is presented on Figure 5. The temporary casing will be removed after well installation is completed.

Samples will be collected during drilling at 2.5-foot intervals using 24-inch split spoon samplers. The samplers will be decontaminated by steam cleaning if waste adheres to the split spoons. The sampling interval may be modified in the field based on the characteristics of the native material. Samples will be collected less frequently if the material is homogeneous, and more frequently if variations are observed. The native soil samples collected from each boring will be submitted to a laboratory for grain size analysis. The number of samples to be submitted for grain size analysis may be reduced in the field if the native material is homogeneous. Field decisions will be made in consultation with USEPA and WDNR. Subsurface lithology will be logged and classified by the hydrogeologist overseeing the drilling. The lithologic description will include the following:

- approximate percentages of major and minor grain-size constituents,
- color.
- geologic origin.
- moisture (qualitative),
- visual presence of secondary permeability,
- voids or layering, and
- any other pertinent observations.

2.2 Soil Gas Monitoring Points

Soil gas monitoring points will be used to collect data during the MRDS pilot test. The soil gas monitoring points are designed to evaluate the SVE system parameters horizontally and vertically over the test area, and to obtain information for evaluating air flow in three-dimensions. Five monitoring points (MP-1, MP-2, MP-3, MP-4, and MP-5) will be located between VW-1 and VW-2, as shown on Figures 2 and 3. These five monitoring points will be used to record pressure changes and collect vapor samples during the SVE tests at VW-1 and VW-2. Four

monitoring points (MP-6, MP-7, MP-8, and MP-9) will be used to record pressure changes and collect vapor samples during the SVE test at VW-3, as shown on Figures 2 and 4. Each monitoring point will contain nested probes placed at depths relative to the screened zone of the vapor extraction well(s) it is intended to monitor.

Monitoring point MP-1 will contain three nested soil gas monitoring probes. The upper probe will be screened at 20 feet, the middle probe at 30 feet, and the bottom probe at 45 feet below grade. Monitoring points MP-2 through MP-5 will contain four nested soil gas monitoring probes. The four monitoring probes will be screened at 20 feet, 30 feet, 45 feet, and 55 feet below grade. Monitoring point MP-6 will contain four nested soil gas monitoring probes. The uppermost probe at MP-6 will be screened entirely within the waste, and the sand filter pack for this probe will not extend into the cover material above the waste or the native soil below. The remaining probes at MP-6 will be screened at approximately 5 feet, 15 feet, and 25 feet below grade. The 5-foot probe will be screened entirely within the native soil beneath the waste. The difference in pressures measured at the uppermost probe and the 5-foot probe at MP-6 will provide qualitative information on the permeability of the waste and whether vapor extraction can establish air flow through the waste. Monitoring points MP-7 through MP-9 will contain three nested soil gas monitoring probes at MP-7, MP-8, and MP-9 will be screened entirely within the native soil beneath the waste.

All monitoring points will be constructed using a 6.25-inch inner diameter hollow stem auger. A temporary 12-inch steel outer casing will be installed to at least two feet below the waste at each drilling location. The 12-inch casing will be installed in a 15-inch borehole drilled using mud-rotary methods. Hollow stem augering will then proceed through the outer casing to the required depth. The probes will be one-inch diameter Schedule 40 PVC terminating with a sixinch long section of slotted screen. The probe screens will be segregated by hydrated bentonite seals and imbedded in a #30 silica sand filter pack. Each probe will have a quick-connect, airtight coupling for use with a vacuum gauge. A detail of the soil gas monitoring point construction is shown on Figure 6. Each probe will be labeled to identify its depth. The temporary casing will be removed after monitoring point installation is completed.

Samples will be collected at 2.5-foot intervals using 24-inch split spoon samplers when drilling MP-1. MP-5, and MP-6. The samplers will be decontaminated by steam cleaning if waste adheres to the split spoons. Subsurface lithology will be logged and classified by the hydrogeologist overseeing the drilling, as previously discussed. A five-foot (or greater) sampling interval may be used for the remaining monitoring points (MP-2, MP-3, MP-4, MP-7, MP-8, and MP-9) if the hydrogeologist determines that the native material is homogeneous. Sampling would resume at more frequent intervals if the hydrogeologist observes any variations in the material. The native soil samples collected from each boring will be submitted to a laboratory for grain size analysis. The number of samples to be submitted for grain size analysis may be reduced in the field if the native material is homogeneous. Field decisions will be made in consultation with USEPA and WDNR.

2.3 Soil Vapor Extraction Equipment

The SVE test will be conducted using a trailer mounted spark-proof positive displacement vacuum blower (Roots Model 33 URAI or equivalent) with a 5 horsepower explosion-proof motor and switch. This blower has an airflow range of 25 to 175 scfm at a maximum vacuum of ten inches of mercury. The trailer mounted rig is also equipped with a vacuum relief valve, an 80-gallon condensate separator tank, a discharge silencer. four manifold two-inch hose connections, vacuum gauges, and airflow meters. The blower is belt-driven and can be sheaved to provide nominal flow rates at 25 scfm increments. A bleed valve can be adjusted to provide minor flow rate variations. Vapor extraction well VW-2 has a 30-foot screen, and it may be necessary to use two or more blowers in parallel to extract a sufficient air flow to establish a subsurface vacuum at a reasonable distance from this well. The need to use additional blowers will be determined during implementation of the pilot test at WV-2.

The soil gas permeability test will run for a maximum of 8 hours at an air flow rate of approximately 100 scfm. More than 100 scfm may be extracted at VW-2 if necessary to obtain adequate data. USEPA and WDNR would be consulted before increasing the air flow rate at VW-2. The VOC data collected during the first phase of the SVE pilot test will be evaluated to determine whether emission controls would be required for longer term operation. The exhaust gas from the second phase of the SVE pilot test will be treated with activated carbon if necessary to comply with the limitations specified in Chapter NR 445. Wisconsin Administrative Code. Returnable 55 gallon (200 lb) carbon vessels will be staged on-site and used as necessary.

2.4 Field Instrumentation and Measurements

2.4.1 Concentrations of Volatile Organic Compounds

Soil gas and exhaust gas samples will be collected from the monitoring points and SVE wells using disposable aluminum vacuum sample canisters with a capacity of approximately 280 cc. The samples will be analyzed for the VOCs listed in Table 1, which were detected in samples collected at the MRDS, using a portable gas chromatograph (GC) (Photovac GC Model #10S50) equipped with a photoionization detector (PID). The portable GC operation procedures are presented in Appendix A.

2.4.2 Pressure/Vacuum Monitoring

Changes in soil gas pressure during the air permeability test will be measured at the soil gas monitoring points using MagnehelicTM or equivalent gauges with pressure ranges of 0 to 100 inches water. TygonTM or equivalent tubing will be used to connect the pressure/vacuum gauge to the quick-disconnect at the top of each soil gas monitoring probe. Pressure gauges on the SVE unit will also be monitored during the test. The gauges are sealed and calibrated at the factory and will be zeroed before each test.

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All attachments to the monitoring probes will be air tight. Vacuum measurement will be to two digits of accuracy (e.g. 0.01 to 0.99, 1.0 to 9.9, and 10 to 99 inches water column).

2.4.3 Air Flow Rate Measurement

Air extraction flow rates will be measured by the flowmeter mounted on the SVE unit during the pilot test. The flowmeter has a capacity up to 180 scfm and will be zeroed before each test. All flow rates will be corrected to standard temperature and ambient pressure conditions.

2.4.4 Temperature and Relative Humidity Measurements

Temperature and relative humidity will be measured at the extraction wells using a portable measurement device.

3.0 PILOT TEST PROCEDURES

3.1 Soil Gas Permeability Test Procedures

The first phase of the SVE test will gather data to estimate the soil gas permeability (k) and the radius of influence (ROI) of the vapor extraction well, to obtain qualitative information on the permeability of the waste, and to evaluate the need for VOC emission controls during the second phase of the test. The analytical method that will be used to estimate these parameters is presented in Appendix B.

3.1.1 System Check

Soil gas samples will be collected from the vapor extraction well and soil gas monitoring points before the test begins. The samples will be analyzed for the VOCs listed in Table 1 to establish baseline concentrations. After the SVE unit has been connected to the vapor extraction well, air will be extracted for 10 to 15 minutes to ensure proper operation of the SVE unit, pressure gauges, and air flow gauges. Pressure response at each monitoring point will be measured before and after the system check period.

3.1.2 Test Procedures

The soil gas permeability test will begin after all monitoring point pressures have returned to zero after the system check. The test will be conducted independently at the three test areas. Samples will be collected and analyzed in accord with the schedule described in Section 4.1.

The first SVE test will be conducted at VW-2. If the data indicate that air extraction rates above the capacity of the blower are required to evaluate SVE at this location, the test would be discontinued to allow one or more additional blowers to be obtained. In the meantime, the SVE tests would be conducted at VW-1 and/or VW-3 to avoid delaying the overall pilot test project schedule.

The soil gas permeability test procedures are summarized below:

- 1. Connect the pressure gauge to the top of each monitoring probe and return the gauges to zero.
- 2. Turn the SVE unit on to extract approximately 100 scfm.
- 3. Record the pressure at each soil monitoring probe at one-minute intervals. (Note: VW-1 will be considered to be a monitoring probe during the test at VW-2).
- 4. After 10 minutes, record the pressure reading at the well head, the temperature and relative humidity of the exhaust gas, and the flow rate from the SVE unit. Record the pressure at each monitoring probe at two-minute intervals.
- 5. After 20 minutes, record the pressure at each monitoring probe at three-minute intervals. Continue to record all SVE unit data at ten-minute intervals.
- 6. Continue to record the pressure at each monitoring probe at three-minute intervals until the change in pressure is less than 0.1 inches of water. At this time, a 5- to 20-minute interval will be used.
- 7. Review results to ensure that accurate data are collected during the first 30 minutes of the test. If the quality of the data is in question (i.e. if a stronger vacuum is measured in monitoring points distant from the test well than in monitoring points closer to the well, there is a problem with the pressure gauges), turn off the blower, allow all monitoring probes to return to zero pressure, and restart the test.

Note: If the data indicate negligible pressure changes during the VW-2 test, additional blowers may be obtained and used in parallel to achieve sufficient data. USEPA and WDNR would be consulted before increasing the air flow rate at VW-2.

- 8. Continue to collect data for four hours to a maximum of eight hours. The test will be terminated when pressure readings do not change by more than 10 percent over a one-hour interval (assumed to represent steady state conditions), or after the test has run for eight hours.
- 9. Collect soil gas samples from the vapor extraction well and soil gas monitoring probes every two hours during the test and immediately after completing the soil gas permeability test. The samples will be analyzed for the VOCs listed in Table 1 using a portable GC.
- 10. Obtain hourly barometric pressure data and record information on weather conditions (precipitation events, presence of frost, etc.) for consideration during data evaluation. Meteorological data will be obtained from the Chippewa Valley Regional Airport in Eau Claire, Wisconsin.
- 11. Calculate k based on the test data using the method outlined in Appendix B. The steady state equations will be used if the pressure readings did not change by more than ten percent over a one-hour period. The dynamic equations will be used if the test does not reach representative steady state conditions and is ended after eight hours. The emissions of VOCs from the SVE system will be calculated and compared to the NR 445 limitations to evaluate the need for emission controls during the second phase SVE test and for full-scale system design.
- 12. The pilot test data will be evaluated using a three-dimensional model to determine the ROI of the vapor extraction wells.

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Any condensate collected by the SVE pilot system will be drummed. Headspace analysis using a portable GC will be performed on one condensate sample from each drum. The condensate will be analyzed for TCA, TCE, PCE, 1,1-DCE, and 1,1-DCA, and the results will be compared to the discharge limits established for these VOCs under the on-site groundwater interim remedial action (IRA). The headspace analysis of the condensate will be performed in accord with the USEPA-approved portable GC method for VOC screening of water samples (Eder, August 1991). The condensate will be hauled to the IRA cascade aerator at the MRDS for treatment and discharge to the Chippewa River if the VOC discharge limits are met. The condensate will be transported to an off-site treatment/disposal facility if the VOC discharge limits are exceeded.

3.2 Soil Vapor Extraction Test

The second phase SVE test will be conducted at the three test areas independently to evaluate longer term SVE system response and effectiveness. The air extraction rate will be determined based on an evaluation of the soil gas permeability test results.

The second phase SVE test results will be evaluated and the test discontinued when the data suggest that further operation would not provide useful information. The decision to discontinue the test at each location will be made in conjunction with USEPA and WDNR. It is expected that the three extended tests would be completed within a total of six weeks.

3.2.1 System Installation

The trailer mounted SVE system will be used for the extended SVE test. A carbon adsorption unit will be used to treat emissions if the exhaust-gas sampling results from the soil gas permeability test show that estimated benzene, individual VOC, and/or total VOC emissions would exceed the NR 445 allowable limits.

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3.2.2 System Operation and Maintenance

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The blower will be operated continuously throughout the extended SVE test periods. The system will be checked daily to ensure that the blower is operating within the intended flow rate, pressure, and temperature range. Maintenance such as replacing filters or gauges, or draining condensate, will be performed as necessary. Serious problems, such as motor or blower failures, will be corrected to ensure the credibility of SVE test.

Any condensate recovered by the SVE pilot system will be drummed. Headspace analysis using a portable GC will be performed on one condensate sample from each drum. The condensate will be analyzed for TCA, TCE, PCE, 1.1-DCE, and 1.1-DCA, and the results will be compared to the discharge limits established for these VOCs under the on-site groundwater IRA. The headspace analysis of the condensate will be performed in accord with the USEPA-approved portable GC method for VOC screening of water samples (Eder, August 1991). The condensate will be hauled to the IRA cascade aerator at the MRDS for treatment and discharge to the Chippewa River if the VOC discharge limits are met. The condensate will be transported to an off-site treatment/disposal facility if the VOC discharge limits are exceeded.

3.2.3 System Monitoring

Soil gas and exhaust gas sampling will be conducted to monitor the progress of the extended SVE tests. All SVE unit parameters, such as pressure, air extraction flow rates, temperature, and relative humidity will also be recorded. Information on barometric pressure and weather conditions (precipitation events, presence of frost, etc.) will be noted and considered when evaluating the data. Meteorological data will be obtained form the Chippewa Valley Regional Airport. A detailed sampling and analytical schedule is presented in Section 4.1.

4.0 SAMPLING AND ANALYSIS

4.1 Analytical Schedule

4.1.1 Soil Gas Permeability Test

The soil gas permeability test will be conducted for a maximum of eight hours. Samples will be collected from the vapor extraction well and soil gas monitoring probes before the test at each location. Samples will be collected every two hours during the test. At the end of the test, samples will be collected from the SVE system exhaust and soil gas monitoring probes immediately after the system is shut down. All samples will be analyzed for the VOCs listed in Table 1 using a portable GC.

4.1.2 Soil Vapor Extraction Test

The extended SVE test will run for approximately two weeks at each location. The three locations will be tested independently, so the second phase of testing is expected to run for approximately six weeks. Three samples of the SVE exhaust gas will be collected on the first three days of operation (at the beginning, middle, and end of each day) at each location. Samples will then be collected twice a week for the remainder of the test. The monitoring probes will be sampled at the beginning, middle, and end of the test period. All samples will be analyzed by portable GC for the VOCs listed in Table 1. The SVE system parameters, such as air flow rates, pressures, temperature, and relative humidity will be monitored during each sampling event. Information on weather conditions and barometric pressure will be recorded and considered when evaluating the data. Meteorological data will be obtained from teh Chippewa Valley Regional Airport.

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4.2 Sample Collection and Handling

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Soil vapor samples and exhaust gas samples will be collected from the monitoring probes and SVE wells. respectively, using disposable aluminum vacuum sample canisters which have a capacity of approximately 280 cc. At least two probe volumes of air will be purged using a portable pump before collecting samples from each soil gas monitoring probe. Once the samples are collected, the aluminum sample canisters will be sealed in a cooler, stored in a dark environment and preserved at 4°C for a maximum 48-hour holding time before analysis. The samples collected during the soil permeability tests will be analyzed on-site. The samples collected from the extended test will be shipped via overnight courier service to Eder's New York office, where they will be analyzed for the VOCs listed in Table 1 using a portable GC (Photovac GC Model #10S50) equipped with a PID. The data quality objective (DQO) for the sample analysis is Level 2 (field analyses).

4.3 Quality Control/Quality Assurance (QA/QC) Procedures

All instruments (i.e. - portable GC and pressure gauges) will be calibrated daily prior to use. The portable GC calibration will be checked at the end of each day. The calibration will follow standard manufacturers' instructions to assure that the equipment is functioning within the tolerances established by the analytical requirements. The following QA/QC procedures will be used during the collection and shipment of soil gas samples:

- Use disposable latex gloves when collecting soil vapor samples.
- Place all samples on ice in a cooler following collection. Analyze samples collected during the soil gas permeability tests using a portable GC brought onsite. Ship samples collected during the extended test via overnight courier service to Eder's New York office for portable GC analysis in accord with the procedures outlined in Appendix A.

• Complete a chain of custody record to accompany each sample shipment.

Field duplicates and trip blanks will be collected and analyzed to monitor sampling reproducibility, to check for cross-contamination in the sampling system, and to check for procedural contamination due to analyte migration during shipment and storage of samples. Field duplicate samples will be collected every 20 samples, or at least once a day if less than 20 samples are collected. One trip blank (vacuum canister filled with Ultra Zero air) will be used per sample shipment.

Appendix C presents the example chain of custody and field measurement and record forms.

5.0 PILOT TEST REPORT

A report will be prepared after the SVE pilot test is completed and the data are evaluated. The report will include the following as specified in the Wisconsin Soil Venting Guidance:

5.1 Discussion

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- A description of the test and final conclusions based on the objective to evaluate whether SVE would be effective in creating a vapor barrier. The text will include dates, weather (ambient temperature, wind, etc.), and any other pertinent field observations from the pilot test. The barometric pressure and whether climbing or falling will also be listed.
- Three-dimensional air flow modeling results. A 3-D model selected in consultation with USEPA and WDNR will be used to evaluate the pilot test data and develop a conceptual SVE system design.
- Conceptual full-scale SVE system design

5.2 Figures

- A site map drawn to scale (horizontal accuracy to +/- one foot). The map will indicate:
 - Locations of soil vapor extraction wells and soil gas monitoring points.
 - Zone of soil contamination.
 - Paved areas, buildings, and structures.
 - Buried utility trenches.

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- Scale, north arrow, title block, site name, key or legend, and date(s) of pilot test.
- Any other pertinent site information.
- Graphs representing subsurface vacuum at a distance from each soil vapor extraction well.
- A water table map of the site for the day of the pilot test.
- Cross sections showing screened intervals, geological units, contour lines of steady state vacuum readings, and soil monitoring points for each soil vapor extraction well.
- Plots of mass removal of VOCs vs. time for each soil vapor extraction well.

5.3 Tables

- Tabulated flow rates, vacuum distribution, soil gas temperatures, times of readings, ambient barometric pressure, and the ambient temperature.
- Water levels in existing groundwater monitoring wells.

5.4 Appendices

- A complete description of the field equipment and field procedures that were used.
- Sampling methods and procedures.
- Analytical methods, analytical results, and lab reports.

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• Boring logs and well construction diagrams for the soil vapor extraction wells and the soil gas monitoring points.

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• Grain size analysis results.

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- Engineering calculations.
- Any other pertinent field data.

6.0 TEST SCHEDULE

Figure 7 presents the proposed schedule for the SVE system pilot test.

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7.0 HEALTH AND SAFETY

The SVE pilot test will be conducted in accordance with the Health and Safety Plan presented in Appendix D.

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NATIONAL PRESTO INDUSTRIES, INC. EAU CLAIRE, WISCONSIN

TABLE 1

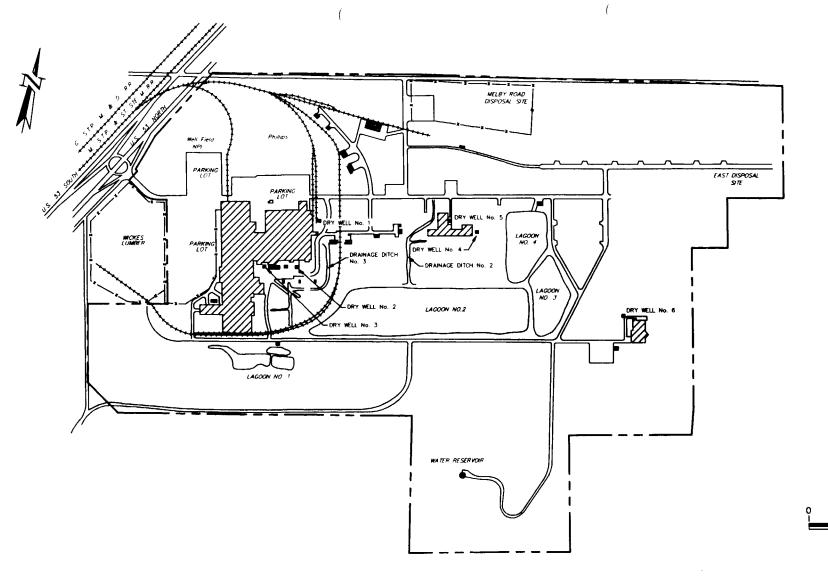
SVE PILOT TEST

VOCs TO BE ANALYZED BY PORTABLE GAS CHROMATOGRAPH

1,1-Dichloroethane 1,1,1-Trichloroethane Tetrachloroethylene Trichloroethylene 2-Butanone Benzene Toluene Xylenes (Total) Ethylbenzene Acetone Methylene Chloride Chloroethane 2-Hexanone 4-Methyl-2-pentanone 1.2-Dichlorobenzene 1.1-Dichloroethylene Styrene

NOTE:

The GC results will be added to estimate total VOC emission rates.



SITE PLAN NATIONAL PRESTO INDUSTRIES, INC. SITE EAU CLAIRE, WISCONSIN

500'

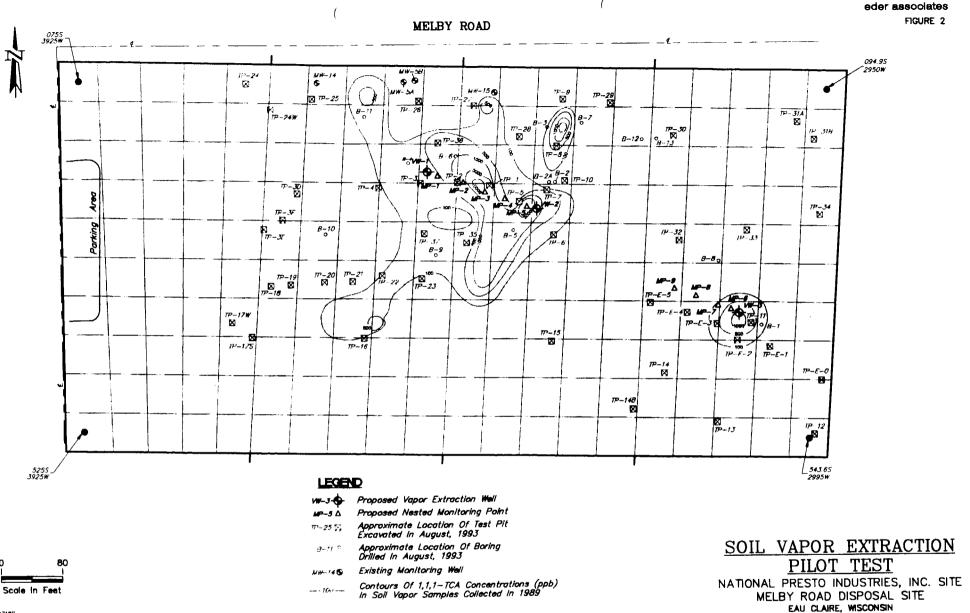
eder associates FIGURE 1

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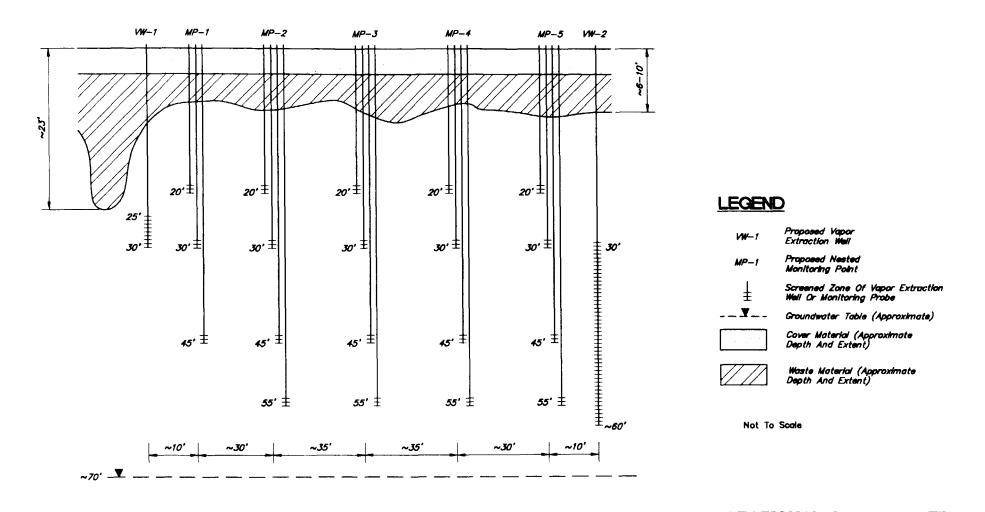
SK 497188 022494

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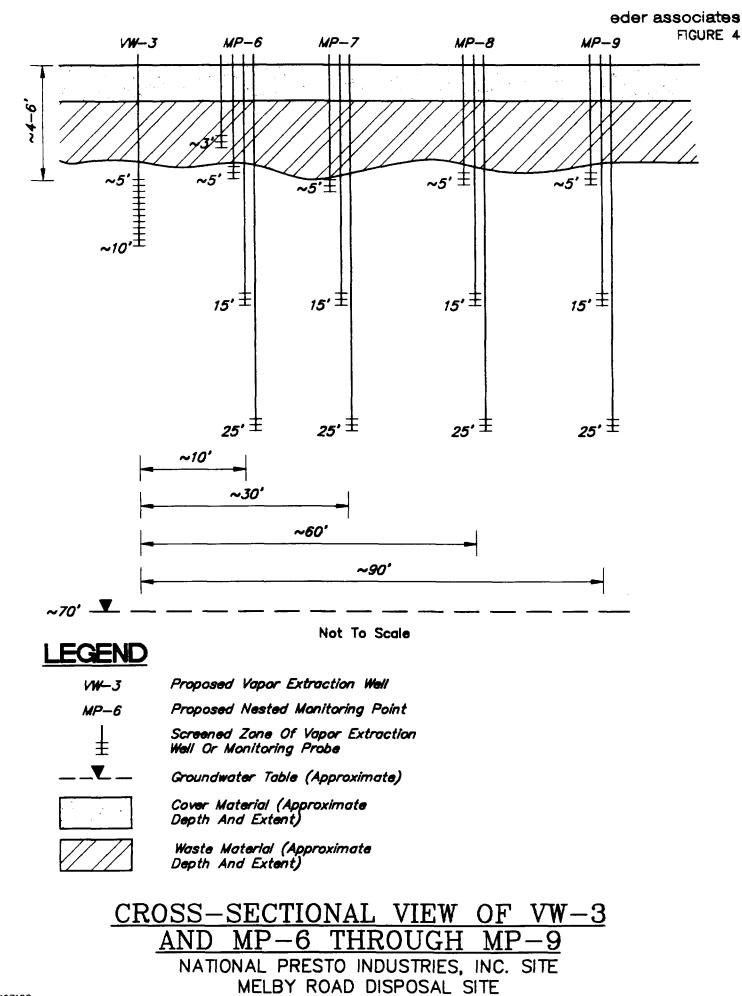
CROSS-SECTIONAL VIEW OF VW-1. VW-2, AND MP-1 THROUGH MP-5NATIONAL PRESTO INDUSTRIES, INC. SITE MELBY ROAD DISPOSAL SITE EAU CLAIRE, WISCONSIN

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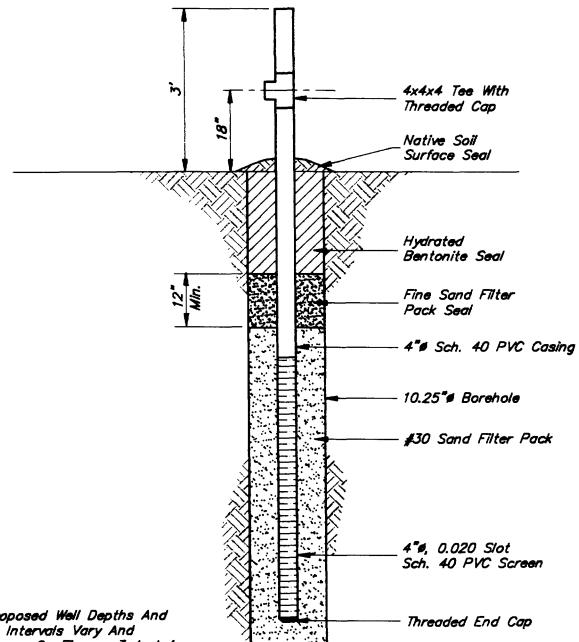
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EAU CLAIRE, WISCONSIN

iK 49718G) 31694

eder associates FIGURE 5

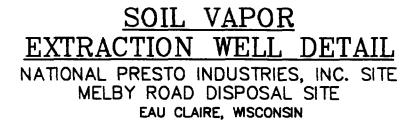


NOTE:

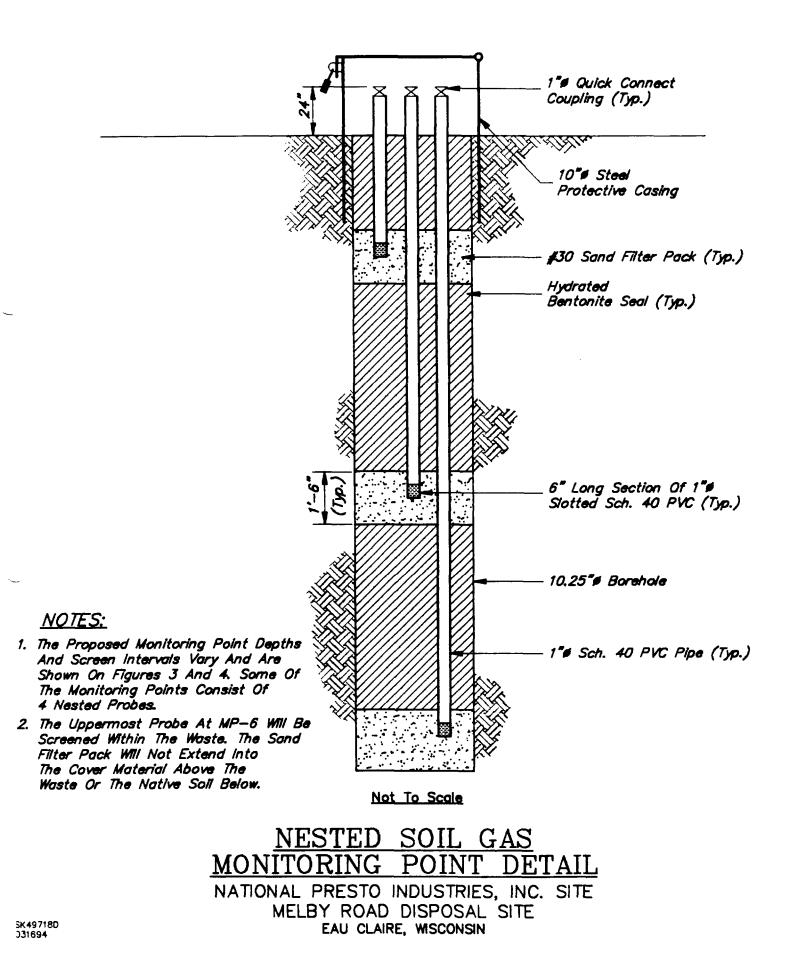
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The Proposed Well Depths And Screen Intervals Vary And Are Shown On Figures 3 And 4.

Not To Scale



eder associates FIGURE 6



PROPOSED SOIL VAPOR EXTRACTION PILOT TEST SCHEDULE NATIONAL PRESTO INDUSTRIES, INC. SITE MELBY ROAD DISPOSAL SITE EAU CLAIRE, WISCONSIN PROJECT TIME SCHEDULE (WEEKS)																										
														CHE	DUL	 	 ;)	 								
TASKS1. DEVELOP DRAFT SVE PILOT TEST WORK PLAN2. USEPA AND WDNR REVIEW OF DRAFT WORK PLAN3. REVISE WORK PLAN3. REVISE WORK PLAN4. USEPA AND WDNR OF REVISED WORK PLAN5. FINALIZE WORK PLAN6. USEPA AND WDNR APPROVAL OF WORK PLAN7. SOLICIT CONTRACTOR BIDS8. INSTALL SVE PILOT TEST SYSTEM9. CONDUCT SOIL GAS PERMEABILITY TESTS10. EVALUATE DATA			RY 4 31		BRU				ARC					95 ·	2 9	 1AY 6 2	06	JUN 3 20		7		-	25 1	-	UGL	JST 5 22
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5. FINALIZE WORK PLAN																										
6. USEPA AND WONR APPROVAL OF WORK PLAN		┼╌┼										1														
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10. EVALUATE DATA												Τ														
11. CONDUCT EXTENDED SVE TESTS																										
12. EVALUATE PILOT TEST DATA AND PREPARE DRAFT SUMMARY REPORT																										

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APPENDIX A

PORTABLE GAS CHROMATOGRAPH OPERATION PROCEDURES

PORTABLE GC OPERATION PROCEDURES

Preparation of Calibration Standards

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Cylinders of certified calibration gas mixtures are widely used as gas standards for the Photovac Model 10S50 Portable Gas Chromatograph (GC). These mixtures are generally stable and accurate to within a \pm 5 percent range. Cylinders are available in a variety of sizes. The gas standards will be purchased from Matheson Gas Products, Inc., Twinsburg, Ohio, or others.

Gas standards can also be prepared by diluting saturated headspace vapor above pure liquid compound in a static volume container containing Ultra Zero air. This method is efficient, and considerably less expensive but more time-consuming than purchasing cylinders of gas standards. The apparatus and steps required to prepare gas standards with pure chemicals are outlined below.

1. Apparatus and Reagents

- (a) Ultra Zero air
- (b) One-liter Tedlar air bag
- (c) Gas-tight syringe
- (d) Adjustable micropipette with disposable tips
- (e) 5 ml mini-vial with a Teflon faced septum cap
- (f) ANTOINE equation for estimating vapor pressure at various temperature

2. Procedure

(a) Spike two ml of a pure compound into a glass mini-vial with a Teflon faced septum cap. Allow about 40 minutes to form an equilibrated headspace vapor above the liquid. (b) Record the ambient temperature and use the ANTOINE equation to calculate vapor pressure of each compound:

 $\log P^{\circ} = A - B/(C + T)$

where, P° = Compound vapor pressure, Kpa T = Ambient temperature, °C, and A, B, C are characteristic constants of the compound.

For benzene, for example, the values of A, B, and C are 6.01905, 1204.637, and 220.069, respectively.

(c) Calculate the volume of headspace vapor required to prepare a certain vapor concentration in ppmv for a known volume of Ultra Zero air diluent:

 $V_{h} = (101.325/P^{\circ}) \ge C \ge V_{L}$

where, $V_h = V$ olume of saturated headspace, μl C = Desired concentration, ppmv, and $V_L = V$ olume of the static volume dilution container = one liter.

(d) Fill a clean one-liter Tedlar bag with the Ultra Zero air. Inject appropriate volume of the saturated vapor, as calculated in Step (c), into the Tedlar bag with a gas-tight syringe. Be careful not to touch the liquid with needle of the syringe when withdrawing the vapor from the mini-vial containing the compound. Since such precise volume cannot be measured, record exact volume injected and back calculate the final concentration.

(e) Repeat Steps (a) through (d) for all compounds of interest. Allow approximately 30 minutes for standards to reach equilibrium.

System Calibration

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The limit of detection for each compound should be determined in advance. At least two successive serial dilutions of working stock are performed to bring the standard into the low ppb range. Once a limit of detection has been established, a calibration standard containing all of the compounds of interest at suitable concentrations will be prepared.

The concentration levels of compounds in the mixed standard are such that the peaks of all compounds are clearly visible during a single analysis. An off-scale peak will be adjusted in concentration so that they are in a half height of a full-scale appearance.

Instrument Set-Up

The instrument settings will be determined by the sensitivity of the detector to the compounds, the compound's elution rate, and the detection limit required. Variables that need to be controlled during the instrument set-up include carrier gas flow rate which is typically between five and 15 ml per minute, column temperature which is typically in the range of 30 to 50 °C, analysis cycle which is typically from two minutes up to one hour, signal gain setting which is typically set between 20 to 100, and retention time window which is typically set to two to ten percent. The settings for all of these parameters (except retention time window) can be printed out with each chromatogram.

The chromatogram column used in the Photovac 10S50 is a Cp-Sil5CB column. This is a wide bore capillary column coated with a two micrometer stationary phase of 100 percent dimethyl polysiloxane. Other columns are available for analyzing special groups of compounds. The carrier gas used is hydrocarbon-free (Ultra Zero) air.

Daily Operation and Maintenance

The carrier gas flow rate will be set using a dual flowmeter with Teflon tubing. After the instrument and the internal column oven have been allowed to stabilize (approximately 30 minutes), an Ultra Zero air blank will be run to determine if the column condition is acceptable. If the column appears to be contaminated, the oven temperature and the carrier gas flow rate need to be increased to purge the column until the contamination is no longer detected.

When the GC has been warmed up and a stable baseline has been achieved, 40 to 100 μ l of the calibration standard headspace will be injected. The appropriate GC library will be updated with new retention times and peak areas of the compounds. A standard will be run every ten samples, or more frequently to ensure continued calibration of the portable GC.

A GC blank will be run at the start of each day's operation to ensure a steady baseline. This is done by starting GC analysis without injecting a sample. Blank or syringe blanks will also be analyzed following the completion of the standard calibration to determine if there is carry-over from previous samples and/or standards. The syringe blanks should be analyzed following all highly contaminated samples. Contaminants detected in the gas-tight syringe will be cleaned using methanol and distilled water.

Air samples are taken through Teflon tubing into a one-liter Tedlar bag. A volume of 40 to 100 μ l of air is withdrawn with a gas-tight syringe and the sample is directly injected into the GC and analyzed.

Care should be taken not to inject liquid into the GC. If a sample is highly contaminated, the injecting volume can be reduced accordingly. The dilution factor can then be calculated by the ratio of injected sample volume to injected calibration volume.

Quality Control

In addition to the blank analysis and calibration analysis, duplicate analysis will be run once every 20 samples or as necessary. The calibration analysis will be performed twice a day or as frequently as necessary. The portable GC calibration will be checked at the end of each day. The retention time of each peak in the chromatogram shall be checked for peak drift to avoid misidentification.

The integrity of the gas-tight syringe will be maintained. Once the syringe plungers loosen up after prolonged use, the Teflon plunger tip will be replaced. The injection port septa in the GC will be changed regularly (typically after 20 injections) to reduce contamination and to prevent air leaking which could affect the integrity of the GC results.

References

- Boublik, T., V. Fried, and E. Hala 1984. The Vapor Pressure of Pure Substances. (2nd edition), Elsevier Science Publication. The Netherlands, 1984.
- PHOTOVAC Inc. 1990. Calculated Headspace Volumes for Preparation of Vapor Standards using Pure Chemicals. PHOTOVAC Technical Bulletin # 21.

APPENDIX B

PARAMETER ESTIMATION METHOD FOR SOIL GAS PERMEABILITY TEST

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The field drawdown method is based on Darcy's Law and equations for steady-state radial flow to or from a vent well. A full mathematical development of this method and supporting calculations are provided by Johnson et al. (1990). A computer program known as HyperVentilateTM has been developed by USEPA for storing field data and computing soil permeability, k. This program will be used to evaluate the soil gas permeability test data, and will speed the calculation and data presentation process. The two solution methods for k are presented below. The first solution is based on carefully measuring the dynamic response of the soil to a constant extraction rate. The second solution for k is based on steady-state conditions and the measurement or estimation of ROI at steady state. The limitations and recommended application of each method are presented below. Whenever possible, field data will be collected to support both solution methods, because one or both of the solution methods may be appropriate, depending on site-specific conditions.

Dynamic Method

This test method requires that air be extracted at a constant rate for a single venting well, while measuring the pressure changes at several soil gas monitoring points throughout the contaminated soil volume. The equation:

$$P' = \frac{Q}{4\pi m (k/\mu)} \left[-0.5772 - \ln\left(\frac{r^2 e\mu}{4kP_{atm}}\right) + \ln(t) \right]$$
(1)

is used to describe the dynamic changes in soil gas pressure/vacuum where:

Ρ'	=	"gauge" pressure measured at distance r from the vent well at time t (g/cm-
		s ²)
m	=	stratum thickness, generally the vent well screened interval (cm)
r	=	radial distance from monitoring point to vent well (cm)
k	=	soil permeability (cm ²)
μ	=	viscosity of air $(1.8 \times 10^{-4} \text{ g/cm-s at } 18^{\circ}\text{C})$
e	=	soil effective porosity (dimension-less)
t	=	time from the start of the test (sec)
Q	=	volumetric flow rate from the vent well (cm ³ /s)
Patm	=	ambient pressure (at sea level 1.013 x 10 ⁶ g/cm-s ²)

Equation (1) predicts that the dynamic of P'-vs-ln(t) is a straight line with a slope of A where:

$$A = \frac{Q}{4\pi m (k/\mu)}$$
(2)

Solving Equation (2) for k gives

$$k = \frac{Q\mu}{4A\pi m}$$

The HyperVentilateTM model is based on the dynamic method and a determination of the slope, A. This method of determining k requires accurate field measurements of Q at the vent well and P'-vs.-time at each monitoring point. It is most appropriately applied at sites with less permeable soils where change in P' occur over a longer time period (10 minutes or more to monitoring points steady state). This method can be accurate for fine sandy soils where the screened interval extends to depths of over 10 feet and when monitoring points are screened at depths of 10 ft or greater. It is less accurate for sites where a high water table or shallow contamination limits the total depth of the vent well screen and monitoring points to less than 10 feet. In shallow and coarse-grained soils, vacuum or pressure levels reach steady state too rapidly to accurately plot P'-vs.-ln(t). Venting systems on shallow sandy sites are subject to higher vertical airflow which is not as accurately described by this one-dimensional radial flow equation.

Steady-State Method

I.

This method for determining k can be used in situations where the dynamic method is inappropriate. This method is based on the steady-state solution to Equation (1).

$$k = \frac{\mathcal{Q}\mu \ln\left(\frac{R_{w}}{ROI}\right)}{H\pi P_{w} \left[1 - \left(\frac{P_{atm}}{P_{w}}\right)^{2}\right]}$$
(3)

where Q, m, μ , and P_{aun} have been previously defined, and

R.,,	=	the radius of the venting well (cm)
Н	=	screen length (cm)
ROI	=	the maximum radius of venting influence at steady state (cm)
P _w	=	the absolute pressure at the venting well (g/cm-s ²)

The value of ROI can be determined by actually measuring the outer limit of vacuum/pressure influence under steady-state conditions, or by plotting the vacuum/pressure at each monitoring point vs. the log of its radial distance from the vent well and extrapolating the straight line to one percent of the vacuum or pressure at the vapor extraction well. The latter method will be used to estimate the ROI value to be used in calculating k. Three-dimensional modeling will be used to determine the ROI of vapor extraction wells when developing a conceptual full-scale system.

References

Johnson, P.C., M.W. Kemblowski, and J.D. Colthart, 1990, "Quantitative Analysis for the Cleanup of Hydrocarbon-Contaminated Soils by In Situ Soil Venting", Groundwater, 28(3): 413-429.

Personal Communication with P.C. Johnson, March 15-17, 1994.

APPENDIX C

EXAMPLE CHAIN OF CUSTODY AND FIELD MEASUREMENT RECORD FORMS



eder associates, consulting engineers, p.c.

65 FOREST AVENUE, LOCUST VALLEY, NEW YORK 11560 8000 EXCELSIOR DRIVE, SUITE 302, MADISON, WISCONSIN 53717 1914 315 W, HURON STREET, SUITE 220240, ANN ARBOR, MICHIGAN 48104 t--

CHAIN OF CUSTODY RECORD

LABORATORY NAME:						PROJECT NO.		0.	PROJECT NAME:			·····						
RESULIS TO:	RESULIS TO:						NO. OF	NO. OF CON- TAINERS		ANALYSIS					REMARKS			
INVOICE TO:							CON-							T	SAMPL	E DESCRIPTIONS 8		
SAMPLE I.D. NO	DATE	TIME	FLTERED	FIXED	SAMPLE LOCATION	SAMPLE METHOD			4		//				//	/	01	THER ANALYSES
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Relinquished by (Signa	iture)			/i	me Agent of:		Rec'd.			ra)		·		Date/Tin		Agent of:	···· ·	
Printed Name Relinquished by (Signature) Date/Time Agent of:			r		Printed Name					_	Date/Tin		Agent of:					
Printed Name				oc'd. by (Signature) Inted Name			-			Agent of								
Relinquished by (Signature) Date/Time Received for Laboratory by: (Signature)		Loboratory by:			/Time]	Instruc	tions:	1			I	- .					
Printed Name				1	Printed Name			1										
Sampler (Signature)					Samplers Nar	ne (Printed)						(



SOIL VAPOR EXTRACTION SYSTEM MONITORING FORM

PROJECT NO.:	SAMPLED BY:
SAMPLE I.D.:	SAMPLING POINT:
SAMPLING DATE:	SAMPLING TIME:

ANAL	YTICAL INFORMATION
ANALYTES:	
AIR FLOWRATE	
V. CUUM-PRESS	
TENDERAT URE:	
LABORATORY:	
GENERAL COMMENTS:	

SITE	TYPE OF TEST
DATE	TEST DATE
SAMPLER(S)	TIME

-

Pressure/Vacuum ("H₂O)

Distance from Vent Well (ft)						Distand Vent	æ from Well			
Time	In(t)	MP1	MP2	мрз	MP4	Time	in(t)	LIP6	MP7	MP8
							1			
•										
			5							

APPENDIX D

HEALTH AND SAFETY PLAN MELBY ROAD DISPOSAL SITE SOIL VAPOR EXTRACTION PILOT TEST

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HEALTH AND SAFETY PLAN NATIONAL PRESTO INDUSTRIES, INC. EAU CLAIRE, WISCONSIN

EDER PROJECT DIRECTOR:	Gary A. Rozmus, P.E. NAME	DATE
	Senior Vice President TITLE	
EDER PROJECT MANAGER:	William Warren NAME	DATE
	Vice President TITLE	
EDER SITE SAFETY		
OFFICER:	Darrell Dallman NAME	DATE
	Environmental Technician TITLE	

AMENDMENTS CONTAINED IN ATTACHMENT A

AMENDMENT 1. DATE _____

AMENDMENT 2. DATE _____

AMENDMENT 3. DATE _____

LLV3760

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ATTACHMENTS

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TABLES

- No. Description
- 1-1 Health and Safety Training Records
- 9-1 Field Equipment/Spill Control Equipment

GLOSSARY OF ACRONYMS

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ANSI	-	AMERICAN NATIONAL STANDARDS INSTITUTE
APR	-	AIR PURIFYING RESPIRATOR
ACGIH	-	AMERICAN CONFERENCE OF GOVERNMENTAL INDUSTRIAL
		HYGIENISTS
CFR	-	CODE OF FEDERAL REGULATIONS
CGI	-	COMBUSTIBLE GAS INDICATOR
CSEP	-	CONFINED SPACE ENTRY PERMIT
HEPA	-	HIGH EFFICIENCY PARTICULATE AIR
HNU-PID	-	HNU PHOTOIONIZATION DETECTOR
HOT ZONE	-	EXCLUSION ZONE
IDLH	-	IMMEDIATELY DANGEROUS TO LIFE & HEALTH
MREM/hr	-	MILLI-ROENTGENS EQUIVALENT IN MAN PER HOUR
NIOSH	-	NATIONAL INSTITUTE FOR OCCUPATIONAL SAFETY & HEALTH
OSHA	-	OCCUPATIONAL SAFETY AND HEALTH ADMINISTRATION
OVA	-	ORGANIC VAPOR ANALYZER
PEL	-	PERMISSIBLE EXPOSURE LIMIT
PPB	-	PARTS PER BILLION
PPE	-	PERSONAL PROTECTION EQUIPMENT
PPM	-	PARTS PER MILLION
SCBA	-	SELF-CONTAINED BREATHING APPARATUS
SOP	-	STANDARD OPERATING PROCEDURE
SPCC	-	SPILL PREVENTION CONTROLS & COUNTERMEASURES
TLV	-	THRESHOLD LIMIT VALUE
TWA	-	TIME WEIGHTED AVERAGE

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STATEMENT OF COMMITMENT TO WORKER HEALTH AND SAFETY

Eder Associates (Eder) employees may be exposed to evident or potential risk from hazardous conditions. Eder's policy is to minimize the possibility of work-related injury through aware and qualified supervision, health and safety training, medical monitoring and the use of appropriate personal protective equipment. Eder has established a worker health and safety program to protect its personnel to the maximum reasonable extent. The Corporate Health and Safety Program is documented in Appendix A of the Eder Employee Handbook, which is issued to each employee.

This site-specific Health and Safety Plan (HASP) applies to Eder personnel and others at the National Presto Industries, Inc. (NPI) site where the site operations involve employee exposure or the reasonable possibility of employee exposure to safety or health hazards. This HASP describes emergency response procedures and actual and potential chemical hazards at the work site that have been identified by Eder. This HASP does not cover the hazards from operating machinery which is the responsibility of the operating contractor. This HASP provides information and guidance to contractors retained by Eder and to other parties who are outside of Eder's ability to control. Notwithstanding the intent of this HASP as site-specific hazard information and guidance, all contractors at the site are retained as independent contractors and are responsible for assuring the work site safety of their employees and others retained by them. This HASP is made available to all parties, however, Eder can not control the actions of others and all parties enter the work site with this understanding.

Eder will require that its personnel take certain safety precautions in accord with this HASP and Eder requests that others protect their personnel in a similar manner to assure work site safety. To assure work site safety, Eder may shut down the work site and request that any party leave the work site.

1

1.0 INTRODUCTION AND SITE ENTRY REQUIREMENTS

This document describes the health and safety guidelines developed for the NPI site to protect on-site personnel, visitors, and the public from physical harm and exposure to hazardous materials or wastes. In accordance with the Occupational Safety and Health Administration (OSHA) 29 CFR Part 1910.120 Hazardous Waste Operations and Emergency Response Final Rule, this HASP, including the Attachments, addresses potential and/or actual safety and health hazards associated with each phase of site operations, except that hazards from machinery operations are addressed by the operating contractor.

This site-specific HASP is based on information available at the time the plan was prepared. The HASP may be revised following an initial site visit by the Site Manager and Eder Site Safety Officer, and when new information is received or conditions change. A written amendment will document all changes made to the plan, and will be included in Attachment A. All amendments will be acknowledged by the Site Manager, Eder Project Manager, and Eder Site Safety Officer.

<u>1.1</u> Training Requirements

All personnel entering the exclusion zone or decontamination zone (Section 6.1, Work Zones) must have completed the OSHA 29 CFR 1910.120(e) training requirements.

Documentation of Eder personnel training is maintained on file, and each Eder field personnel's record of 40-Hour OSHA Training, 8-Hour Refresher Training, and if applicable, Supervisor Training certificates will be maintained in the field office.

1.2 Medical Monitoring Requirements

All personnel (including visitors) entering the exclusion zone or decontamination zone must have completed the medical monitoring requirements under OSHA 29 CFR 1910.120(f).

Documentation of medical monitoring is the responsibility of each employer. If there are additional medical monitoring requirements for this site, evidence of compliance must also be included. Documentation of Eder personnel medical monitoring is maintained on file.

1.3 Fit-Testing Requirements

All personnel (including visitors) entering the exclusion zone or decontamination zone using a negative pressure air purifying respirator must have successfully passed a qualitative respirator fit-test in accordance with OSHA 29 CFR 1910.134 or the American National Standards Institute.

Documentation of fit-testing is the responsibility of each employer. Documentation of Eder personnel fit-testing is maintained on file.

NATIONAL PRESTO INDUSTRIES, INC. SITE EAU CLAIRE, WISCONSIN

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TABLE 1-1

HEALTH AND SAFETY TRAINING RECORDS

Eder Personnel	Attendance Date 40-Hour Health and Safety Training Course	Attendance Date 8-Hour Health and Safety Training Refresher Course	Attendance Date Supervisors Training	Date of Last Physical	Date of Last Fit Test
N. Brew	06/25/93	12/04/93		08/12/93	06/25/93
J. Barish	02/05/88	12/04/93	2/27/93	09/07/93	11/21/92
K. Butler	08/27/92	12/04/93		09/14/93	05/08/93
K. McHale	06/16/89	12/04/93	02/27/93	09/02/93	11/21/92
S. O'Brien	01/15/93	12/04/93		02/01/94	01/15/93
B. Pendergast	05/04/90	12/04/93	02/27/93	09/14/93	11/21/92
T. Perotto	10/16/91	12/04/93		01/27/94	10/12/92
V. Raykin	05/04/90	12/04/93	02/27/93	08/17/93	11/21/92
K. Savo	03/26/93	12/04/93		01/27/94	03/26/93
J. Valenti	09/14/90	01/27/94	02/27/93	12/02/93	01/27/94
D. Dallman	04/02/92	04/03/93	04/03/92	04/93	04/03/93
T. Miller	02/13/92	04/03/93		03/03/93	04/03/93
C. Donais	06/22/92	04/03/93		01/93	04/03/93
J. Piper	09/90	04/03/93		02/93	04/03/93
R. Seymour	1988	04/03/93	11/20/89	02/93	04/03/93
R. Wolske	06/93			06/93	06/93
D. Olig	03/90	04/03/93		03/93	03/90

1.4 Site Safety Plan Acceptance Acknowledgement

The Eder Site Safety Officer shall be responsible for informing all personnel entering the exclusion zone or decontamination zone of the contents of this plan and will request that each person sign the Safety Plan Acknowledgment Form in Attachment B. By signing the Safety Plan Acknowledgment Form, personnel recognize the hazards associated with the site and the policies and procedures that Eder will take to minimize exposure or adverse effects.

1.5 Daily Safety Meetings

Daily safety meetings will be held to ensure that all on-site personnel understand site conditions and operating procedures, to ensure that personal protective equipment is being used correctly, and to address questions and concerns that on-site personnel may have regarding health and safety. The meetings will be led by the Eder Site Safety Officer. All personnel trained and prepared to enter the exclusion and decontamination zones will attend the daily safety meetings. The initial meeting will be detailed to explain all site safety issues to all site workers, while the time allocated for subsequent meetings will be kept to a minimum.

1.6 Key Personnel

The Eder Principal in Charge for this project is Gary A. Rozmus, Senior Vice President. The Eder Project Manger is William Warren, and the Site Safety Officer is Darrell Dallman. Any changes in Eder's key project personnel will be recorded with the HASP amendments in Appendix A.

1.7 Roles and Responsibilities

The Principal in Charge is responsible for overall project administration. The Site Safety Officer will oversee daily safety issues. Each contractor (as an employer under OSHA) is responsible for the health and safety of its employees.

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The Eder Site Safety Officer is also responsible for coordinating health and safety standards for all individuals on-site. The Site Safety Officer will meet the emergency response and hazardous materials handling training requirements of OSHA 29 CFR Part 1910.120, will have completed supervisors training, and will have appropriate experience pertinent to the on-site work. The Site Safety Officer has the authority to order any and all site work to be suspended based on safety concerns, and is responsible for:

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- the indoctrination of all personnel with regard to all of the information in this HASP and any 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 medical situations;
- coordination with the Project Manager and Site Manager regarding site safety decisions;
- maintenance of the designation between the exclusion, decontamination, and support zones;
- 4. monitoring the condition and status of on-site hazards, and maintenance and implementation of the air quality monitoring program specified in this HASP;
- 5. maintenance of records of safety problems encountered, mitigative actions taken, and documentation of any chemical exposures or physical injuries of workers.

Any person who observes safety concerns or potential hazards that have not been addressed in the daily safety meetings should immediately report observations/concerns to the Eder Site Safety Officer or other appropriate key personnel.

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2.0 SITE BACKGROUND AND SCOPE OF WORK

Site Background

Background information on the NPI site is presented in Section 1.1 of the soil vapor extraction pilot test work plan.

Scope of Work

A soil vapor extraction (SVE) pilot test will be conducted to evaluate the feasibility and effectiveness of SVE at the Melby Road Disposal Site (MRDS).

Three MRDS locations will be tested as described in the pilot test work plan. Three vapor extraction wells and nine nested soil monitoring points will be installed. A trailer mounted vacuum blower will be used to extract vapors at each test location. Subsurface pressure drops will be recorded at the monitoring points. Vapor samples will be collected from the SVE well exhaust and the soil monitoring points in accord with the procedure and schedule described in the work plan. The vapor samples will be analyzed using a portable gas chromatograph.

SITE NAME:
ALTERNATE PROCEDURES:
ALTERNATE PROCEDURES:
ALTERNATE PROCEDURES:
ALTERNATE PROCEDURES:
REQUIRED CHANGES IN PPE:
NPI PROJECT SUPERINTENDENT (DATE)
EDER HEALTH AND SAFETY OFFICER (DATE)

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3.0 HAZARD ASSESSMENT

This Hazard Assessment identifies the activity-specific hazards associated with site operations and the standard operating procedures (SOPs) that should be implemented to reduce the hazards. This section identifies general physical hazards that can be expected at most sites, and presents an analysis of documented or potential chemical hazards at the site. Every effort will be made to reduce or eliminate these hazards. Hazards that cannot be eliminated must be guarded against by use of engineering controls and/or personal protective equipment.

3.1 Activity-Specific Hazards and Standard Operating Procedures

Sample Collection

<u>Hazards</u>

- Inhalation of volatile vapors and particulate contaminants;
- Skin contact with contaminated soils, sediments, and water through cuts, abrasions, and dermal absorption;
- Heat stress; and
- Cold stress.

SOPs to Avoid Hazards

- Wear appropriate respiratory protection;
- Wear protective gloves during sampling;
- Monitor air in the breathing zone with OVA; and
- Ensure all site personnel are familiar with the symptoms of heat stress and cold stress outlined in Attachment C.

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Sampling Equipment Decontamination

<u>Hazards</u>

- Inhalation of volatile vapors and mists;
- Skin contact with contaminants from splash;
- Slipping on wet surfaces;
- Heat stress; and
- Cold stress.

SOPs to Avoid Hazards

- Wear appropriate respiratory protection if deemed necessary;
- Wear protective gloves during decontamination;
- Ensure that all wash and rinsewater are properly drained from the decontamination area;
- Monitor air with OVA; and
- Ensure all site personnel are familiar with the symptoms of heat stress and cold stress outlined in Attachment C.

Operation of Heavy Equipment

The contractor will adhere to the specific guidelines for operating heavy equipment as outlined by OSHA in 29 CFR 1926.602. The drilling contractor's general guidelines will be amended to this report prior to the start of site work. The drilling contractor will be responsible for safety around the machinery.

3.2 General Site Hazards

The following discussion is provided as additional information, although not all of these hazards may be encountered at the NPI site.

Shock-Electrocution

All electrical power must have a ground fault circuit interrupter as part of the circuit. All equipment must be suitable and approved for the class of hazard. Applicable OSHA 29 CFR 1926 Subpart K standards for use of electricity shall apply.

Fall from Heights

Work in which a fall potentially exists will be performed using appropriate ladders and/or protection (i.e. body harness and lifeline). All work at this site is expected to be conducted at the ground surface.

Cold Stress

Cold stress is a function of cold, wetness and wind. A worker's susceptibility to cold stress can vary according to his/her physical fitness, degree of acclimatization to cold weather, age, and diet. Workers shall wear several layers of clothing and adequately cover extremities. Workers shall have access to break periods and warm beverages as necessary. When feasible, personnel shall be rotated and work performed during the warmer hours of the day. A heated enclosure for workers shall be provided close to the work area if conditions warrant. All personnel routinely working on-site (including the support zone) shall be familiar with the symptoms and signs, and care associated with cold stress as discussed in Attachment C of this HASP.

Heat Stress

When the temperature exceeds 70°F and personnel are wearing protective clothing, a heat stress monitoring program shall be implemented as appropriate. Employees shall have access to break periods and beverages as necessary. All personnel routinely working on-site (including the support zone) shall be familiar with the symptoms and signs, and emergency care associated with heat stress, heat exhaustion, and heat stroke as discussed in Attachment C of this HASP.

Facial Injury and Inhalation

In accordance with 29 CFR 1910.151(c), all operations involving the potential for eye injury, splash, etc., must have approved eye wash units locally available. Protective eye wear shall be donned in Level D, when appropriate. (The full-face air purifying respirator required by Level C and the pressure demand self-contained breathing apparatus mask required by Level B serve as eye protection.)

<u>Fire</u>

Operations involving the potential for fire hazards shall be conducted in a manner such that risk will be minimized. Non-sparking tools and fire extinguishers shall be used or available as appropriate. Sources of ignition shall be removed from work areas. When necessary, explosion-proof instruments and/or bonding and grounding will be used to prevent fire or explosion.

Overhead and underground utilities shall be identified and/or inspected prior to conducting operations involving potential contact or interference.

3.3 Chemical Hazards

The NPI site Remedial Investigation (RI) indicated the presence of VOCs at the MRDS. The most commonly found VOCs were 1,1,1-trichloroethane (TCA), trichloroethylene (TCE), tetrachloroethylene (PCE), 1,1-dichloroethylene (1,1-DCE), and 1,1-dichloroethane (1,1-DCA). Air monitoring of organic vapor concentrations during drilling using either a PID or an FID (Micro Tip/HNu or Foxoboro OVA), is required. It must be assumed that benzene is being read by the field PID instruments, since calibration will be to benzene. Polynuclear aromatic hydrocarbons (PAHs) and metals are also present in waste at the NPI site. Precautions will be taken to protect against inadvertent ingestion of contaminants during the pilot test.

A summary table of these chemicals containing pertinent health and safety information is contained in Attachment D. As additional chemicals are identified, Attachment D will be supplemented with appropriate information.

4.0 PERSONAL PROTECTIVE EQUIPMENT

The selection of personal protective equipment (PPE) shall be conducted in accordance with the site air monitoring program, OSHA 29 CFR 1910.120(c) and (g), and 1910.132. Protective equipment shall be NIOSH-approved and its use for 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.

4.1 Activity-Specific Levels of Personal Protection

The required level of PPE is specific to the activity being conducted, and is based on air monitoring results (Section 5.0) and properties of identified contaminants and contaminants expected to be encountered (Section 3.3).

The work at the MRDS will be initially performed in Level D. The air monitoring data collected during the August 1993 soil boring work conducted at the MRDS showed organic vapor readings on the order of 100 to 1000 ppm for some of the split spoon samples collected from the borings, however none of the readings in the breathing zone were sustained above background levels. Air monitoring will be conducted continuously and recorded to determine whether Level C or Level B is required, or if Level D conditions are suitable. If during drilling or sampling activities, air monitoring in the breathing zone indicates sustained VOC readings above background but less than 5 ppm above background, the field team must shut down work or upgrade PPE to Level C. If air monitoring indicates sustained VOC readings in the breathing zone above background, the field team must shut down work or upgrade PPE to Level B. A gas chromatograph may be used to identify specific compounds present in the breathing zone. If the identified compounds have Level C PPE capability, then work can be conducted in Level C. If

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this is done, then air samples for gas chromatograph analysis will be collected 4 times a day to verify that the condition is sustained, or as determined by the Site Safety Officer.

4.2 Level D

Level D PPE will be used when atmospheric conditions permit and the work precludes splashes, immersion or the potential for unexpected contact with harmful chemicals. Level D PPE consists of:

- Standard work uniform or coveralls (or tyvek, as needed);
- Steel toe and steel shank work boots;
- Hard hat;
- Gloves as needed; and
- Safety glasses as needed.

<u>4.3 Level C</u>

Level C PPE shall be donned when sustained concentrations of known total organic vapors in the breathing zone exceed background concentrations but are less than 5 ppm above background using a portable PID or FID. The compounds present will be determined using a gas chromatograph. The air purifying filter cartridges must be appropriate for contaminants identified or expected to be encountered. Level C PPE shall be donned when the contaminant identified has adequate warning properties and criteria for the use of APR have been met. The appropriate PPE level will be established by the Site Safety Officer. Unknowns are adequately defined as judged by the Site Safety Officer. Level C PPE consists of:

- Chemical resistant or coated tyvek coveralls;
- Steel toe and steel shank work boots;
- Chemical resistant overboots or disposable boot covers;
- Disposable inner gloves (surgical gloves);

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- Disposable outer gloves;
- Full-face APR fitted with organic vapor/dust and mist filters or filters appropriate for the identified contaminants expected to be encountered;
- Hard-hat;
- Splash shield, as needed; and,
- Ankles/wrists taped with duct tape.

4.4 Level B

Level B PPE shall be donned when sustained concentrations of measured total organic vapors in the breathing zone are greater than 5 ppm above background using a portable PID or FID. Level B PPE shall be donned if the IDLH of a known contaminant is exceeded. If a contaminant is identified or is expected to be encountered for which NIOSH and/or OSHA recommend the use of a positive pressure self-contained breathing apparatus (SCBA), Level B PPE shall be donned, even though the total organic vapors in the breathing zone may not exceed background by 5 ppm. If Level B PPE is required for a task, then at least two Eder personnel shall wear Level B at all times during the performance of that task and a third person will remain out of the work zone on standby with Level B equipment should a problem arise. Level B PPE consists of:

- Chemical resistant coveralls;
- Steel toe and steel shank work boots;
- Chemical resistant over boots or disposable boot covers;
- Disposable inner gloves;
- Disposable outer gloves;
- Supplied air SCBA or airline system with 5-minute egress system;
- Hard-hat; and,
- Ankles/wrists taped.

5.0 AIR MONITORING AND ACTION LEVELS

According to 29 CFR 1910.120(h) air monitoring shall be employed to identify and quantify airborne levels of hazardous substances and health hazards.

5.1 Routine Air Monitoring Requirements

Air monitoring using a either a portable PID or FID (Micro Tip/HNu or Foxoboro OVA) and a combustible gas indicator/oxygen meter shall be used when any of the following conditions apply:

- Initial site entry;
- The possibility of an IDLH condition or flammable atmosphere has developed;
- Work begins on a different portion of the site;
- Contaminants other than those previously identified have been discovered;
- A different task or activity is initiated;

All air monitoring data will be documented in a site log book. Air monitoring instruments will be calibrated and maintained in accordance with the manufacturer's specifications.

The following site entry and monitoring guidelines regarding actions to be taken based on routine air monitoring shall be applied. These are:

- CGI readings < 10% LEL: continue.
- CGI readings of 10 to 20% LEL: proceed with caution.
- CGI readings > 20% LEL: stop work.
- Organic vapor readings in breathing zone sustained at background: continue in Level D.

- Organic vapor readings in breathing zone exceed background but are less than 5 ppm above background: Level C PPE.
- Organic vapor readings in breathing zone sustained at levels greater than 5 ppm above background: Level B PPE or stop work.

The compounds present may be determined using a gas chromatograph. If air purifying filter cartridges are appropriate for the identified contaminants, Level C may be used when the contaminants identified have adequate warning properties and criteria for the use of APR are met. If this is done, then air samples for gas chromatograph analysis must be collected 4 times a day to verify that the conditions are sustained, or as determined by the Site Safety Officer.

Real-time measurements of organic vapor concentrations will be made using a PID or a FID. Air monitoring locations will be at four locations on the perimeter of the exclusion zone and at the site boundary (one location upwind and two locations downwind). All monitoring data will be recorded in the field notebook.

Real-time air sampling can be conducted at locations in addition to those specified in this HASP as conditions warrant. For example, if sustained elevated organic vapor concentrations are measured near the work area, additional readings will be taken as the air sampling personnel moves away from the work area in the downwind direction towards the NPI site property line (the Melby Road fence line) to evaluate the dissipation of contaminant concentrations in the air.

6.1 Work Zones

The Eder Site Safety Officer shall designate an exclusion zone, a decontamination zone, and a support zone in step with the work being performed.

The exclusion zone is the area within which tasks requiring the OSHA 40-hour Hazardous Waste Operations and Emergency Response Operations training are carried out.

The removal of protective equipment shall occur within the designated decontamination zone. Disposable protective equipment shall be stored in receptacles kept in the decontamination zone, and non-disposable equipment will be decontaminated according to the procedures outlined in Section 7.0. All personnel and equipment will exit the exclusion zone via the decontamination zone. First aid equipment, an eye wash unit, and drinking water shall be kept in the decontamination trailer.

The support zone will be used for the command post, for vehicle parking, daily safety meetings, and supply storage. No decontamination will be permitted in the support zone. This HASP, the HASP attachments, a site map indicating the three work zones, and a telephone will be kept at the command post.

6.2 General Field Safety and Standard Operating Procedures

It is Eder's policy to practice administrative hazard control for all site areas by restricting entrance to exclusion zones to essential personnel, and by implementing SOPs.

• Personnel not specifically authorized to enter the exclusion zone will remain in the support zone.

- Prior to entering the exclusion or decontamination zones, all personnel must be familiar with emergency incident procedures (Section 9.0); the locations of site safety, first aid and communication equipment; the directions to the hospital; and the list of emergency telephone numbers.
- The "buddy system" will be used at all times by all field personnel in the exclusion zone. No one is to perform field work alone. When in Level C, visual contact or radio contact should be maintained at all times. When in Level B, visual contact should be maintained at all times, and radio contact should be maintained with the decontamination or support zone.
- Whenever possible, avoid contact with contaminated and potentially contaminated surfaces. Walk around (not through) puddles and discolored surfaces. Do not kneel on the ground or place equipment on the ground. Protect equipment from contamination.
- All personnel exiting the exclusion zone must exercise the decontamination procedures described in Section 7.0 of this HASP.
- Facial hair that interferes with respirator fit will preclude admission to the exclusion zone. Contact lenses shall not be worn in the exclusion or decontamination zones, or if the worker may be expected to enter these zones under routine or emergency situations.
- Eating, drinking, or smoking is permitted only in designated areas in the support zone.
- Each worker must be supplied with and maintain his/her own PPE.

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7.0 DECONTAMINATION PROCEDURES

In general, everything that enters the exclusion zone must either be decontaminated or properly discarded upon exit from the exclusion zone. All personnel must enter and exit the exclusion zone through the decontamination area. Due to the nature of the work to be conducted, the exclusion and decontamination zones may "float". Bags used to discard disposable personal protective clothing and equipment will be carried from location to location.

All boots and other potentially contaminated garments which have come in contact with waste material and contaminated soil will be cleaned with detergent/water solution and rinsed with water in wash tubs. The wash water, rinse water and residues will be collected and properly stored until sampling results are received and final disposition of the waste can be determined. Disposable PPE, including spent respirator cartridges and canisters, will be properly bagged and disposed of. All contaminated boots, clothing, and equipment (eg. leather boots, equipment carrying straps) which cannot be decontaminated will be disposed of with the disposable garments.

The minimum measures for Level B removal and decontamination are:

- Deposit equipment on plastic drop cloths;
- Scrub outer boots and gloves with a solution of water and detergent and rinse off;
- Remove outer boots and outer gloves. Dispose of any disposable outer garments in waste receptacle provided;
- Remove tyvek/outer garment and place in receptacle provided;

- Remove inner gloves and deposit in receptacle provided;
- Remove SCBA and face piece and place on rack provided; and,
- Full shower including washing hair, face and hands.

The minimum measures for Level C removal and decontamination are:

- Deposit equipment on plastic drop cloths;
- Scrub outer boots and gloves with a solution of water and detergent and rinse off;
- Remove outer boots and outer gloves. Dispose of any disposable outer garments in receptacle provided;
- Remove tyvek/outer garment and place in receptacle provided;
- Remove first pair of inner gloves;
- Remove respirator (with "clean" inner gloves) and place on rack provided;
- Remove last pair of inner gloves and deposit in receptacle provided; and,
- Full shower including washing hair, face and hands.

The second to last item to be removed should be the APR, and the last item to be removed should be the last of several pairs of surgical gloves. Wearing several pairs of inner gloves permits layers to be removed as needed during various stages of the doffing procedure, and, in the event that the APR has inadvertently become contaminated, wearing inner gloves to remove the APR guards against bare hands contacting the APR.

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8.0 CONFINED SPACE

In general, a confined space is defined as a space or work area not designed or intended for normal human occupancy, with limited means of access and poor natural ventilation. Confined space entry is not anticipated at the NPI site. In the event a confined space entry is to be necessary, the HASP will be amended and the requirements for a confined space entry will be followed, as per OSHA standard 29 CFR 1910.146.

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9.0 EMERGENCY RESPONSE CONTINGENCY PLAN

It is essential that site personnel be prepared for an emergency. Emergencies can take many forms; illnesses or injuries, chemical exposure, fires, explosions, spills, leaks, releases of harmful contaminants, or sudden changes in the weather.

The list of emergency telephone numbers and directions to the hospital will by posted in the command post. Site personnel should be familiar with the emergency incident procedures, and the locations of site safety, first aid, and communication equipment.

9.1 Emergency Equipment On-Site

Private Telephones:	Eder mobile phone. Nearest residence.
Two-Way Radios:	Eder site personnel.
Emergency Alarms	On-site vehicle horns [•] .
First Aid Kits:	On-site Eder vehicle.
Fire Extinguisher:	On-site Eder vehicle, work zones.**

* Horns: Air horns will be supplied to personnel at the discretion of the Site Manager or Site Safety Officer.

****** Work Zones - There must be a chemical fire extinguisher present in the hot zone where Level B PPE is worn.

9.2 Emergency Telephone Numbers and Hospital Information

	EMERGENCY NUMBER	NON-EMERGENCY
Eau Claire Fire Department	911	839-5012
Eau Claire Police Department	911	839-4972
Sacred Heart Hospital (Trauma Center	r) 839-4222	839-4222
Luther Hospital	839-3242	839-3242
(Poison Control Center)	835-1515	835-1515
National Response Center	800-424-8802	800-424-8802

Emergency Route to Local Hospitals

Sacred Heart Hospital 900 West Clairemont Avenue Eau Claire, WI

Directions: Take Highway 53 south to Clairemont Avenue (approx 5 miles). Turn right (west) on Clairemont Avenue and continue approximately one mile to hospital.

Luther Hospital 1221 Whipple Street Eau Claire, WI

Directions: Take Highway 53 south to Main Street. Go right (west) on Main Street to Farewell Street. Go left (south) on Farewell Street to Lake Street. Go right on Lake Street over the bridge to Fifth Avenue. Make right on 5th Avenue to Chestnut Street. Go left on Chestnut Street and hospital is on the corner.

POST A COPY OF THIS PAGE IN THE OFFICE TRAILER FIELD VEHICLE

9.3 Personnel Responsibilities During an Emergency

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As the administrator of the project, the Project Manager has primary responsibility for responding to and correcting emergency situations. In the absence of the Project Manager, the Site Safety Officer shall act as the Project Manager's on-site designee. Their responsibilities include:

- Take appropriate measures to protect personnel including: exit from the exclusion zone, total evacuation and securing of the site or up-grading or down-grading the level of protective clothing and respiratory protection;
- 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. In the event of an air release of toxic materials, the local authorities should be informed in order to assess the need for evacuation;
- Ensure that appropriate decontamination treatment or testing for exposed or injured personnel is obtained;
- Determine the cause of the incident and make recommendations to prevent the recurrence; and,
- Ensure that all required reports have been prepared.

9.4 Medical Emergencies

Any person who becomes ill or injured in the exclusion zone must be decontaminated to the maximum extent possible. If the injury or illness is minor, full decontamination should be completed and first aid administered prior to transport. First aid should be administered while waiting for an ambulance or paramedics. Any person transporting an injured/exposed person to

a clinic or hospital for treatment should take the directions to the hospital and information on the chemical exposure.

9.5 Fire or Explosion

In the event of a fire or explosion, the fire department should be summoned immediately. Upon their arrival, the project manager or designated alternate will advise the fire commander of the location, nature and identification of the hazardous materials on-site. The contractor shall have access to stockpiled sand which will be used to extinguish fires by smothering them, prior to arrival of the fire department.

9.6 Evacuation Routes

Evacuation routes established by work area locations for this site will be highlighted on a site map and periodically reviewed during the daily safety meetings. As the work areas change, the evacuation route and map will be updated accordingly, and the new route will be reviewed during the daily safety meetings.

Under conditions of extreme emergency, evacuation should be conducted immediately and without regard for equipment. The evacuation signal will be a continuous blast of a vehicle horn, if possible, and/or by verbal/radio communication. All site personnel shall:

- Keep upwind of smoke, vapors or spill location.
- Exit through the decontamination corridor if possible.
- If evacuation via the decontamination corridor is not possible, site 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 Project Manager, Site Manager, or Site Safety Officer will conduct a head count to assure all personnel have been evacuated safely. The head count will be verified with the site and/or exclusion zone entry/exit log.
- In the event that emergency site evacuation is necessary, all personnel are to escape the emergency situation and decontaminate to the maximum extent practical.

9.7 Spill Control Procedures

In the event of a leak or a release, on site personnel will:

- Inform their supervisor immediately;
- Locate the source of the spillage and stop the flow if it can be done safely; and,
- Begin containment and recovery of the spilled materials. Equipment on-site shall be sufficient to handle any spills that may occur. Equipment shall be diked and containerized appropriately.

No guests, owner's representatives, or other non-trained on-site personnel shall respond to an emergency as outlined above, but shall immediately evacuate the site in such an event.

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TABLE 9-1

FIELD EQUIPMENT/SPILL CONTROL EQUIPMENT

Equipment	Use (Zone)
Steam Cleaner	Decontamination - interface between hot and cold zone
Mobile-Decontamination Unit Personnel Hygiene Trailer	Decontamination - field office
MicroTip	Exclusion zone - air monitoring
ονΑ	Exclusion zone - air monitoring
SA-CGI	Exclusion zone - air monitoring
Mobile phone	Cold zone - communication
Hand held radios	Exclusion zone - communication
Absorbent pads	Drum Sampling ARCA - Drum storage roll-off container

ATTACHMENT A

SITE SAFETY PLAN AMENDMENTS

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SITE SAFETY PLAN AMENDMENT #		
SITE NAME:		
REASON FOR AMENDMENT:		
ALTERNATE PROCEDURES:		
REQUIRED CHANGES IN PPE:		
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NPI PROJECT SUPERINTENDENT (DATE)		
EDER HEALTH AND SAFETY OFFICER	(DATE)	
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TE NAME:		
EASON FOR AMENDMENT:		
LTERNATE PROCEDURES:		
EQUIRED CHANGES IN PPE:		
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ATTACHMENT B

SITE SAFETY ACKNOWLEDGMENT FORM

SITE SAFETY PLAN ACKNOWLEDGMENT FORM

I have been informed and understand the procedures set forth in the HASP and Amendments for the NPI Soil Vapor Extraction Pilot Test Project.

Printed Name	Signature	Representing	Date
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ATTACHMENT C

HEAT STRESS/COLD STRESS

HEAT RELATED EMERGENCIES

Good judgment is essential. Pace yourself by knowing your limitations. Avoid over exertion. You are your best gauge for heat related emergencies. When in doubt, get out!

HEAT EXPOSURE

The human body stubbornly defends its constant core temperature of 99.6°F. To maintain this constant temperature, heat loss must equal heat gain. If heat loss exceeds heat gain, the body temperature will fall; conversely, if heat production exceeds heat loss, the temperature will rise. In a heat related emergency, the body's mechanisms for temperature regulation are overwhelmed. The body can no longer regulate core temperature, and the core temperature begins to rise. As this rise occurs, the body will begin to show the signs and symptoms of heat related emergencies. The sequence of illness may start with heat Cramps and progress into a more severe case or may go straight to Heat Stroke. The degree of illness will vary from person to person, depending on the nature of the exposure, physical conditioning and inherited traits.

PREVENTION

General

- Maintain good physical conditioning and control your blood pressure (avoid weight gain, smoking, etc.).
- Eat regularly and properly. Increase salt intake through food consumption during the hot season or hot spells and avoid the use of salt tablets.
- Regulate alcohol intake if you are going to be working in hot environments, either from ambient conditions or through the wearing of Chemical Protective Clothing.
- Obtain basic First Aid and CPR training.
- If you are on medication or have a chronic medical history, consult a physician prior to working in a hot environment.

On-Site/Scene

- If you anticipate field work, get acclimated and conditioned prior to working in high temperatures.
- Sufficient quantities of water should be consumed to help avoid heat related emergencies.

HEAT RELATED EMERGENCIES SIGNS AND SYMPTOMS EMERGENCY CARE

The work will involve the wearing of PPE at Level B and heat will have a great influence. slowing the work. If site related chemicals are such that splash conditions are unlikely than the site safety officer may downgrade the Level B dress to include cotton coveralls. The conditions of summer work dictate that special precautions be taken to ensure that heat related injuries are avoided. All field personnel are encouraged to drink plenty of liquids (i.e., gatorade or equivalent). This section presents signs and symptoms for heat related conditions and limitations on the work. Workers at the NPI Sites will be monitored for internal temperature hourly when temperatures are over 85°F.

Heat Rash

Also known as prickly heat, this is a condition affecting the skin. The condition occurs in situations where the skin remains wet most of the time. The sweat ducts become plugged, and a skin rash soon appears.

Signs and Symptoms

- 1. Skin rash over affected areas of the body.
- 2. Tingling or prickling sensation on the affected areas.

Emergency Care

- 1. Take shower after working in heat.
- 2. Dry the skin thoroughly.
- 3. Change underwear as needed.
- 4. Stay in cool place after work hours.
- 5. Avoid repeated exposure to heated environment until condition improves, when possible.

Heat Cramps

Heat cramps are muscle pains, usually in the lower extremities, the abdomen, or both, which occur secondary to profuse sweating with accompanying salt depletion. Heat cramps most often afflict people in good physical condition, who overwork in conditions of high temperature and humidity. Untreated, heat cramps may progress to heat exhaustion.

Treatment of heat cramps is aimed at eliminating the exposure and restoring the loss of salt and water.

Signs and Symptoms

1. Cramps in the extremities and abdomen which come on suddenly during vigorous activity. Heat cramps can be mild with only slight abdominal cramping and tingling in the extremities, but more commonly present intense and incapacitating pain in the abdomen and extremities.

- 2. Respiration rate will increase, decreasing after the pain subsides.
- 3. Pulse rate will increase.
- 4. Skin will be pale and moist.
- 5. Body temperature will be normal.
- 6. Loss of consciousness or airway maintenance are seldom problems with this condition.
- 7. Generalized weakness will be noted as the pain subsides.

Emergency Care

- 1. Move the worker to a cool environment. Have him lie down if he feels faint.
- 2. <u>If the worker is not nauseated</u>, he may be given 1 or 2 glasses of an electrolyte solution. Have the worker drink slowly. The use of salt tablets is not recommended, as they may precipitate nausea.
- 3. <u>If the worker is nauseated</u>, avoid giving anything by mouth until the nausea subsides.
- 4. Avoid massaging the cramping muscles. This rarely helps and may actually aggravate the pain.
- 5. As the salt and water level is replenished, the worker's pain will subside. He may wish to return to work, however, this is NOT recommended for a period of 12 hours. Further exertion may lead to heat exhaustion or heat stroke.

HEAT EXHAUSTION

Heat exhaustion represents a somewhat more severe response to salt and water loss, as well as an initial disturbance in the body's heat-regulating system. Like heat cramps, heat exhaustion tends to occur in persons working in hot environments. Heat exhaustion is likely in dehydrated and hypertensive people. Untreated Heat Exhaustion may progress to Heat Stroke.

Treatment of heat exhaustion is similar in principle to that of heat cramps.

Signs and Symptoms

- 1. Heat Exhaustion may come on suddenly or may be present with a headache, fatigue, dizziness, nausea with occasional abdominal cramping.
- 2. Sweating will be profuse.
- 3. Pulse rate will be rapid and weak.
- 4. Respiration rate will be rapid and shallow.
- 5. The skin will be pale and clammy.
- 6. The body temperature will be normal or decreased.
- 7. The worker could be irritable and restless.
- 8. Monitor the worker's level of consciousness and airway.

Emergency Care

- 1. Move the worker to a cool environment, take off as much of his clothing as possible, and place him in a supine position with his legs elevated.
- 2. Sponge the worker with cool water. If you fan the worker, avoid chilling. When the body chills, the muscles generate energy. When the body shivers, this energy is released in the form of heat and actually can increase the body temperature.
- 3. If this is a true medical emergency, prompt intervention by Emergency Medical Services is recommended.

HEAT STROKE

Heat Stroke is caused by a severe disturbance in the body's heat-regulating mechanism and is a profound emergency, with a mortality rate ranging from 25 to 50 percent. It is most common in men over 40, especially in alcoholics. It can also occur in people of any age having too much exposure to the sun or prolonged confinement in a hot atmosphere. Heat stroke comes on suddenly. As the sweating mechanism fails, the body temperature begins to rise precipitously, reaching 106°F (41°C) or higher within 10 to 15 minutes. If the situation is not corrected rapidly, the body cells - especially the very vulnerable cells of the brain - are literally cooked, and irreversible central nervous system damage occurs.

The treatment for Heat Stroke is aimed at maintaining vital functions and causing as rapid a temperature fall as possible.

Signs and Symptoms

- 1. The worker's pulse will be strong and pounding.
- 2. The skin will be hot, dry and flushed.
- 3. The worker may experience headache, dizziness, and dryness of mouth.
- 4. Seizures and coma occur.
- 5. Loss of consciousness and airway maintenance problems can occur.

Emergency Care

- 1. Establish an open airway.
- 2. Move the worker to a cool environment. Take off as much clothing as possible, and place him in a semi-reclining position with the feet elevated.
- 3. Use any means to cool the worker. Improvise with whatever is available. A bathtub filled with cold water and ice cubes is ideal. Remember, speed is essential; delay may result in permanent brain damage. Vigorous efforts to cool the worker must continue until the body temperature is below 103°F (38.9°C).
- 4. This is a true medical emergency; prompt intervention by Emergency Medical Services is recommended.

These are only guidelines for the care of Heat Related Emergencies. Actual training in emergency medical care or basic first aid is recommended.

HEAT STRESS

1. Heart rate (HR) should be monitored by the radial pulse for 30 seconds as soon as possible in the resting period.

If at the beginning of the rest period a worker's radial pulse is measured and his heart rate exceeds 100 beats per minute, the worker's next work period should be reduced by 33%. Therefore, if the original work period was one hour, the following work cycle should be reduced to 40 minutes.

- 2. Administering salt tablets to prevent heat stress is not recommended due to a number of reasons: (a) sweat is hypotonic, therefore, adding salt to the body would only increase the body's need for water; (b) additional salt may interfere with a worker's predisposed physical condition (i.e., high blood pressure); and (c) increasing the sodium content in the body may cause an imbalance in the body's potassium content. Unless a physician recommends the use of soil tablets, individuals naturally obtain the necessary salt in their normal diet.
- 3. Heat Stroke is a true medical emergency. First aid should be directed toward immediate measures to cool the body quickly, as well as seeing that the victim receives medical attention as soon as possible.

Prior to medical treatment, remove as much clothing as possible and proceed to cool the victim's body, taking care not to overchill the victim once his temperature falls below 102°F. One of the following cooling measures should be taken: (1) sponge the bare skin with cool water; (b) apply cold packs continuously; (c) wrap the victim in a sheet soaked with water; or (d) immerse the victim in a tub of cold water, while closely monitoring the victim's level of consciousness.

4. Prior to site activity, the field TEAM leader may make arrangements for heat stress monitoring (i.e., monitoring heart rate, body temperature and body water loss) during actual site work if conditions warrant these measures. In addition, he would want to ensure that the team members have been acclimatized to the particular environmental conditions and that personnel are aware of the signs and symptoms of heat illness and have been adequately trained in first aid procedures. As field team leader, one could also make sure there is sufficient personnel on site, so as to rotate work assignments, schedule work during hours of reduced temperatures, and ensure personnel do not consume alcoholic or caffeinated beverages but rather drink moderate levels of an electrolyte solution and eat well prior to commencing site work.

5. The worker could be experiencing a condition of heat rash. Allow workers to rest and relieve the itching associated with heat rash rather than return to work too soon. Itching workers may not follow stringent decon procedures or scratch where it itches on-site and risk cross contamination.

Keeping the skin clean and dry will reduce the incidence of heat rash. This can be accomplished by wearing cotton garments (or other materials that absorb perspiration) underneath protective clothing. Upon removal of the protective clothing, the worker should wash and dry his skin thoroughly.

- 6. The sense of thirst is not an adequate regulator of water replacement during heat exposure. Therefore, as a general rule, the amount of water administered should replace the amount of water lost, and it should be administered at regular intervals throughout the day. For every ½ pound of water loss, 8 ounces of water should be ingested. Water should be replaced by drinking 2-4 ounce servings during every rest period. A recommended alternative to water is an electrolyte drink split 50/50 with water.
- 7. Although there is no specific test given during a baseline physical that would identify a person's intolerance to heat, there are physical factors and personal habits which may indicate possible intolerance to heat, such as, whether or not an individual smokes, one's dietary habit, body weight, as well as predisposed physical conditions such as high blood pressure, heart conditions, diabetes, or one's medication, that may influence an individual's ability to tolerate excessive heat.
- 8. Heat cramps are caused by profuse perspiration with inadequate fluid intake and salt replacement. Heat cramps most often afflict people in good physical condition who overwork in conditions of high temperature and humidity. Heat cramps usually come on suddenly during vigorous activity. Untreated, heat cramps may progress directly to heat exhaustion or heat stroke. First aid treatment: remove victim to a cool place and give sips of salted water (1 teaspoon of salt to 1 quart of water) 4 ounces every 15 minutes over a period of one hour. A commercial preparation, e.g., Gatorade, may be used if split 50/50 with water.

The salted water or solution should mitigate the cramps. Manual pressure should not be applied to the cramped muscles.

TABLE C-1⁽¹⁾

REQUIRED FREQUENCY OF HEAT STRESS MONITORING FOR WORKERS IN IMPERMEABLE CLOTHING

Adjusted ⁽²⁾ Temperature (°F)	Work Time Allowed Before Monitoring Break (min.)
90 or above	15
87.5-90	30
82.5-87.5	60
77.5-82.5	90
72.5-77.5	120

- Adapted from Eastern Research Group and National Institute for Occupational Safety and Health, <u>Occupational Safety and Health Guidance Manual for Super Activities</u>. September 26, 1984, pp. 8-75.
- (2) Calculate the adjusted air temperature (Ta adj) by using this equation:

Ta adj $^{\circ}F = Ta ^{\circ}F + (13 \times \% \text{ sunshine})$

Measure air temperature (Ta) with a standard thermometer, with the bulb shielded from radiant heat. Then estimate percent sunshine (100 percent sunshine = no cloud cover and a sharp, distinct shadow; 0 percent sunshine = no shadows).

TABLE C-2

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Heat Stress Indicator	When to Measure	If Exceeds	Action
heart rate (pulse)	beginning of rest period	110 beats per minute	shorten next work period by 33%
oral temperature	beginning of rest period	99°F (after thermometer is under tongue for 3 minutes) 100.6°F	shorten next work period by 33% prohibit work in impermeable clothing
body weight	1. before workday begins (a.m.) 2. after workday ends (p.m.)		increase fluid intake

TABLE C-3^(b)

SYMPTOMS OF HEAT STRESS

Heat rash results from continuous exposure to heat or humid air.

Heat cramps are caused by heavy sweating with inadequate fluid intake. Symptoms include:

- muscle spasms.
- pains in the hands, feet, and abdomen.

Heat exhaustion occurs when body organs attempt to keep the body cool. Symptoms include:

- pale, cool, moist skin.
- heavy sweating.
- dizziness.

<u>Heat stroke</u> is the most serious form of heat stress. Immediate action must be taken to cool the body before serious injury and death occur. Symptoms are:

- red, hot, dry skin.
- lack of perspiration.
- nausea.
- dizziness and confusion.
- strong, rapid pulse.
- coma.
- (1) Reproduced from <u>Occupational Safety and Health Guidance Manual for Superfund</u> <u>Activities</u> (see Table A-1), p. 8-79.

Cold Stress (Hypothermia)

Cold stress is a function of cold, wetness and wind. A worker's susceptibility to cold stress can vary according to his/her physical fitness, degree of acclimatization to cold weather, age, and diet.

Prevention

Institute the following steps to prevent overexposure of workers to cold:

- 1. Maintain body core temperature at 96.8°F or above by encouraging workers to drink warm liquids during breaks (preferably not coffee) and wear several layers of clothing. Wool is recommended since it can keep the body warm even when the wool is wet.
- 2. Avoid frostbite by adequately covering hands, feet, and other extremities. Clothing such as insulated gloves or mittens, earmuffs, and hat liners should be worn. To prevent contact frostbite (from touching metal and cold surfaces below 20°F), workers should wear anti-contact gloves. Tool handles and control bars should be covered with insulating material.
- 3. Adjust work schedules if necessary, providing adequate rest periods. When feasible, rotate personnel and perform work during the warmer hours of the day.
- 4. Provide a heated enclosure for workers close to their work area. Workers should remove their outer layer(s) of clothing while in the shelter to allow for sweat evaporation.
- 5. In the event that wind barriers are constructed around an intrusive operation (such as drilling), the enclosure must be properly vented to prevent the build-up of toxic or explosive gasses or vapors. Care must be taken to keep any heat source away from flammable substances.
- 6. Using a wind chill chart such as the one in Table E-4, obtain the equivalent chill temperature (ECT) based on actual wind speed and temperature. Refer to the ECT when setting up work warm-up schedules, planning appropriate clothing, etc. Workers should use warming shelters at regular intervals at or below an ECT of 20°F. For exposed skin, continuous exposure should not be permitted at or below an ECT of -25°F.
- 7. Workers who become immersed in water or whose clothing becomes wet (from perspiration, rain, etc.) must immediately be provided a change of dry clothing whenever the air temperature is 25.6°F or below.

8. Maintain an optimal level of worker fitness by encouraging regular exercise, proper diet, etc. If possible, acclimatize workers to site conditions for several days before work begins.

Monitoring

Personnel should be aware of the symptoms of cold stress. If the following symptoms of <u>systemic hypothermia</u> are noticed in any worker, he/she should immediately go the warm shelter:

- heavy, uncontrollable shivering;
- excessive fatigue or drowsiness;
- loss of coordination;
- difficulty in speaking; and,
- frostbite (see below).

<u>Frostbite</u> is the generic term for local injury resulting from cold. The stages of frostbite and their symptoms are as follows:

- 1. frostbite or incipient frostbite:
 - sudden blanching or whitening of the skin.
- 2. superficial frostbite:
 - waxy or white skin which is firm to the touch (tissue underneath is still resilient).
- 3. deep frostbite:
 - tissues are cold, pale, and solid.

TABLE C-4⁽¹⁾

COOLING POWER OF WIND ON EXPOSED FLESH EXPRESSED AS AN EQUIVALENT TEMPERATURE (UNDER CALM CONDITIONS)

		Actual Temperature Reading (°F)P										
Estimated Wind Speed (in mps)	50	40	30	20	10	0	-10	-20	-30	-40	-50	-60
		Equivalent Chill Temperature (°F)										
calm	50	40	30	20	10	0	-10	-20	-30	-40	-50	-60
5	48	37	27	16	6	-5	-15	-26	-36	-47	-57	-68
10	40	28	16	4	-9	-24	-33	-46	-58	-70	-83	-95
15	36	22	9	-5	-18	-32	-45	-58	-72	-85	-99	-112
20	32	18	4	-10	-25	-39	-53	-67	-82	-96	-110	-121
25	30	16	0	-15	-29	-14	-59	-74	-88	-10 4	-118	-133
30	28	13	-2	-18	-33	-48	-63	-79	-94	-10 9	-125	-140
35	27	11	-4	-20	-35	-51	-67	-82	-9 8	-113	-1 29	-145
40	26	10	-6	-21	-37	-53	-69	-85	-100	-116	-132	-148
(Wind speeds greater than 40 mph have little additional effect.)	ln < hr	•	h dry skin. Maximum DANGE false sense of security Danger f		ER from free d flesh wi	-		DANGE) seconds		
Trenchfoot and immersion foot may occur at any point on this chart.												

Developed by U.S. Army Research Institute of Environmental Medicine, Natick, MA.

(1) Reproduced from American Conference of Governmental Industrial Hygienists, <u>Threshold</u> <u>Limit Values and Biological Exposure Indices for 1985-1986</u>, p. 01.

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ATTACHMENT D

CHEMICAL HAZARDS

References:

- Sax, N. Irving and Lewis, Richard J. <u>Dangerous Properties of Industrial Materials -</u> <u>7th Edition</u>, 1989.
- 2. U.S. Department of Health and Human Services. NIOSH Pocket Guide to Chemical Hazards, June 1990.

ACETONE ABC750

25

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SYNS:

L&-DIMETHYL-m-DIOXAN-4-OL
ACETATE
1.6-DIMETHYL-m-DIOXAN-4-YL
ACETATE
DIOXIN (BACTERICIDE) (OBS)
GIV GARD DXN
NCI-C56213
CODEN:
ENMUDM 8(Suppl
7).1.86
ENMUDM 7.677.85
JNCIAM 53,791.74
GCTB** 3/25/77

orl-rat LD50: 1930 mg/kg

IARC Cancer Review: Animal Limited Evidence IMEMDT 15.177.77

THR: An experimental carcinogen. Moderately toxic by ingestion. See also ESTERS. When heated to decomposition it emits acrid smoke and fumes.

ABC500		HR: 2
2'-ACETONAL	PHTHONE	
CAS: 93-08-3		NIOSH: AL 2988000
mf: $C_{12}H_{10}O$	mw: 170.22	
SYNS:		

ORANGE CRYSTALS	METHYL-B-NAPHTHYL KETONE
B-ACETONAPHTHALENE	METHYL-2-NAPHTHYL KETONE
B-ACETYLNAPHTHALENE	B-METHYL NAPHTHYL KETONE
2-ACETYLNAPHTHALENE	142-NAPHTHALENYL)ETHANONE
ACETONAPHTHONE	B-NAPHTHYL METHYL KETONE
B-ACETONAPHTHONE	2-NAPHTHYL METHYL KETONE
2-ACETONAPHTHONE	
TONICETY DATA	CODEN

TOXICITY DATA: skn-hmn 500 mg/24H ori-mus LD50:599 mg/kg CODEN: FCTXAV 13,681,75 MDZEAK 8,244,67

Reported in EPA TSCA Inventory.

THR: Moderately toxic by ingestion. A human skin irritant. When heated to decomposition it emits acrid smoke and fumes.

A BC750		HR: 2
ACETONE		
CAS: 67-64-1		NIOSH: AL 3150000
DOT: 1090		
mf: C ₃ H ₆ O	mw: 58.09	

PROP: Colorless liquid, fragrant mint-like odor. Mp: -94.6°, bp: 56.48°, ulc: 90, flash p: 0°F (CC), lel: 2.6%, uel: 12.8%, d: 0.7972 @ 15°, autoign temp: (color) 869°F, vap press: 400 mm @ 39.5°, vap d: 2.00. Misc in water, alc, and ether.

SYNS:

ACETON (GERMAN DUTCH POL. ISHO DIMETHYLFORMALDEHYDE DIMETHYLKET AL DIMETHYL KETONE KETONE PROPANE B-KETOPROPANE

TOXICITY DATA:

skn-rbt 500 mg/24H MLD eye-rbt 100 mg/24H MOD cyt-smc 200 mmol/tube sin-smc 47600 ppm cyt-ham: fbr 40 g/L ihl-mam TCLo: 31500 µg/m³/ 24H (1-13D preg): REP orl-man TDLo: 2857 mg/kg orl-man TDLo: 2857 mg/kg ivn-rat LD50:5500 mg/kg ihl-man TDLo: 440 μ g/m³/6M ihl-man TDLo: 10 mg/m³/6H orl-mus LD50:3000 mg/kg eye-hmn 500 ppm skn-rbt 395 mg open MLD eve-rot 3950 µg SEV ihl-hmn TCLo: 500 ppm: EYE ihl-man TCLo: 12000 ppm/4H: CNS unk-man LDLo: 1159 mg/kg ihl-rat LCLo: 16000 ppm/4H ipr-rat LDLo: 500 mg/kg ihl-mus LCLo: 110000 mg/m³/ 62M ipr-mus LD50:1297 mg/kg ort-dog LDLo: 24 g/kg ipr-dog LDLo:8 g/kg scu-dog LDLo: 5 g/kg skn-rbt LD50:20 g/kg scu-gpg LDLo: 5000 mg/kg

METHYL KETONE PROPANONE : PROPANONE PYROACETIC ACID PYROACETIC ETHER RCRA WASTE NUMBER L'007

CODEN:

28ZPAK -.42.72 28ZPAK -.42,72 HEREAY 33,457,47 ANYAA9 407,186,83 FCTOD7 22.623.84 GTPZAB 26(6),24,82

AJOPAA 29,1363,46 JIHTAB 25,282,43 AOHYA3 16,73,73	34ZIAG64.69 DIAEAZ 15.810,66 NPIRI* 1.1,74 GISAAA 42(8)42.77 GISAAA 42(8)42.77 PCJOAU 14.162,80 JIHTAB 25.282,43 UCDS** 5/7/70
	UCDS** 5/7/70 AJOPAA 29.1363.46 JIHTAB 25.282.43

85DCAI 2.73.70 AIHAAP 23,95,62 JPPMAB 11,150,59 AGGHAR 5,1,33

SCCUR* -.1.61 AEXPBL 18,218,1884 AEXPBL 18,218,1884 AEXPBL 18.218.1884 UCDS** 5/7/70 AGGHAR 5,1,33

On Community Right To Know List. Reported in EPA TSCA Inventory.

OSHA PEL: TWA 1000 ppm ACGIH TLV: TWA 750 ppm; STEL 1000 ppm DFG MAK: 1000 ppm (2400 mg/m³) NIOSH REL: TWA 590 mg/m³

DOT Classification: Flammable Liquid, Label: Flammable Liquid

THR: Moderately toxic by various routes. A skin and severe eye irritant. Human systemic effects by inhalation: changes in EEG, changes in carbohydrate metabolism, nasal effects, conjunctiva irritation, respiratory system effects, nausea and vomiting, and muscle weakness. Human systemic effects by ingestion: coma, kidney damage, and metabolic changes. Narcotic in high concentration. In industry, no injurious effects have been reported other than skin irritation resulting from its defatting action, or headache from prolonged inhalation. A common air contaminant. Dangerous disaster hazard due to fire and explosion hazard; can react vigorously

ABDOOD ACETONE CHLOROFORM

with oxidizing materials. Potentially explosive reaction with nime acid + sulfuric acid; bromine trifluoride; nitrosyl chlonde + platinum; nitrosyl perchlorate; chromyl chloride; thiotrithiazyi perchlorate; and 2.4.6-trichloro-1.3.5-triazine - water. Reacts to form explosive peroxide products with 2-metnyl-1,3-butadiene; hydrogen peroxide; and peroxomonosulfuric acid. Ignites on contact with activated carbon: chromium moxide; dioxygen difluoride + carbon dioxide; and potassium-tert-butoxide. Reacts violently with bromoform: chloroform + alkalies: bromine: and sulfur dichloride. Incompatible with CrO; (nitric + acetic acid); NOCI; nitryl perchlorate; permonosulfuric acid; NaOBr; (sulfuric acid + potassium dichromate): (thio-diglycol + hydrogen peroxide); trichloromelamine; air: HNO3; chloroform; and H₂SO₄. To fight fire, use CO₂, dry chemical, alcohol foam. For further information see Vol. 4. No. 3 of DPIM Report.

HR: 3 ABD000 ACETONE CHLOROFORM NIOSH: UC 0175000 CAS: 57-15-8

mw: 177.46 mf: C4H7Cl3O

PROP: Crystals, camphor odor. Mp: 97°, bp: 167°.

SYNS:	
CHLORETONE	CHLORBUTANOL
CLORTRAN	CHLORBUTOL
METHAFORM	CHLOROBUTANOL
SEDAFORM	TRICHLORO-un-BUTYL ALCOHOL
B.B.B-TRICHLORO-Len-BUTYL AL-	uen-TRICHLOROBUTYL ALCOHOL
COHOL	LI.I.I.TRICHLORO-2-METHYL-2-
HCP	PROPANOL
ANHYDROUS CHLOROBUTANOL	

CODEN:

TOXICITY DATA:

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MUREAV 90.91.81 mmo-sat 20 µmol/plate HEREAY 33,457,47 cyt-smc 10 mmoi/tube skn-rbt 850 µg MLD XEURAQ MDDC-1715 eve-rbt 9180 µg/30S MLD XEURAO MDDC-1715 orl-dog LDLo: 238 mg/kg AIPTAK 8.77.01 orl-rbt LDLo: 213 mg/kg AIPTAK 8.77.01 par-frg LDLo: 800 mg/kg AIPTAK 8.77.01

Reported in EPA TSCA Inventory.

THR: Poison by ingestion. Moderately toxic by parenteral route. A narcotic. A skin and eye irritant. Mutagenic data. See also CHLORAL HYDRATE, which acts similarly. Dangerous: can react with oxidizing materials. Combustible when exposed to heat or flame. When heated to decomposition it emits toxic fumes of Cl⁻. See also PHOSGENE.

ABD250 HR: 3 ACETONE DIETHYL KETAL CAS: 126-84-1 NIOSH: AL 4900000

mf: C₇H₁₆O₂ mw: 132.23

USAF DO-44

SYNS: 2.2-DIETHOXYPROPANE TOXICITY DATA: ipr-mus LD50:125 mg/kg CODEN: NTIS** AD277-689

Reported in EPA TSCA Inventory.

THR: Poison by intraperitoneal route. When heated to decomposition it emits acrid smoke.

ABD500 HR: 3 ACETONE DIETHYLSULFONE CAS: 115-24-2 NIOSH: TX 3850000 mw: 228.35

mf: $C_7H_{16}O_4S_2$

ori-gpg LDLo:8500 mg/kg

PROP: D: 1.183, mp: 127°-128°, bp: 300° (sl decomp), Sol in water, alc, and ether.

SYNS:

ACETONE BIS(ETHYL SULPONE) 2.2-BIS(ETHYLSULPONYL)PRO- PANE DIETHYLSULPONDIMETHYL- METHANE	PROPANE DIETHYL SULPONE SULFONAL SULPONMETHANE
TOXICITY DATA:	CODEN:
unk-man LDLo: 147 mg/kg orl-dog LDLo: 800 mg/kg orl-rbt LDLo: 3000 mg/kg	85DCAI 2,73,70 12VXA5 8,1003,68 HBAMAK 4,1404,35

THR: A human poison by unspecified route. Moderately toxic by ingestion. Mutagenic data. When heated to decomposition it emits toxic fumes of SO_r.

HBAMAK 4,1404,35

ABD750 ACETONE OIL	HR: 3
	NIOSH: AL 6700000

DOT: 1091

PROP: (a) Standard: light, lemon-yellow. (b) Refined: almost water white. (c) Heavy: dark, orange-yellow. Bp: (a) 75-160°, (c) 80-225°. D: (a) 0.826-0.830, (b) 0.812, (c) 0.885-0.865.

DOT Classification: Flammable Liquid, Label: Flammable Liquid

THR: Dangerous fire and explosion hazard when exposed to heat or flame. Can react vigorously with oxidizing materials. Some carcinogenic activity. To fight fire, use CO₂, dry chemical.

ABE000 ACETONE PEROXIDE

HR: 3

PROP: Liquid. The trimeric form is crystalline. Mp: 97°. THR: No toxicity data. See PEROXIDES, ORGANIC. Flammable by spontaneous chemical reaction; can react vigorously with reducing materials. The trimeric form is shock-sensitive and static-electricity-sensitive and may detonate.

BBL250 BENZENE

pounds. Reacts explosively with aniline at 240°C/7.6 bar. Can react vigorously with oxidizing materials. To fight fire, use water, CO2, water mist or spray, dry chemical. See also ANILINE. For further information. see Vol. 4. No. 4 of DPIM Report.

BBL250

BENZENE

HR: 3 NIOSH: CY 1400000

CAS: 71-43-2 DOT: 1114 mw: 78.12 mf: C₆H₆

PROP: Clear, colorless liquid. Mp: 5.51°, bp: 80.093°-80.094°, flash p: 12°F (CC), d: 0.8794 @ 20°, autoign temp: 1044°F, lel: 1.4%, uel: 8.0%, vap press: 100 mm $(\tilde{a}, 26.1^{\circ}, \text{ vap d: } 2.77, \text{ ulc: } 95-100.$

CODEN:

SYNS:

DIANNULENE	CYCLOHEXATRIENE
BENZEEN (DUTCH)	FENZEN (CZECH)
BENZEN (POLISH)	MINERAL NAPHTHA
BENZIN (OBS.)	MOTOR BENZOL
BENZINE (OBS.)	NCI-C55276
BENZOL (DOT)	NITRATION BENZENE
BENZOLE	PHENE
BENZOLENE	PHENYL HYDRIDE
BENZOLO (ITALIAN)	PYROBENZOL
BICARBURET OF HYDROGEN	PYROBENZOLE
CARBON OIL	RCRA WASTE NUMBER U019
COAL NAPHTHA	

TOXICITY DATA:

AIHAAP 23.95.62 skn-rbt 15 mg/24H open MLD eye-rbt 88 mg MOD AMIHAB 14.387.56 eye-rbt 2 mg/24H SEV 28ZPAK -.23,72 skn-rbt 500 mg/24H MOD 28ZPAK -.23.72 slt-dmg-ori 11250 µmol/L PMRSDJ 5.325,85 oms-hmn:lym 5 µmol/L CNREA8 45.2471,85 mma-mus: emb 2500 mg/L PMRSDJ 5.639.85 cyt-ham:ing 550 mg/L PMRSDJ 5,427,85 sin-ham: ivr 62500 µg/L PMRSDJ 5.397.85 ihl-rat TCLo: 670 mg/m³/24H HYSAAV 33,327,68 (15D pre/1-22D preg):REP ihl-rat TCLO: 50 ppmv24H (7-JHEMA2 24,363.80 14D preg): TER shi-rat TCLO: 150 ppm/24H (7-JHEMA2 24.363.80 14D preg): TER ihl-man TCLo: 200 mg/m³/78W-EJCAAH 7.83.71 I:CAR.BLD ihl-hmn TCLo: 10 ppm/8H/10Y-TRBMAV 37,153,78 I:CAR.BLD ori-rat TDLo: 52 g/kg/52W-I: MELAAD 70,352,79 CAR ihl-rat TCLo: 1200 ppm/6H/10W-PAACA3 25,75,84 I ETA orl-mus LD50:18250 mg/kg/2Y-NTPTR* NTP-TR-C:CAR 289,86 ihl-mus TCLo: 300 ppm/6H/16W-TXAPA9 75.358,84 I:ETA skn-mus TDLo: 1200 g/kg/49W-BJCAAI 16.275,62 1:NEO ipr-mus TDLo: 1200 mg/kg/8W-TXAPA9 82.19.86

1:NEO

par-mus TDLo: 670 mg/kg/19W-KLWOAZ 12.109.33 L: ETA shi-hmn TC:150 ppm/15M/8Y-L: BLOOAW 52.285.78 CAR.BLD orl-rat TD: 52 g/kg/1Y-I:CAR AJIMD8 4.589,83 ori-rat TD:10 g/kg/52W-I:CAR MELAAD 70.352.79 thl-hmn_TC:150 ppm/15M/8Y-I: NEJMAG 271.872.64 CAR.BLD ihl-man TC: 150 ppm/11Y-I: BLUTA9 28.293.74 CAR.BLD ihl-mus TC: 1200 ppm/6H/10W-PAACA3 25,75,84 I:ETA ihl-hmn LCLo:2 pph/SM TABIA2 3.231.33 ori-hmn TDLo: 130 mg/kg: AHYGAJ 31.336,1897 CNS.GIT ihl-hmn LCLo: 20000 ppm/5M 29ZUA8 -.-.53 ihl-hmn TCLo: 210 ppm: 27ZXA3 -.341.63 CNS.GIT ihl-man TCLo: 150 ppm/1Y-I: BLUTA9 28.293.74 MET.BLD hi-hmn TCLo: 100 ppm: INMEAF 17,199,48 CNS.GIT.SKN ihl-hmn LCLo:65 mg/m³/5Y: ARGEAR 44,145,74 BLD orl-rat LD50: 3400 mg/kg NPIRI* 1,5,74 ihl-rat LC50:10000 ppm/7H 28ZRAQ -,113,60 ipr-rat LD50:2890 µg/kg 36YFAG -.302,77 orl-mus LD50:4700 mg/kg HYSAAV 32.349.67 ihl-mus LC50:9980 ppm JIHTAB 25,366,43 NPIRI* 1,5,74 ska-mus LD50:48 mg/kg AGGHAR 18,109.60 ipr-mus LD50:990 µg/kg ori-dog LDLo: 2000 mg/kg HBAMAK 4,1313,35 ihl-dog LCLo: 146000 mg/m³ HBTXAC 1,324,56 HBTXAC 1.324,56 ihl-cat LCLo: 170000 mg/m³ ivn-rbt LDLo:88 mg/kg JTEHD6 HBTXAC 1.42.56

scu-mus TDLo:600 mg/kg/17W-

I:ETA

ipr-gpg LDLo: 527 mg/kg scu-frg LDLo: 1400 mg/kg ihl-mam LCLo: 20000 ppm/5M

IARC Cancer Review: Human Limited Evidence IMEMDT 7,203,74; Animal Inadequate Evidence IMEMDT 7,-203,74; IARC Cancer Review: Animal Limited Evidence IMEMDT 29,93,82; Human Sufficient Evidence IMEMDT 29,93,82; NTP Carcinogenesis Studies (gavage); Clear Evidence: mouse, rat NTPTR* NTP-TR-289,86. EPA Genetic Toxicology Program. Reported in EPA TSCA Inventory. On Community Right To Know List.

OSHA PEL: TWA 1 ppm/8H; Pk 5 ppm/15M ACGIH TLV: TWA 10 ppm (suspected human carcinogen); BEI (total phenol in urine at end of shift) 50 mg/L recommended as a mean value DFG TRK: 8 ppm (26 mg/m³) NIOSH REL: CL 1 ppm/60M DOT Classification: Flammable Liquid, Label: Flammable Liquid THR: A human poison by inhalation. An experimental poi-

son by skin contact, intraperitoneal, intravenous and possibly other routes. Moderately toxic by ingestion and subcuta-

KRANAW 9,403.32

-(Suppl:2),45,77

HBAMAK 4.1313.35

AEPPAE 138,65,28

leous routes. A severe eye and moderate skin irritant. Hunan systemic effects by inhalation and ingestion: euphona. somnolence, changes in REM sleep, changes in motor activ-(v, nausea or vomiting, reduced number of blood platelets, ther unspecified blood effects, dermatitis, and fever. A numan carcinogen which produces myeloid leukemia and lymphomas by inhalation. An experimental carinogen, neoplastigen, tumorigen, and teratogen. Other exremental animal reproductive effects. Human mutagenic data. A narcotic. In industry, inhalation is the primary route of chronic benzene poisoning. Poisoning by skin contact ias been reported. Recent (1987) research indicates that effects are seen at less than 1 ppm. Exposures needed to be reduced to 0.1 ppm before no toxic effects were oberved. Elimination is chiefly through the lungs. A common ur contaminant.

A dangerous fire hazard when exposed to heat or flame. Explodes on contact with diborane; bromine pentafluoride; remanganic acid: peroxomonosulfuric acid: and peroxodisulfuric acid. Forms sensitive, explosive mixtures with ioline pentafluoride; silver perchlorate; nitryl perchlorate; niric acid; liquid oxygen; ozone; arsenic pentatluoride + potassium methoxide (explodes above 30°C). Ignites on contact with sodium peroxide + water: dioxygenyl tetraluoroborate; iodine heptafluoride; and dioxygen difluoride. ligorous or incandescent reaction with hydrogen + Raney nickel (above 210°C); uranium hexafluoride: and bromine rifluoride. Can react vigorously with oxidizing materials. such as Cl_2 ; CrO_3 ; O_2 ; $NClO_4$; O_3 ; perchlorates; $(AlCl_3 +$ $FClO_4$; (H₂SO₄ + permanganates); K₂O₂; (AgClO₄ + aceric acid); Na2O2. Moderate explosion hazard when exposed o heat or flame. Use with adequate ventilation. To fight lire, use foam, CO₂, dry chemical. For further information, see Vol. 4, No. 6 of DPIM Report.

Poisoning occurs most commonly via inhalation of the apor, although benzene can penetrate the skin and cause poisoning. Locally, benzene has a comparatively strong rritating effect, producing ervthema and burning, and, in nore severe cases, edema and even blistering. Exposure to high concentrations of the vapor (3000 ppm or higher) may result from failure of equipment or spillage. Such exposure, while rare in industry, may cause acute poisoning, characterized by the narcotic action of benzene on the central nervous system. The anesthetic action of benzene s similar to that of other anesthetic gases, consisting of a reliminary stage of excitation followed by depression and. if exposure is continued, death through respiratory failure. The chronic, rather than the acute form, of benzene poisonng is important in industry. It is a recognized leukemogen. there is no specific blood picture occurring in cases of chronic benzol poisoning. The bone marrow may be hyposlastic, normal, or hyperplastic, the changes reflected in he peripheral blood. Anemia, leucopenia, macrocytosis, reticulocytosis, thrombocytopenia, high color index, and rolonged bleeding time may be present. Cases of myeloid leukemia have been reported. For the worker, repeated blood examinations are necessary, including hemoglobin determinations, white and red cell counts and differential smears. Where a worker shows a progressive drop in either red or white cells, or where the white count remains low, 5.000/mm₃ or the red count < 4.0 million/mm₃, on two successive monthly examinations, he should be immediately removed from benzene exposure. Elimination is chiefly through the lungs, when fresh air is breathed. The portion that is absorbed is oxidized, and the oxidation products are combined with sulfuric and glycuronic acids and eliminated in the urine. This may be used as a diagnostic sign. Benzene has a definite cumulative action, and exposure to a relatively high concentration is not serious from the point of view of causing damage to the blood-forming system. provided the exposure is not repeated. In acute poisoning, the worker becomes confused and dizzy, complains of tightening of the leg muscles and of pressure over the forehead. then passes into a stage of excitement. If allowed to remain exposed, he quickly becomes stuperied and lapses into coma. In non-fatal cases, recovery is usually complete with no permanent disability. In chronic poisoning the onset is slow, with the symptoms vague; fatigue, headache, dizziness, nausea and loss of appetite, loss of weight and weakness are common complaints in early cases. Later, pallor, nosebleeds, bleeding gums, menorrhagia, petechiae and purpura may develop. There is great individual variation in the signs and symptoms of chronic benzene poisoning.

BBL500 BENZENEACETALDEHYDE

HR: 2

DENZENEACE I ALDEHYDE CAS: 122-78-1 NIC

mf: C_xH₈O mw: 120.16

NIOSH: CY 1420000

PROP: Oily, colorless liquid which polymerizes and grows more viscous on standing. Odor similar to lilac and hyacinth. Has been crystallized, mp: 33-34°, d:(25/25) 1.023-1.030, bp: (10) 78°, n (20/D) 1.524-1.528. Sltly sol in water. Sol in alc, ether. One part is sol in two parts of 80% alc forming a clear solution.

SYNS:

skn-hmn 2%/48H

ori-rat LD50:1550 mg/kg

orl-mus LD50:3890 mg/kg

orl-gpg LD50:3890 mg/kg

HYACINTHIN PH PAA a⁻ PHENYLACETALDEHYDE a⁻ PHENYLACETIC ALDEHYDE TOXICITY DATA:

PHENYLETHANAL a-TOLUALDEHYDE a-TOLUIC ALDEHYDE

> CODEN: FCTXAV 17.377.79 FCTXAV 17.377.79 FCTXAV 17.377.79 FCTXAV 17.377.79

Reported in EPA TSCA Inventory.

THR: Moderately toxic by ingestion. Human skin irritant. When heated to decomposition it emits acrid smoke and irritating tumes. Used in perfumery. See also ALDE-HYDES.

BOV000 4-BUTANOLIDE

T.

TOXICITY DATA:	CODEN:
eye-rbt 83 mg	AIHAAP 19.171.58
ihi-rat LC50:4020 ppm/4H	AIHAAP 19.171.58
ori-rat LD50:1500 mg/kg	AIHAAP 19.171.58
pr-rat LD50: 399 mg/kg	AIHAAP 19.171.58
:hl-mus LC50:2500 ppm/4H	AIHAAP 19.171.58

Reported in EPA TSCA Inventory.

OSHA PEL: TWA 10 ppm ACGIH TLV: TWA 0.5 ppm DFG MAK: 0.5 ppm (1.5 mg/m³) NIOSH REL: (n-Alkane Mono Thiols) CL 0.5 ppm/15M

DOT Classification: Flammable Liquid, Label: Flammable Liquid

THR: Poison by intraperitoneal route. Moderately toxic by inhalation and ingestion. An eye irritant. Dangerous fire hazard by exposure to heat, flame, sparks, or powerful oxidizers. Reacts violently with HNO₃. Incompatible with acids; acid fumes; oxidizing materials; heat; flame; sparks. To fight fire, use alcohol foam. When heated to decomposition it emits toxic SO_r. See also MERCAPTANS.

BOV000		HR: 3
4-BUTANOLID	E	
CAS: 96-48-0		NIOSH: LU 3500000
mf: C₄H ₆ O ₂	mw: 86.10	

PROP: Colorless liquid, mild odor. Mp: -44°, bp: 206°, flash p: 209°F (OC), d: 1.124 @ 25°/4°, vap d: 3.0.

SYNS:

gamma-o480	DIHYDRO-2(3H)-FURANONE
gamma-BL	+HYDROXYBUTANOIC ACID
BLO	LACTONE
BLON	Y-HYDROXYBUTYRIC ACID
BUTYRIC ACID LACTONE	CYCLIC ESTER
V-BUTYROLACTONE	+HYDROXYBUTYRIC ACID -
BUTYRYL LACTONE	LACTONE
a-BUTYROLACTONE	NCI-C55878
4-DEOXYTETRONIC ACID	TETRAHYDRO-2-FURANONE
Y-HYDROXYBUTYROLACTONE	
TOVICEN	CODEN

IOXICITY DATA:	CODEN:
dnd-bcs 20 µL disc	PMRSDJ 1.175.81
otr-ham:kdy 25 mg/L	PMRSDJ 1.638.81
orl-rat TDLo: 25 g/kg (20D	ARANDR 10,239,83
male): REP	
skn-mus TDLo:50 g/kg/42W-I:	JNCIAM 31.41.63
ETA	
orl-rat LD50:1800 mg/kg	85GMAT31,82
ipr-rat LD50: 1000 mg/kg	AITEAT 13,70.65
ipr-mus LD50:1100 mg/kg	AITEAT 13.70.65
ivn-rbt LDLo: 500 mg/kg	AITEAT 13,70,65

IARC Cancer Review: Animal No Evidence IMEMDT 11,231,76. EPA Genetic Toxicology Program. Reported in EPA TSCA Inventory.

THR: Moderately toxic by ingestion, intravenous and intraperitoneal routes. An experimental tumorigen by skin contact. Experimental reproductive effects. Mutagenic data.

Less acutely toxic than B-propiolactone. Combustible when exposed to heat or flame: can react with oxidizing inaterials. To fight fire, use foam, alcohol foam, CO₂, dry chemical, When heated decomposition it emits acrid and imitating fumes. For further information, see gamma Butvrolactone, Vol. 1, No. 3 of DPIM Report.

BOV250

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2-BUTANONE
CAS: 78-93-3
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NIOSH: EL 6475000

HR: 3

DOT: 1193/1232 mf: C_4H_9O mw: 72.12

CH₃CO•CH₂CH₃

PROP: Colorless liquid, acetone-like odor. Bp: 79.57°, fp: -85.9°, lel: 1.8%, uel: 11.5%, flash p: 22°F (TOC), d: 0.80615 @ 20°/20°, vap press: 71.2 mm @ 20°, autoign temp: 960°F, vap d: 2.42, ULC: 85-90.

SYNS:

AETHYLMETHYLKETON (GER-	MEK
MAN)	METHYL ACETONE (DOT)
BUTANONE 2 (FRENCH)	METHYL ETHYL KETONE (ACGIH)
ETHYL METHYL CETONE	METILETILCHETONE (ITALIAN)
(FRENCH)	METYLOETYLOKETON (POLISH)
ETHYLMETHYLKETON (DUTCH)	RCRA WASTE NUMBER U159
ETHYL METHYL KETONE (DOT)	
TOXICITY DATA:	CODEN:

TOXICITY DATA:

eye-hmn 350 ppm JIHTAB 25,282,43 skn-rbt 500 mg/24H MOD ЛНТАВ 25.282.43 TXAPA9 19,276,71 skn-rbt 402 mg/24H MLD skn-rbt 13780 µg/24H open AIHAAP 23,95,62 MLD TXAPA9 19,276,71 eye-rbt 80 mg MUREAV 149.339.85 sla-smc 33800 ppm TXAPA9 28,452,74 ihl-rat TCLo: 3000 ppm/7H (6-15D preg): REP TXAPA9 28,452,74 ihl-rat TCLo: 1000 ppm/(6-15D preg): TER ihl-hmn TCLo: 100 ppm/5M: IRR JIHTAB 25,282,43 TXAPA9 19.699,71 orl-rat LD50:2737 mg/kg ihl-rat LCLo: 2000 ppm/4H JIHTAB 31.343,49 ihl-mus LC50:40 g/m³/2H 85GMAT -,83,82 ipr-mus LD50:616 mg/kg SCCUR* -,6,61 UCDS** 5/7/70 skn-rbt LD50:13 g/kg FCTXAV 15.611,77 ipr-gpg LDLo: 2000 mg/kg

Community Right To Know List. EPA Genetic Toxicology Program. Reported in EPA TSCA Inventory.

OSHA PEL: TWA 200 ppm ACGIH TLV: TWA 200 ppm; STEL 300 ppm DFG MAK: 200 ppm (590 mg/m³) NIOSH REL: (Ketones) TWA 590 mg/m³

DOT Classification: Flammable Liquid, Label: Flammable Liquid

THR: Moderately toxic by ingestion, skin contact and intraperitoneal routes. Human systemic effects by inhalation: conjunctiva irritation and unspecified effects on the nose

ind respiratory system. An experimental teratogen. Experiiental reproductive effects. A strong irritant. Human eye intation (a, 350 ppm). Affects peripheral nervous system and central nervous system. See also KETONES. Dangerus fire hazard when exposed to heat or flame. Moderately xplosive when exposed to flame. Reaction with hydrogen peroxide + nitric acid forms a heat and shock sensitive xplosive product. Ignition on contact with potassium tertutoxide. Mixture with 2-propanol will produce explosive peroxides during storage. Vigorous reaction with chloroform + alkali. Incompatible with chlorosulfonic acid: leum. To fight fire, use alcohol foam, CO₂, dry chemical. when heated to decomposition it emits acrid smoke and fumes.

10V625

HR: 3 **2-BUTANONE OXIME HYDROCHLORIDE** CAS: 4154-69-2 $f: C_4H_{10}ClNO$ mw: 123.58

SYN: 32 HYDROXYLIMINIOBUTANE CHLORIDE:

"HR: Decomposes violently above 50°C. When heated to ecomposition it emits toxic fumes of Cl⁻ and NO_r.

ROV750 HR: 3 **JUTAZOLIDINE SODIUM** NIOSH: UQ 8300000

CAS: 129-18-0 mf: $C_{19}H_{20}N_2O_2 \cdot Na$

SYNS:

mw: 331.40 -BUTYL-1.2-DIPHENYL-3.5-PYRA-PHENYLBUTAZONE SODIUM ZOLIDINEDIONE SODIUM SALT SODIUM BUTAZOLIDINE 5-DIOXO-1.2-DIPHENYL-4-N-BU-SODIUM PHENYLBUTAZONE

ZONE

SODIUM SALT of PHENYLBUTA-

TYLPYRAZOLIDIN SODIUM JIPHENYLDIOXOBUTYLPYRAZO-LIDINE-BUTAZOLIDINE-SO-DUM

	COXICITY DATA:	CODEN:
-	ori-wmn LDLo:16 mg/kg	AIMEAS 39.1096.53
	cu-rat LD50:360 mg/kg	ARZNAD 8.229,58
	ri-mus LD50:476 mg/kg	RPOBAR 2,314,70
	pr-mus LD50:169 mg/kg	RPOBAR 2,314,70
	ivn-rat LD50:113 mg/kg	FRPSAX 13,922,58
	scu-mus LD50:271 mg/kg	FRPSAX 12.521,57
	vn-mus LD50:94 g/kg	FRPSAX 13.922.58

THR: A human poison by ingestion. Human systemic effects by ingestion: respiratory system damage, agranulocyosis, and dermatitis. An experimental poison via subcutaneous, intravenous, and intraperitoneal routes. An anti-inflammatory drug. When heated to decomposition it mits toxic fumes of NO, and Na₂O.

BOV800

BUTEA FRONDOSA, seed extract

NIOSH: EM 2455000

HR: 3

PROP: Indian plant belonging to the family Leguminosae (UEBA6 11.43.73).

HR: 3

HR: 3

NIOSH: GP 9625000

NIOSH: DN 9205000

SYN: PALASH SEED EXTRACT

TOXICITY DATA:	CODEN
ort-rat TDLo: 250 mg/kg (1-5D	UPPAZ 13.239.69
pregi:REP	
ori-mus TDLo: 50 mg/kg (1-5D	UPPAZ 13.239.69
preg): REP	
orl-mus LD50:7500 mg/kg	UPPAZ 13.239.69
ipr-mus LD50:20 mg kg	UEBA6 11.43.73

THR: Poison by intraperitoneal route. Experimental reproductive effects.

mw: 516.72

BOV825

BUTEDRIN CAS: 5716-20-1

mf: $C_{24}H_{38}N_2O_4 \cdot H_2O_4S$

SYNS:

BAMETAN SULFATE CYCLATE BAMETHAN SULFATE ECLERIN BASCURAT GARMIAN BUPATOL PERIPHETOL BUTTBATOL ROTESAR a-0 BUTYLAMINO METHYLI-p-HY-VASCULAT DROXYBENZYL ALCOHOL SUL-VASCULIT VASCUNICOL FATE BUTYLNORSYMPATOL VASKULAT TOXICITY DATA: ipr-mus LD50:210 mg/kg

CODEN: NURDN 6.585.82 NURDN 6.585,82 NURND 6.585.82

THR: Poison by intravenous and intraperitoneal routes. Moderately toxic by subcutaneous route. A vasodilator. When heated to decomposition it emits toxic fumes of SO_r and NO₁. See also SULFATES.

BOW000

trans-2-BUTENAL

scu-mus LD50:422 mg/kg

ivn-mus LD50:72 mg/kg

CAS: 123-73-9 mf: C₄H₆O mw: 70.10

PROP: Water-white, mobile liquid; pungent, suffocating odor. Bp: 104°, fp: -76.0°, lel: 2.1%, uel: 15.5%, flash p: 55°F, d: 0.853 @ 20°/20°, vap d: 2.41, autoign temp: 450°F.

SYNS: ALDEHYDE CROTONIQUE ETHYLENE GLYCOL DIPROPIO-(FRENCH) NATE (SCI) (E)-2-BUTENAL ETHYLENE PROPIONATE CROTENALDEHYDE B-METHYL ACROLEIN CROTONALDEHYDE (ACGIH) NCI-C56279 CROTONIC ALDEHYDE PROPYLENE ALDEHYDE 1.2-ETHANEDIOL DIPROPANOATE RCRA WASTE NUMBER U053 TOPANEL. (9CD)

TOXICITY DATA:

eve-hmn 45 ppm skn-rbt 500 mg open MLD mmo-sat 250 µg/plate

CODEN: AIHAAP 28.561.67 UCDS** 4/21/67 ENMUDM 5(Suppl 1),3.83

SYNS:

0-126-DICHLOROANILINO PHE-VERIODIN NYLIACETIC ACID MONOSO-PROPHENATIN DIUM SALT SODIUM (0+2.6-DICHLOROANILI-1+16-DICHLOROPHENYL) NOIPHENYLIACETATE AMINO-BENZENEACETIC ACID SODIUM (0-(12.6-DICHLORO-PHENYLIAMINOIPHENYLIACE-MONOSODIUM SALT DICHRONIC TATE TSUDOHMIN DICLOFENAC SODIUM DICLOPHENAC SODIUM VALETAN VOLTAREN OP 45840 KRIPLEX VOLTAROL TOXICITY DATA: CODEN: OYYAA2 27.117.84 orl-rat TDLo:1 mg/kg (21D preg): TER orl-rat TDLo: 312 mg/kg (22W KSRNAM 6.1521.72 male): REP orl-rat TDLo:6 mg/kg (9-14D KSRNAM 6.1673.72 preg): TER orl-rat LD50:76500 µg/kg OYYAA2 16.353.78 NIIRDN 6.311.82 ipr-rat LD50:25 mg/kg scu-rat LD50:83 mg/kg IYKEDH 5.106.74 ivn-rat LD50:117 mg/kg IYKEDH 5.106.74 ARZNAD 34.280.84 orl-mus LD50:125 mg/kg ipr-mus LD50:130 mg/kg IYKEDH 5.106.74 scu-mus LD50: 390 mg/kg NIIRDN 6.311.82 ivn-mus LD50:116 mg/kg IYKEDH 5.106.74 KSRNAM 6.1521.72 orl-dog LD50:59 mg/kg ivn-dog LD50:42 mg/kg KSRNAM 6.1521.72 orl-rbt LD50:157 mg/kg KSRNAM 6.1521.72

THR: Poison by ingestion, intravenous, intraperitoneal and subcutaneous routes. An experimental teratogen. Experimental reproductive effects. An anti-inflammatory agent. When heated to decomposition it emits very toxic fumes of Cl^- . Na₂O and NO₇.

DEP400 HR: 3 2-DICHLOROARSINOPHENOXATHIIN

CAS: 63834-20-8 NIOSH: SP 7000000 mf: C12H7AsCl2OS mw: 345.07

SYN: TL 472

ş

TOXICITY DATA:	CODEN:
orl-rat LDLo: 250 mg/kg	NCNSA6 5,13.53
ihl-mus LCLo:400 mg/m ³ /10M	NDRC** NDCrc-
-	132,Dec.42

Arsenic and its compounds are on the Community Right To Know List.

OSHA PEL: TWA 0.5 mg(As)/m³

THR: Poison by ingestion and inhalation. See also AR-SENIC COMPOUNDS. When heated to decomposition it emits very toxic fumes of As, Cl⁻ and SO_r.

DEP600	BENZENE	HR: 3
CAS: 95-50-1		NIOSH: CZ 4500000
v DOT : 1591		
mf: $C_{9}H_{4}Cl_{2}$	mw: 147.00	

PROP: Clear liquid. Mp: -17.5° , bp: 180-183°, fp: -22° , tlash p: 151°F. d: 1.307 @ 20°/20°, vap d: 5.05, autoign temp: 1198°F. lel: 2.2%, uel: 9.2%.

SYNS: CHLOROBEN CHLORODEN CLOROBEN DCB >-DICHLORBENZENE J-DICHLOR BENZOL 1.2 DICHLOROBENZENE DICHLOROBENZENE, ORTHO, liguid (DOT) DILANTIN DB DELATIN DB

DIZENE DOWTHERM E NC1-C54944 ODB ODCB ORTHODICHLOROBENZENE ORTHODICHLOROBENZOL RCRA WASTE NUMBER U070 SPECIAL TERMITE FLUID TERMITKIL.

TOXICITY DATA:	CODEN:
eye-rbt 100 mg/30S ms MLD	AMIHAB 17.180.58
spm-rat-ipr 250 mg/kg	JACTDZ 4(2),224.85
ihl-rat TCLo: 200 ppm/6H	FAATDF 5.190.85
(6-15D preg): TER	
ipr-rat TDLo: 50 mg/kg (1D	JACTDZ 4(1),224,85
male): REP	-
ori-rat LD50:500 mg/kg	WRPCA2 7.135,68
ihi-rat LCLo:821 ppm/7H	AMIHAB 17,180,58
ipr-rat LD50:840 mg/kg	MEPAAX 20.519,69
orl-mus LD50:4386 g/kg	YKYUA6 32.471.81
ivn-mus LDLo: 400 mg/kg	JPBAA7 44.281.37
orl-rbt LD50:500 mg/kg	85ARAE 3.32,76/77
ivn-rbt LDLo: 250 mg/kg	JPBAA7 44.281.37
orl-gpg LDLo: 2000 mg/kg	14CYAT 2,1336,63
uhl-gpg LCLo: 800 ppm/24H	JPBAA7 44,281,37

IARC Cancer Review: Human Inadequate Evidence IMEMDT 29,213,82; Animal Inadequate Evidence IMEMDT 29,213,82. Reported in EPA TSCA Inventory. Community Right To Know List.

OSHA PEL: CL 50 ppm ACGIH TLV: CL 50 ppm (skin) DFG MAK: 50 ppm (300 mg/m^3)

DOT Classification: ORM-A, Label: None; IMO: Poison B; Label: St. Andrews Cross

THR: Poison by ingestion and intravenous routes. Moderately toxic by inhalation and intraperitoneal routes. An eye, skin, and mucous membrane irritant. Causes liver and kidney injury. An experimental teratogen and suspected carcinogen. Experimental reproductive effects. Mutagenic data. A pesticide. Flammable when exposed to heat or flame. Can react vigorously with oxidizing materials. To fight fire, use water, foam, CO₂ dry chemical. Slow reaction with aluminum may lead to explosion during storage in a sealed aluminum container. When heated to decomposition it emits toxic fumes of Cl⁻. See also BENZENE CHLO-RIDE and CHLORINATED HYDROCARBONS, ARO-MATIC.

DFF809 1,1-DICHLOROETHANE

hl-rat TCLo: 57 mg/m ⁻ 4H (22W pre): TER	GISAAAA 39(7).25.74
pieriek	
ori-rat LD50:1120 mg/kg	HYSAAV 32.349.67
orl-mus LD50:625 mg/kg	HYSAAV 32.349.67
hl-mus LCLo: 10 g m	GISAAA 20(8),19.55
skn-rbt LD50: 3890 me/kg	UCDS** 3/23/70
unr-mam LD50:807 mg/kg	GTPZAB 26(4).26.82

THR: Moderately toxic by ingestion, skin contact and possibly other routes. Mildly toxic by inhalation. An experimental teratogen. Other experimental reproductive effects by inhalation. When neated to decomposition it emits very toxic fumes of CIT. See also 1.2-DICHLOROETHANE: and CHLORINATED HYDROCARBONS, ALIPHATIC.

k	DFF809	H R : 3
*	1.1-DICHLOROETHANE CAS: 75-34-3 DOT: 2362 mf: C ₂ H ₄ Cl ₂ mw: 98.96	NIOSH: KI 0175000
	PROP: Lei: 5.6%. uei: 11.4%	
	SYNS: AETHYLDENCHLORD (GERMAN) CHLORINATED HYDROCHLORIC ETHER CHLORURE & ETHYLDENE (FRENCH) CLORURO DI ETILIDENE (ITAL- IAN) 1.1-DICHLOORETHAAN (DUTCH)	1 I-DICHLORAETHAN (GERMAN) I.I-DICHLORETHANE I.I-DICLOROETANO (ITALIAN) ETHYLIDENE CHLORIDE ETHYLIDENE DICHLORIDE NCI-CO4535 RCRA WASTE NUMBER U076
	TOXICITY DATA: ihl-rat TCLo: 6000 ppm/7H (6-15D preg): TER orl-mus TDLo: 185 g/kg/78 W-I: ETA orl-mus TD : 1300 g/kg/78 W-I: ETA orl-rat LD50: 725 mg/kg	CODEN: TXAPA9 28,452.74 NCTTR* NCI-CG-TR- 66.78 NCTTR* NCI-CG-TR- 66.78 HYSAAV 32,349.67

EPA TSCA Chemical Inventory. NCI Carcinogenesis Bioassay (gavage); Inadequate Studies: mouse, rat NCITR* NCI-CG-TR-66,78.

OSHA PEL: TWA 100 ppm

DOT Classification: Flammable Liquid; Label: Flammable Liquid

THR: Moderately toxic by ingestion. An experimental tumorigen and teratogen. A suspected carcinogen. When heated to decomposition it emits very toxic fumes of Cl⁻. See also 1.2-DICHLOROETHANE: and CHLORINATED HYDROCARBONS, ALIPHATIC.

D FF900		HR: 3
1,2-DICHLORO	ETHANE	
CAS: 107-06-2		NIOSH: KI 0525000
DOT: 1184		
$mf: C_2H_4Cl_2$	mw: 98.96	

PROP: Colorless liquid, pleasant odor, sweet taste. Bp: 83.5°, ulc: 60-70, lel: 6.2%, uel: 15.9%, fp: -35.7°, flash p: 56°F. d: 1.257 @ 20°/4°, autoign temp: 775°F, vap press: 100 mm (a. 29.4°, vap d: 3.35.

SYNS:

511.0.
AETHYLENCHLORID (GERMAN)
BICHLOROETHANE
BICHLORURE D'ETHYLENE
FRENCH
BORER SOL
BROCIDE
CHLORURE D'ETHYLENE
FRENCH
CLORURO DI ETHENE (ITALIAN)
1.2-DCE
DESTRUXOL BORER-SOL
1.2-DICHLOORETHAAN (DUTCH)
1.2-DICHLOR-AETHAN (GERMAN)
DICHLOREMULSION
DI-CHLOR-MULSION
DICHLORO-1.2-ETHANE (FRENCH)
a.B-DICHLOROETHANE

TOXICITY DATA:

skn-rbt 600 mg open MLD eye-rbt 63 mg SEV mmo-sat 40 µmol/plate msc-hmn: lym 100 mg/L sit-mus-ipr 300 mg/kg otr-ham: emb 200 µL/plate ihl-rat TCLo: 300 ppm/7H (6-15D preg): REP orl-rat TDLo: 5286 mg/kg/69W-I:CAR ihl-rat TCLo: 5 ppm/7H/78W-I: ETA ori-mus TDLo: 3536 mg/kg/78W-I:CAR ihl-mus TCLo:5 ppm/7H/78W-I: ETA skn-mus TDLo: 1120 g/kg/74W-I:NEO orl-rat TD :38 g/kg/78W-I:CAR 55.78 orl-mus TD :76 g/kg/78W-I: CAR, TER 55,78 ori-rat TD :18 g/kg/78W-I:CAR 55,78 ori-mus TD : 38 g/kg/78W-I: 55,78 CAR.TER ihl-hmn TCLo: 4000 ppm/H: CNS.PNS.GIT orl-hmn TDLo: 428 mg/kg: GTT.CNS.PUL orl-man TDLo:892 mg/kg: GIT.LIV ori-hma LDLo: 286 mg/kg: **GIT,LIV** orl-man LDLo:714 mg/kg: CNS,CVS,PUL ori-rat LD50:670 mg/kg ihl-rat LC50:1000 ppm/7H scu-rat LDLo:99 mg/kg ori-mus LD50:489 mg/kg

1.1. DICHLOROFTHANE DICHLOROETHYLENE 2 DICLOROETANO (ITALIAN) DUTCH LIOUD DUTCH OIL EDC ENT 1.656 ETHANE DICHLORIDE ETHYLEENDICHLORIDE (DUTCH) ETHYLENE CHLORIDE ETHYLENE DICHLORIDE (ACGIH. DOD 1.2-ETHYLENE DICHLORIDE GLYCOL DICHLORIDE NCI-C00511 RCRA WASTE NUMBER 0077

CODEN: UCDS** 3/23/70 UCDS** 3/23/70 CBINA8 20.1.78 MUREAV 142,133,85 MUREAV 117,201,83 EVSRBT 25,75,82 BANRDU 5,149,80 BANRDU 5.35,80 BANRDU 5,3,80 BANRDU 5,35,80 BANRDU 5,3,80 JJIND8 63,1433,79 NCITR* NCI-CG-TR-NCITR* NCI-CG-TR-NCITR* NCI-CG-TR-NCITR* NCI-CG-TR-PCOC** -,500,66 SOMEAU 22,132,58 WILEAR 28,983,75 CLCEAL 86,203,47 KLWOAZ 48.822,70 FMCHA2 -. C99.83 AMIHBC 4,482,51 AMPLAO 51.346,51 TOXID9 1.26.81

1148

DFH600).O-DI(2-CHLOROETHY METHYLCOUMARIN-1		SYNS: CHLORURE de VINY FRENCH)
CAS: $321-55-1$ nt: $C_{14}H_{14}C_{13}O_{k}P$ mw:	NIOSH: GN 5250000	TEDCE TEDICHLOROETHE NCI-C54262 RCRA WASTE NUMI
SYNS: DO-BIS(2 CHLOROETHYL::OD) HLORO::4:METHYL TCOL MARINYL: PHOSPHATE CHLOROETHANOL HYDROGEN PHOSPHATE ESTER with 3- CHLORO:T:HYDROXY:4- METHYLCOUMARIN CHLOROETHANOL PHOSPHATE DIESTER ESTER with 3-CHLORO- THYDROXY:4-METHYLCOLMA- RIN CHLORO:T:HYDROXY:4- METHYLCOUMARIN BIS(2) CHLOROETHYL:PHOSPHATE CHLORO:4:METHYL-UMBELLI- PERONE BIS(2-CHLOROETHYL) PHOSPHATE FOXICITY DATA:	DI-(2-CHLOROETHYL)-3-CHLORO- A-METHYL, T-COUMARINYL PHOSPHATE DI-(2-CHLOROETHYL)-3-CHLORO- A-METHYLCOUMARIN-T-YL PHOSPHATE EUSTIDIL GALLOXON GALOXANE 30H60 HALOXON HELMIRANE HELMIRON HELMIRONE LOXON LUXON CODEN:	TOXICITY DA mmo-sat 5 pph mma-sat 3 pph/2H dnd-rat-ihl 10 ppr dns-mus-ihl 50 pp dns-mus-orl 200 r orl-rat TDLo: 200 preg): TER ihl-rat TCLo: 80 preg): TER ihl-rat TCLo: 55 pre): REP ihl-rat TCLo: 55 pre): REP ihl-rat TCLo: 55 skn-mus TDLo: 4 NEO ihl-rat TC: 150 pp
Jni-hmn: oth 10 mg/L orl-rat LD50: 900 mg/kg pr-ckn LD50: 800 mg/kg orl-dom LD50: 763 mg/kg	TTEHD6 10.143.82 FAZMAE 17.108.73 BCPCA6 16.1183.67 AJVRAH 41.1857.80	ETA ihl-mus TC:55 p ETA ihl-mus TC:55 p ETA ihl-rat TC:55 pp

THR: Moderately toxic by ingestion and intraperitoneal routes. Human mutagenic data. An anthelmintic. When neated to decomposition it emits very toxic fumes of PO, and Cl⁻. See also other coumarin entries.

D FH800	HR: 2
DICHLOROETHYLENE	
CAS: 25323-30-2	NIOSH: KV 9250000
DOT: 1150	
mf: $C_2H_2Cl_2$ mw: 96.94	
TOXICITY DATA:	CODEN:
hl-mus LCLo: 76 g/m ^{-/} 2H	AEXPBL 83,235,18

DOT Classification: Flammable Liquid; Label: Flammable Liquid

AEXPBL 83.235.18

UGLAAD 121.375.59

ihl-gpg LCLo: 155 g/m³/1H

orl-mam LDLo: 2500 mg/kg

THR: Moderately toxic by ingestion. Mildly toxic by inhalation. Flammable when exposed to heat or flame. When heated to decomposition it emits toxic fumes of Cl⁻. See also 1.1-DICHLOROETHYLENE.

D F1000		J.	HR: 3
1,1-DICHLORO	DETHYLENE -	ҝ	
CAS: 75-35-4		NIOSH: K	V 9275000
mf: $C_2H_2Cl_2$	mw: 96.94		

PROP: Colorless, volatile liquid. Bp: 31.6°, lel: 7.3%, uel: 16.0%, fp: -122°, flash p: 0°F (OC), d: 1.213 @ 20°/4°, autoign temp: 1058°F.

21.92:	
CHLORURE de VINYLIDENE	SCONATEX
FRENCH)	VDC
T DCE	VINYLIDENE CHLORIDE (II)
DICHLOROETHENE	VINYLIDENE CHLORIDE ACGIH)
NCI-C54262	VINYLIDENE DICHLORIDE
RCRA WASTE NUMBER U078	VINYLIDINE CHLORIDE
TOXICITY DATA:	CODEN:
mmo-sat 5 pph	MUREAV 57.141,78
mma-sat 3 pph/2H	MUREAV 58.183.78
dnd-rat-ihi 10 ppm	TXAPA9 53.357.80
dns-mus-ihi 50 ppm	TXAPA9 53.357.80
dns-mus-orl 200 mg/kg	TXCYAC 36.199.85
orl-rat TDLo: 200 mg/kg (6-15D	TXAPA9 49.189.79
preg): TER	
ihi-rat TCLo: 80 ppmv7H (6-15D	TXAPA9 49.189.79
preg): TER	
ihl-rat TCLo: 55 ppm/6H (55D	JTEHD6 3.965.77
pre):REP	
ihl-rat TCLo: 55 ppm/6H/52W-I:	JTEHD6 4.15.78
ETA	
ihl-mus TCLo: 25 ppm/4H/52W-I	MELAAD 68.241.77
skn-mus TDLo: 4840 mg/kg:	JJIND8 63.1433.79
NEO	
ihl-rat TC:150 ppm/4H/52W-I:	MELAAD 68.241.77
ETA	
ihl-mus TC: 55 ppm/6H/13W-I:	JTEHD6 7,909,81
ETA	
ihl-mus TC:55 ppm/6H/52W-I:	EVHPAZ 21.25,77
ETA	
ihl-rat TC:55 ppmv6H/52W-I:	EVHPAZ 21.25.77
ETA	
ihi-rat TC:150 ppm/4H/52W-I:	EVHPAZ 21.45.77
ETA	
thi-rat TC:55 ppm/6H/28W-I:	JTEHD6 7.909,81
ETA	
ihl-hmn TCLo:25 ppm:	CHINAG (11),463,76
CNS,LIV.KID	
orl-rat LD50: 200 mg/kg	DCTODJ 1.63.77
ihl-rat LCLo: 10000 ppm/24H	EXMPA6 20,187,74
ihl-mus LC50:98 ppm/22H	JTEHD6 3.913.77
orl-dog LDLo: 5750 mg/kg	QJPPAL 7.205.34
ivn-dog LDLo: 225 mg/kg	QJPPAL 7.205.34
scu-rbt LDLo: 3700 mg/kg	QJPPAL 7.205.34

LARC Cancer Review: Human Inadequate Evidence IMEMDT 39,195.86; Animal Sufficient Evidence IMEMDT 19,439,79; Animal Limited Evidence IMEMDT 39,195,86; Human Inadequate Evidence IMEMDT 19,-439,79. EPA Genetic Toxicology Program. Reported in EPA TSCA Inventory. Community Right To Know List.

ACGIH TLV: TWA 5 ppm; STEL 20 ppm

THR: Poison by inhalation, ingestion and intravenous routes. Moderately toxic by subcutaneous route. An experimental carcinogen, neoplastigen, tumorigen and teratogen. Human systemic effects by inhalation: general anesthesia, liver and kidney changes. Experimental reproductive effects. Mutagenic data. See also VINYL CHLORIDE. A very dangerous fire hazard when exposed to heat or flame. Moderately explosive in the form of gas when exposed to heat or flame. It forms explosive peroxides upon exposure

DF200 CIS-DICHLOROETHYLENE

to air. Potentially explosive reaction with chlorotrifluoroethylene at 180°C. Reaction with ozone forms dangerous products. Explosive reaction with perchloryl fluoride when heated above 100°C. Also can explode spontaneously. Reacts violently with chlorosulfonic acid: HNO₃; oleum. Can react vigorously with oxidizing materials. To fight fire, use alcohol foam, CO₂, dry chemical. When heated to decomposition it emits toxic fumes of Cl⁻.

 DF1200
 HR: 1

 cis-DICHLOROETHYLENE

 CAS: 156-59-2
 NIOSH: KV 9420000

 mf: C3H3Cl3
 mw: 96.94

HCC1=CHC1

PROP: Colorless liquid, pleasant odor. Mp: -80.5°, bp: 59°, lel: 9.7%, uel: 12.8%, flash p: 39°F, d: 1.2743 @ 25°/4°, vap press: 400 mm @ 41.0°, vap d: 3.34.

SYN: 12 DICHLOROETHYLENE

TOXICITY DATA:	CODEN:
mmo-smc 100 mmol/L	TCMUD8 4.365.84
mma-smc 40 mmoi/L	TCMUD8 4,365.84
mrc-smc 100 mmol/L	TCMUD8 4.365.84
dns-rat: lvr 4300 µmol/L	CRNGDP 5.1629.84
ihl-mus LCLo:65000 mg/ m ³ /2H	AHBAAM 116,131,36
ihl-cat LCLo: 20000 mg/ m ³ /6H	AHBAAM 116.131.36

Reported in EPA TSCA Inventory.

THR: Mildly toxic by by ingestion and inhalation. In high concentration it is irritating and narcotic. Has produced liver and kidney injury in experimental animals. A suspected carcinogen. Mutagenic data. Sometimes thought to be nonflammable, however, it is a dangerous fire hazard when exposed to heat or flame. Reaction with solid caustic alkalies or their concentrated solutions produces chloracetylene gas which ignites spontaneously in air. Reacts violently with N₂O₄; KOH; Na; NaOH. Moderate explosion hazard in the form of vapor when exposed to flame. Can react vigorously with oxidizing materials. To fight fire, use water spray, foam, CO₂, dry chemical. When heated to decomposition it emits toxic fumes of Cl⁻. See also 1,1-DICHLOROETHYLENE and CHLORINATED HY-DROCARBONS, ALIPHATIC. For further information. see Vol. 4, No. 3 of DPIM Report.

DF1600

HR: 1

CAS: 156-60-5 NIOSH: KV 9400000 mf: C₂H₂Cl₂ mw: 96.94

PROP: Flash p: 35.6°F, lel: 9.7%, uel: 12.8%.

SYN: TARE-ACETYLENE DICHLORIDE TARE-DICHLOROETHYLENE

trans-1,2-DICHLOROETHYLENE

 TOXICITY DATA:
 CODEN:

 :hl-hmn TCLo: 4800 mg/m³/10M
 AHBAAM 116,131,36

 ipr-rat LD50: 7536 mg/kg
 TXCYAC 7,141,77

 thl-mus LCLo: 75000 mg/m3/2H
 AHBAAM 116,131,36

 ipr-mus LD50: 4019 mg/kg
 TXCYAC 7,141,77

 thl-cat LCLo: 43000 mg/m3/6H
 AHBAAM 116,131,36

Reported in EPA TSCA Inventory.

THR: Mildly toxic to humans by inhalation. Mildly toxic experimentally by inhalation and intraperitoneal routes. A very dangerous fire hazard when exposed to heat or flame. Violent reaction with diffuoromethylene dihypofluorite. Forms shock-sensitive explosive mixtures with dinitrogen tetraoxide. Reaction with solid caustic alkalies or their concentrated solutions produces chloracetylene gas which ignites spontaneously in air. Reacts violently with N_2O_4 ; KOH; Na; NaOH. Moderate explosion hazard in the form of vapor when exposed to flame. Can react vigorously with oxidizing materials. To fight fire, use water spray, foam, CO_2 , dry chemical. When heated to decomposition it emits toxic fumes of Cl⁻. See also 1.1-DICHLOROETHYLENE and CHLORINATED HYDROCARBONS. ALIPHATIC. For further information, see Vol. 4, No. 3 of DPIM Report.

DF1800 HR: 3 1,2-DICHLOROETHYLENE CARBONATE CAS: 3967-55-3 NIOSH: JH 7400000

CH0. 3701-33-3		
mf: $C_3H_2Cl_2O_3$	mw: 156.95	

SYN: 4.5-DICHLORO-2-OXO-1,3-DIOXOLANE

TOXICITY DATA:	CODEN:
scu-mus TDLo:648 mg/kg/54W-	JNCIAM 48,1431,72
I:ETA	

THR: An experimental tumorigen. When heated to decomposition it emits toxic fumes of Cl^{-} .

DF J000 HR: 3 DICHLORO(ETHYLENEDIAMMINE)PLATINUM(II) CAS: 14096-51-6 NIOSH: TP 2497100

 $mf: C_2H_8Cl_2N_2Pt \qquad mw: 326.11$

SYNS: ethylenediaminedichloride platinum (II)	PLATINUM ETHYLENEDIAMMINE DICHLORIDE
TOXICITY DATA:	CODEN:
mmo-sat 2 µg/plate	MUREAV 77.45.80
mma-sat 2 µg/plate	MUREAV 77.45.80
dnd-esc 100 mmol/L	CBINA8 16,39.77
dni-hmn: oth 25 µmol/L	UCNAW 6,207,70
oms-mam:lym 10 µmol/L	BCPCA6 23,1659.74
ipr-mus LDLo: 14 mg/kg	BCPCA6 2,187,73

THR: Poison by intraperitoneal route. Human mutagenic data. See also PLATINUM COMPOUNDS. When heated to decomposition it emits very toxic fumes of Cl^- and NO_r .

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TOXICITY DATA: ms-rat TDLo: 50 mg/kg:ETA CODEN: CNREA8 29,506.69

THR: An experimental tumorigen. When heated to decomposition it emits acrid smoke and irritating fumes.

EG**P**500 ETHYL BENZENE CAS: 100-41-4 DOT: 1175 mf: C.H₁₀ mw: 106.18 HR: 2

NIOSH: DA 0700000

PROP: Colorless liquid, aromatic odor. Misc in alcohol and ether, insol in NH₃; sol in SO₂. Bp: 136.2°, fp: -94.9°, flash p: 59°F, d: 0.8669 @ 20°/4°, autoign temp: 810°F, vap press: 10 mm (a) 25.9°, vap d: 3.66, lel: 1.2%, uel: 6.8%.

SYNS:

ETILBENZENE (ITALIAN)
ETYLOBENZEN (POLISH)
NCT-C56393
PHENYLETHANE
CODEN

TOXICITY DATA:	CODEN:
skn-rbt 15 mg/24H open MLD	AIHAAP 23.95.62
eye-rbt 100 mg	AJOPAA 29,1363,46
sce-hmn:lym 1 mmol/L	MUREAV 116,379,83
ihi-rat TCLo:97 ppm/7H (15D	BATTL* JAN.81
preg): REP	
ihl-rat TCLo: 985 ppm/7H	BATTL* JAN,81
(1-19D preg): TER	
ihl-rat TCLo:96 ppm/7H (1-19D	BATTL* JAN,81
preg): TER	
ihl-hma TCLo: 100 ppm/8H:	AIHAAP 31,206,70
EYE.CNS.PUL	
orl-rat LD50:3500 mg/kg	AMIHAB 14.387.56
ihl-rat LCLo: 4000 ppm/4H	AIHAAP 23,95,62
ihl-mus LCLo: 50 g/m3/2H	GTPZAB 5(5),3.61
ipr-mus LD50: 2272 mg/kg	ARTODN 58,106.85
skn-rbt LD50:17800 mg/kg	FCTXAV 13,803.75
ihl-gpg LCLo: 10000 ppm	PHRPA6 45.1241.30

Reported in EPA TSCA Inventory. EPA Genetic Toxicology Program. Community Right To Know List.

OSHA PEL: TWA 100 ppm (skin)

- ACGIH TLV: TWA 100 ppm; STEL 125 ppm; BEI: 2 g/L (mandelic acid in urine at end of shift) DFG MAK: 100 ppm (440 mg/m³)
- DOT Classification: Flammable Liquid, Label: Flammable Liquid

THR: Moderately toxic by ingestion and intraperitoneal route. Mildly toxic by inhalation and skin contact. An experimental teratogen. Human systemic effects by inhalation: eye, sleep and pulmonary changes. An eye and skin irritant. Human mutagenic data. The liquid is an irritant to the skin and mucous membranes. A concentration of 0.1% of the vapor in air is an irritant to human eyes, and a concentration of 0.2% is extremely irritating at first, then causes dizziness, irritation of the nose and throat and a sense of constriction in the chest. Exposure of guinea pigs to 1% concentration has been reported as causing ataxia, loss of consciousness, tremor of the extremities and finally death through respiratory failure. The pathological findings were congestion of the brain and lungs with edema. No data are available regarding the effect of chronic exposure.

A very dangerous fire and explosion hazard when exposed to heat or flame; can react vigorously with oxidizing materials. To fight fire, use foam, CO2, dry chemical. When heated to decomposition it emits acrid smoke and irritating fumes. For further information, see Vol. 2, No. 6 of DPIM Report.

EGQ000 a-ETHYLBENZENEMETH	HR: 2
CAS: $93-54-9$ mf: $C_9H_{12}O$ mw: 136.21	NIOSH: DO 5470000
SYNS: ejtbil a-ethylbenzyl alcohol ethyl phenyl carbinol felicur felitrope fenicol a-hydroxypropylbenzene Livonal	PHENICOL PHENYCHOLON PHENYLAETHYLCARBINOL (GER- MAN) 1-PHENYLPROPANOL 1-PHENYL-1-PROPANOL 1-PHENYLPROPYL ALCOHOL SH 261
TOXICITY DATA: orl-rat LD50:1600 mg/kg orl-mus LD50:500 mg/kg scu-mus LD50:700 mg/kg	CODEN: ARZNAD 12,347,62 AIPTAK 116,154,58 AIPTAK 116,154,58

THR: Moderately toxic by ingestion and subcutaneous route. When heated to decomposition it emits acrid smoke and irritating fumes.

<i>EGR000</i> ETHYL BENZO	ነ ልፐፑ	H R : 2
CAS: 93-89-0		NIOSH: DH 0200000
mf: $C_9H_{10}O_2$	mw: 150.19	
DDOD: Coloriers	aromatic liquid	Mo: - 34 6° bo: 213 4°

PROP: Colorless, aromatic liquid. Mp: -34.6°, bp: 213.4°, flash p: >204°F, d: 1.048 @ 20°/20°, vap press: 1 mm @ 44.0°, vap d: 5.17, autoign temp: 914°F. Insol in water; misc in petroleum, alcohol, chloroform, and ether.

SYNS: BENZOIC ETHER

ESSENCE OF NIOBE

TOXICITY DATA:	CODEN:
ska-rbt 10 mg/24H open MLD	AMIHBC 10,61,54
eye-rbt 500 mg open	AMIHBC 10.61.54
ori-rat LD50:2100 mg/kg	JPETAB 84.358.45
skn-cat LDLo: 10 g/kg	JPETAB 84,358,45
ori-rot LD50:2630 mg/kg	JPETAB 84,358,45

Reported in EPA TSCA Inventory.

EHH000 ETHYL CHLORIDE CAS: 75-00-3 DOT: 1037 mf: C₂H₃Cl mw: 64.52

HR: 1 NIOSH: KH 7525000

ALIAN)

PROP: Colorless liquid or gas; ether-like odor. sol in water at 0.45; misc in alcohol and ether, burning taste. Bp: 12.3°, lel: 3.8%, uel: 15.4%, fp: -139°, flash p: -58°F (CC), d: 0.9214 (ā 0°/4°, autoign temp: 966°F, vap press: 1000 mm $(\tilde{a} 20^{\circ} \text{ vap d}; 2.22)$.

SYNS:

AETHYLCHLORID (GERMAN)	CLOROETANO (ITALIAN)
AETHYLIS	CLORURO DI ETILE (ITALIA
AETHYLIS CHLORIDUM	ETHER CHLORATUS
ANODYNON	ETHER HYDROCHLORIC
CHELEN	ETHER MURIATIC
CHLOORETHAAN (DUTCH)	ETYLU CHLOREK (POLISH)
CHLORETHYL	HYDROCHLORIC ETHER
CHLORIDUM	KELENE
CHLOROAETHAN (GERMAN)	MONOCHLORETHANE
CHLOROETHANE	MURIATIC ETHER
CHLORURE D'ETHYLE (FRENCH)	NARCOTTLE
CHLORYL	NCI-C06224
CHLORYL ANESTHETIC	

TOXICITY DATA:	CODEN:
ihl-rat LC50:160 g/m ³ /2H	85GMAT - ,66,82
ihl-mus LC50: 146 g/m ³ /2H	85GMAT -,66,82
ihl-gpg LCLo: 40000 ppm/45M	XPHBAO 185,1,29

Reported in EPA TSCA Inventory. Community Right To Know List.

OSHA PEL: TWA 1000 ppm ACGIH TLV: TWA 1000 ppm

DOT Classification: Flammable Liquid, Label: Flammable Liquid; Flammable Gas; Label; Flammable Gas

THR: Mildly toxic by inhalation. An irritant to skin, eyes and mucous membranes. The liquid is harmful to the eyes and can cause some irritation. In the case of guinea pigs, the symptoms attending exposure are similar to those caused by methyl chloride, except that the signs of lung irritation are not as pronounced. It gives some warning of its presence because it is irritating, but it is possible to tolerate exposure to it until one becomes unconscious. It is the least toxic of all the chlorinated hydrocarbons. It can cause narcosis, although the effects are usually transient. A priority pollutant.

A very dangerous fire hazard when exposed to heat or flame; can react vigorously with oxidizing materials. Severe explosion hazard when exposed to flame. Reacts with water or steam to produce toxic and corrosive fumes. Incompatible with potassium. To fight fire, use carbon dioxide. When heated to decomposition it emits toxic fumes of phosgene and Cl⁻. See also CHLORINATED HYDROCARBONS. ALIPHATIC. For further information, see Vol. 1, No. 4 of DPIM Report.

FHH500

ETHYL CHLORO BENZENE

CAS: 1331-31-3

mw: 140.62 $mf: C_{*}H_{0}Cl$

PROP: Clear. colorless liquid. Mp: -62.6°, bp: 184.3°, flash p: 147°F, d: 1.05 @ 25°/25°, vap press: 1 mm @ 19.2°, vap d: 4.86.

SYN: CHLOROETHYLBENZENE

TOXICITY DATA: CODEN: skn-rbt 10 mg/24H open JIHTAB 30.63.48 eye-rbt 500 mg AJOPAA 29.1363,46 orl-rat LD50: 5000 mg/kg JIHTAB 30.63.48 skn-rbt LD50:18 g/kg **JIHTAB 30.63.48**

THR: Mildly toxic by ingestion and skin contact. A skin and eye irritant. Flammable when exposed to heat or flame; can react vigorously with oxidizing materials. To fight fire, use foam, CO₂, dry chemical. When heated to decomposition it emits acrid smoke and irritating fumes. See also CHLORINATED HYDROCARBONS, AROMATIC, and CHLOROBENZENE.

EHI500

7-(2-(ETHYL-2-CHLOROETHYL)AMINOETHYL-AMINO)BENZ(c)ACRIDINE DIHYDROCHLO-RIDE

CAS: 4310-69-4	NIOSH: CU 3440000
mf: $C_{23}H_{24}ClN_3 \cdot 2ClH$	mw: 450.87
SYNS:	
N'-BENZ(c)ACRIDEN-7-YL-N-(2-	7-12-22-CHLOROETHYLEHTYL
CHLOROETHYL - N-ETHYL-1.2-	AMINO)ETHYLAMINO)BENZ(c)
ETHANEDIAMINE DIHYDRO-	ACRIDINE DIHYDROCHLORIDE
CHLORIDE	ICR 311
TOXICITY DATA:	CODEN:
ivn-mus TDLo:4500 µg/kg: NEO	CNREA8 36.2423.76

ive-mus LDLo:4500 µg/kg CNREA8 36,2423,76

THR: Poison by intravenous route. An experimental neoplastigen. When heated to decomposition it emits very toxic fumes of NO_x and Cl^- .

EH1000

7-(3-(ETHYL-2-(CHLOROETHYLAMINO)PROPYL-AMINO))BENZ(c)ACRIDINE DIHYDROCHLO-RIDE

CAS: 4251-89-2	NIOSH: CU 3460000
mf: C ₂₄ H ₂₆ ClN ₃ •2ClH	mw: 464.90

SYN: ICR 292

TOXICITY DATA:	CODEN:
mmo-sat 500 ng/plate	MUREAV 136,185.84
pic-esc 60 ng/plate	CNREA8 43.2819.83

HR: 1

HR: 3

HR: 3

NIOSH: CZ 0700000

H⁻T500 HR: 3 2. 6.2',4'.6'-HEXANITRODIPHENYLAMINE CAS: 131-73-7 NIOSH: JJ 9275000 I SO 24 I SO 24

 $m^{f_1}C_{12}H_5N-O_{12}$ mw: 439.24

S NS:	
BLS(2.4.6-TRINITRO-PHENYL)	HEXANTTRODIPHENYLAMINE
AMIN (GERMAN) D	HEXANITRODIPHENYLAMINE
E. NITRODIFENILAMINA IITAL-	2.2.4.4 .6.6 HEXANITRODIPHE-
LAN)	NYLAMINE
HEXANITRODIFENYLAMINE	HEXYL (GERMAN, DUTCH)
(UTCH)	
1 UXICITY DATA:	CODEN:
mmo-sat 228 nmol/plate	MUREAV 136.209.84
r. ia-sat 456 nmol plate	MUREAV 136,209.84
c rat TDLo: 14 g/kg/76W-C: NEO	NATUAS 180,509.57

P-ported in EPA TSCA Inventory.

IR: An experimental neoplastigen. Mutagenic data. A powerful and violent explosive used as a booster explosive; i use is superior to TNT. It is not as good for this purpose i tetryl, but is extremely stable and much safer to handle. See also NITRO COMPOUNDS OF AROMATIC HY-DROCARBONS.

HET675 HR: 3 "EXANITROETHANE AS: 918-37-6 mf: C₂N₆O₁₂ mw: 300.06

THR: A powerful oxidant which explodes above 140°C. xplosive reaction with boron. Hypergolic reaction with dimethyl hydrazine or other strong organic bases. Forms rowerfully explosive mixtures with nitrogen containing or-

anic compounds (e.g., 2-nitroaniline). Upon decomposiuon it emits toxic fumes of NO_x . See also NITRO COM-POUNDS.

HEU000 HEXANOIC ACID	H R : 2
AS: 142-62-1	NIOSH: MO 5250000
JOT : 1706	
mf: $C_6H_{12}O_2$ mw: 116.1	8

'ROP: Oily, colorless liquid; odor of Limburger cheese. Jp: 205.0°, fp: -3.4° , flash p: 215°F (COC), d: 0.9295 @ 20°/20°, vap press: 0.18 mm @ 20°, vap d: 4.0, autoign emp: 716°F. Slightly sol in water; very sol in ethanol, :ther.

SYNS:

JUTYLACETIC ACID
CAPROIC ACID
n-CAPROIC ACID
CAPRONIC ACID
HEXACID 698

n-HEXANOIC ACID n-HEXOIC ACID PENTIPORMIC ACID PENTYLFORMIC ACID TOXICITY DATA: skn-rbt 10 mg/24H open MLD skn-rbt 465 mg open MLD eye-rbt 695 µg SEV oms-nmi:oth 10 mmol/L cyt-nmi:oth 10 mmol/L orl-rat LD50: 3000 mg/kg orl-mus LD50: 5 g/kg ihl-mus LC50: 4100 mg/m³/2H 1pr-mus LD50: 3180 mg/kg scu-mus LD50: 3180 mg/kg

skn-gpg LD50:4635 mg/kg

mf: $C_8H_{14}O_2$

2-HEXANONE HEVOOD

CODEN: AMIHBC 10.61.54 UCDS** 11/2/71 AJOPAA 29.1363.46 CHROAU 40.1.73 CHROAU 40.1.73 JIHTAB 26.269.44 85GMAT -.32.82 JPPMAB 21.85.69 JPPMAB 21.85.69 AMIHBC 10.61.54 JIHTAB 26.269.44

Reported in EPA TSCA Inventory.

DOT Classification: Corrosive Material; Label: Corrosive

THR: Moderately toxic by ingestion, skin contact, intraperitoneal, and subcutaneous routes. Mutagenic data. Corrosive. A skin and severe eye irritant. Combustible when exposed to heat or flame; can react with oxidizing materials. To fight fire, use CO_2 , dry chemical, fog, mist. When heated to decomposition it emits acrid smoke and fumes.

HEU500 HR: 1 HEXANOIC ACID, VINYL ESTER (MIXED ISO-MERS)

NIOSH: MO 8450000

EN:
AP 23.95.62
AP 23.95,62
AP 23.95,62

mw: 142.22

THR: Mildly toxic by ingestion and inhalation. A skin irritant. When heated to decomposition it emits acrid smoke and fumes. See also ESTERS.

HEV000 HR: 3 2-HEXANONE K CAS: 591-78-6 NIOSH: MP 1400000 mf: C₆H₁₂O mw: 100.18

PROP: Clear liquid. Mp: -56.9° , bp: 127.2°, lel: 1.22%, uel: 8.0%, flash p: 95°F (OC), d: 0.830 @ 0°/4°, vap press: 10 mm @ 38.8°, vap d: 3.45, autoign temp: 991°F. Slightly sol in H₂O; sol in alc, ether.

SYNS: BUTYL METHYL KETONE METHYL B-BUTYL KETONE (ACGIH) -BUTYL METHYL KETONE HEXANONE-2 MNBK MBK CODEN: TOXICITY DATA: AMIHBC 10,61,54 eye-rbt 100 mg open EESADV 5.291.81 ihl-rat TCLo: 1000 ppm/6H (1-21D preg): TER EESADV 5.291.81 ihl-rat TCLo: 2000 ppm/6H (1-21D preg): REP NPIRI* 1,78,74 ihl-hmn TCLo: 1000 ppm: EYE.CNS.GIT

orl-rat LD50: 2590 mg/kg	AMIHBC 10.61.54
thl-rat LC50:8000 ppm/4H	NPIRI* 1.78.74
pr-rat LDLo:914 mg/kg	RaIRL# 01MAR74
ori-mus LDLo: 1000 mg/kg	UCPHAQ 2.217.49
skn-rbt LD50:4800 mg/kg	NPIRI* 1.78.74
orl-gpg LDLo: 914 mg/kg	RaIRL# 01MAR74

Reported in EPA TSCA Inventory.

OSHA PEL: TWA 100 ppm ACGIH TLV: TWA 5 ppm (skin) NIOSH REL: (Ketones) TWA 4 mg/m³

THR: Moderately toxic by ingestion and intraperitoneal routes. Mildly toxic by inhalation and skin contact. Experimental teratogenic and reproductive effects. Human systemic effects by inhalation: unspecified eye effects, headache, nausea or vomiting. An eye irritant. Dangerous fire and explosion hazard when exposed to heat or flame; can react with oxidizing materials. To fight fire, use alcohol foam. CO₂, dry chemical. See also KETONES.

HEV500	HR: 2
3-HEXANONE	
CAS: 589-38-8	NIOSH: MP 1576000
mf: C ₆ H ₁₂ O mw: 100.18	
PROP: Colorless liquid Bp: 124	e d 0 813 @ 21 8°/4°

PROP: Coloriess líquía. Bp: 124°, d: 0.813 @ 21.874°, flash p: 57.2°F (OC).

SYNS: AETHYLPROPYLKETON (GER-ETHYL PROPYL KETONE MAN TOXICITY DATA: CODEN: TXAPA9 28.313.74 orl-rat LD50: 3360 mg/kg ihl-rat LCLo: 4000 ppm/4H TXAPA9 28,313,74 skn-rbt LD50:3170 mg/kg TXAPA9 28.313.74

Reported in EPA TSCA Inventory.

scu-gpg LDLo: 700 mg/kg

THR: Moderately toxic by ingestion, skin contact and subcutaneous routes. Mildly toxic by inhalation. A very dangerous fire hazard when exposed to heat or flame; can react vigorously with oxidizing materials. To fight fire, use foam, CO₂, dry chemical. When heated to decomposition it emits acrid smoke and fumes. See also KETONES.

HEXANOYLETHYLENEIMINE

HEW000		HR: 3
1-HEXANOYLA	ZIRIDINE	
CAS: 45776-10-1	l	NIOSH: CM 7890000
mf: C ₈ H ₁₅ NO	mw: 141.24	
CVAIC.		

SYNS: I-CAPROYLAZIRIDINE CAPROYLETHYLENEIMINE

TOXICITY DATA CODEN:

IOMCITI DATA.	CODEN.
cyt-rat-ipr 50 mg/kg	BJPCAL 9,306,54
scu-rat TDLo: 495 mg/kg/19W-I:	BJPCAL 9,306,54
NEO	

THR: An experimental neoplastigen and tumorigen. Mutagenic data. When heated to decomposition it emits toxic fumes of NO...

HEY000 HR: 3 HEXAPYRIDINEIRON(II) TRIDECACARBONYL TETRAFERRATE(2⁻⁻)

CAS: 23129-50-2 mf: $C_{43}H_{30}Fe_5N_6O_{13}$ mw: 1117.9

SYN: HEXAKIS(PYRIDINE) IRON(II) TRIDECACARBONYL TETRAFER-RATE(2-)

THR: Ignites spontaneously in air. When heated to decomposition it emits toxic fumes of NOr. See also CAR-BONYLS.

HEXASODIUM TETRAPHOSPHATE

CAS: 14986-84-6 NIOSH: XF 1700000 mf: $Na_6O_{13}P_4$ mw: 469.82

SYNS:

HEY500

HEXANATRIUMTETRAPOLY-PHOSPHAT (GERMAN)

HEXASODIUM TETRAPOLY-PHOSPHATE

TOXICITY DATA: ori-mus LD50: 3920 mg/kg scu-mus LD50:875 mg/kg

CODEN: ARZNAD 7,445,57 ARZNAD 7,445,57

Reported in EPA TSCA Inventory.

THR: Moderately toxic by ingestion and subcutaneous routes. When heated to decomposition it emits toxic fumes of Na₂O and PO₇. See also PHOSPHATES.

HEZ000 1.3.5-HEXATRIENE

Line and the second sec		
CAS: 2235-12	2-3	NIOSH: MP 5425000
mf: C ₆ H ₈	mw: 80.14	

SYN: DIVINYLETHYLENE

TOXICITY DATA:	CODEN:
eye-rbt 369 mg	IHFCAY 6,1,67
ori-rat LD50:210 mg/kg	IHPCAY 6,1,67
ihl-rat LCLo: 100000 ppm/15M	IHPCAY 6,1,67
skn-rbt LD50:6730 mg/kg	IHPCAY 6,1,67

Reported in EPA TSCA Inventory.

THR: Poison by ingestion. Mildly toxic by skin contact and inhalation. An eye irritant. When heated to decomposition it emits acrid smoke and fumes.

HR: 2

HR: 3

• 3

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THR: A poison by ingestion, inhalation, and skin contact. a skin irritant. When heated to decomposition it emits a...d smoke and fumes. See also ACETYLENE COM-POUNDS and ALCOHOLS.

F 'F300	HR: 3
4-HEXEN-1-YN-3-ONE	
CAS: 13061-80-8	NIOSH: MQ 0350000
r : C ₂ H ₂ O mw: 94.12	

SYN: 4-HEXENE-1-YNE-3-ONE

DXICITY DATA:	CODEN:
 i-rbt 10 mg 24H open MLD 	AIHAAP 23.95.62
ori-rat LD50:71 mg/kg	AIHAAP 23.95.62
h1-rat LCLo: 13 ppm/4H	AIHAAP 23.95.62
h-rbt LD50:100 mg/kg	AIHAAP 23.95.62

HR: A poison by ingestion, inhalation, and skin contact. A skin irritant. When heated to decomposition it emits rid smoke and fumes. See also ACETYLENE COM-. JUNDS and KETONES.

FF500	H R : 3
EXOBENDINE DIHYDROCHLORIDE	

CAS: 50-62-4	NIOSH: DI 0230000
$f: C_{30}H_{44}N_2O_{10}\cdot 2ClH$	mw: 665.68
YNS:	

REOXYL ANDIAMINE N.N'-DIMETHYL-N.N.-BIS(3-(3')-4'.5'-TRIMETHOXYBENZOXY) PROPYL) ETHYLENEDIAMINE DEHYDROCHLORIDE **OXICITY DATA:** CODEN: -l-rat LD50:2550 mg/kg **GNRIDX 3.77.69** scu-rat LD50:930 mg/kg **GNRIDX 3.77.69 GNRIDX 3.77.69** ivn-rat LD50:52 mg/kg GNRIDX 3.77.69 rl-mus LD50:682 mg/kg :u-mus LD50:328 mg/kg **GNRIDX 3,77,69** ivn-mus LD50:35200 µg/kg **GNRIDX 3.77.69**

"HR: Poison by subcutaneous and intravenous routes. Modrately toxic by ingestion. When heated to decomposition it emits toxic fumes of NO_x and HCl. See also ESTERS.

1FG000		HR: 3
JEXOCYCLIUM		
CAS: 6004-98-4		NIOSH: TM 3150000
nf: $C_{1}H_{36}N_{2}O_{5}S$	mw: 428.61	

2ROP: Crystals. Mp: 200-210°, sltly sol in chloroform, insol in ether, sol in H_2O .

SYNS + B-CYCLOHEXYL-B-HYDROXY-TRAL PHENETHYL)-1,1-DIMETHYL PI-PERAZINIUM SULFATE

TOXICITY DATA:	CODEN:
mmo-sat 32 µg/plate	JEPTDQ 4.345.80
orl-mus LD50:600 mg/kg	27 ZIAQ -,-,65

pr-mus LD50:55 mg/kg scu-mus LD50:360 mg/kg ivn-mus LD50:11 mg/kg

27ZIAQ - (19.73 27ZIAQ - 119.73 27ZIAO - 119.73

THR: Poison by intraperitoneal, subcutaneous, and intravenous routes. Moderately toxic by ingestion. Mutagenic data. When heated to decomposition it emits very toxic fumes of NO_r and SO_r.

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HFG500 HEXONE CAS: 108-10-1 DOT: 1245 mf: $C_{5}H_{12}O$

HR: 3

NIOSH: SA 9275000

mw: 100.18

CH₃CO•CH₂CH(CH₃)₇

PROP: Clear liquid. Bp: 118°, lel: 1.4%, uel: 7.5%, flash p: 62.6°F, d: 0.803, fp: -80.2°, autoign temp: 858°F, vap press: 16 mm @ 20°, d: 3.45.

SYNS:	
HEXON (CZECH)	4-METHYL/PENTAN/2/ON
SOBUTYL METHYLKETON	(DUTCH. GERMAN)
CZECHI	2 METHYL + PENTANONE
ISOBUTYL METHYL KETONE	+METHYL-2-PENTANON (CZECH)
ISOPROPYLACETONE	4-METHYL-2-PENTANONE
METHYL ISOBUTYL CETONE	METILISOBUTILCHETONE (TTAL-
FRENCH	LAN
METHYLISOBUTYLKETON	4-METTLPENTAN-2-ONE (ITALIAN)
DUTCH, GERMAN)	MOBK
METHYL ISOBUTYL KETONE	MER
ACGIH.DOT)	RCRA WASTE NUMBER U161
METYLOIZOBUTYLOKETON (POL-	SHELL MUBK
ISH)	

CODEN:

JIHTAB 28.262,46

28ZPAK -.42.72

UCDS** 4/25/58

28ZPAK -.42.72

UCDS** 4/25/58

28ZPAK -.42.72

SCCUR* -.7.61

TOXICITY DATA: eye-hmn 200 ppm/15M skn-rbt 500 mg/24H MLD eye-rbt 40 mg SEV eve-rbt 500 mg/24H MLD orl-rat LD50:2080 mg/kg ihi-rat LC50:8000 ppm/4H ipr-mus LD50:268 mg/kg

Community Right To Know List.

OSHA PEL: TWA 100 ppm ACGIH TLV: TWA 50 ppm; STEL 75 ppm NIOSH REL: (Ketones) TWA 200 mg/m³

DOT Classification: Flammable Liquid: Label: Flammable Liquid

THR: A poison by intraperitoneal route. Moderately toxic by ingestion. Mildly toxic by inhalation. Very irritating to the skin, eyes and mucous membranes. A human systemic irritant by inhalation. Narcotic in high concentration. Dangerous fire hazard when exposed to heat, flame or oxidizers. Ignites on contact with potassium-tert-butoxide. Moderately explosive in the form of vapor when exposed to heat or flame. May form explosive peroxides upon exposure to air. Can react vigorously with reducing materials.

HEXONE HEGSOD

CAR

1880

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To fight fire, use alcohol foam, CO_2 , dry chemical. Incompatible with air; potassium-tert-butoxide. See also KE-TONES.

HFG550		HR: 3
HEXOPAL CAS: 6556-11-2		NIOSH: NM 7535400
mf: C ₄₂ H ₃₀ N ₆ O ₁₂	mw: 810.71	

PROP: Crystals. Mp: 254.3-254.9°. Practically insol in water; sol in dil acids.

SYNS:	
DILCIT	HEXANICIT
DILEXPAL	HEXANICOTINOYL INOSITOL
ESANTENE	HEXANICOTOL
INOSITOL HEXANICOTINATE	HEXA-3-PYRIDINECARBOX-
m-INOSITOL HEXANICOTINATE	YLATE-myo-INOSITOL (9CI)
meso-INOSITOL HEXANICOTINATE	LINODIL
myo-INOSITOL HEXANICOTINATE	MESONEX
INOSITOL NIACINATE	MESOTAL
INOSITOL NICOTINATE	PALOHEX
HAMOVANNID	
TOXICITY DATA:	CODEN:
scu-rat LD50:1180 mg/kg	NIIRDN 6,77,82

scu-rat LD50:1180 mg/kg ivn-rat LD50:268 mg/kg ipr-mus LD50:6400 mg/kg ivn-mus LD50:345 mg/mg CODEN: NIIRDN 6.77,82 NIIRDN 6.77,82 OYYAA2 7.149,73 NIIRDN 6.77,82

THR: Poison by intravenous route. Moderately toxic by subcutaneous route. When heated to decomposition it emits toxic fumes of NO_r .

HFG600 HR: 3 HEXOPRENALINE DIHYDROCHLORIDE

CAS: 4323-43-7 NIOSH: DO 6349000 mf: C₂₂H₃₂N₂O₆•2ClH mw: 493.48

SYNS:

N.N'-BIS(2-(3',4'-DIHYDROXYPHE-NYL)-2-HYDROXYETHYL)HEXA-METHYLENEDIAMINE DIHY-DROCHLORIDE

TOXICITY DATA:	CODEN:
ivn-rat TDLo:12 mg/kg (9-14D preg):TER	KSRNAM 6,983.72
orl-mus TDLo:12 mg/kg (7-12D preg):REP	KSRNAM 6.983,72
ivn-mus TDLo: 1200 µg/kg	KSRNAM 6,983,72
(7-12D preg): TER	
orl-rat LD50:10 g/kg	OYYAA2 26,811,83
ipr-rat LD50:139 mg/kg	KSRNAM 6,1286,72
scu-rat LD50:143 mg/kg	KSRNAM 6,1286,72
ivn-rat LD50:58 mg/kg	OYYAA2 26,811,83
orl-mus LD50:2036 mg/kg	KSRNAM 6,1286,72
ipr-mus LD50:133 mg/kg	KSRNAM 6,1286,72
scu-mus LD50:110 mg/kg	KSRNAM 6,1286,72
ivn-mus LD50:88 mg/kg	KSRNAM 6,1286,72

THR: Poison by subcutaneous, intravenous and intraperitoneal routes. Moderately toxic by ingestion. An experimental teratogen. Experimental reproductive effects. When heated to decomposition it emits toxic fumes of NO_x and HCl.

<i>HFG650</i> HEXOPRENALINE SULF	HR: 3	
CAS: 32266-10-7	NIOSH: DO 6350000	
mf: $C_{22}H_{32}N_2O_6 \cdot H_2O_4S$	mw: 518.64	
SYNS: N.N'-BIS(2-(3',4'-DIHYDROXYPHE- NYL)-2-HYDROXYETHYL)HEXA- METHYLENEDIAMINE SULFATE	ST-1512 SULFATE	
TOXICITY DATA:	CODEN:	
orl-rat TDLo:66 mg/kg (7-17D preg):TER	OYYAA2 27,239,84	
orl-mus TDLo:6 mg/kg (7-12D preg):TER	KSRNAM 6,983,72	
ipr-rat LD50:145 mg/kg	NIIRDN 6,745,82	
scu-rat LD50:150 mg/kg	NIIRDN 6,745,82	
ipr-mus LD50:159 mg/kg	NIIRDN 6,745,82	
scu-mus LD50:274 mg/kg	NIIRDN 6.745,82	
THR: Poison by subcutaneous and intraperitoneal routes.		

An experimental teratogen. Experimental reproductive effects. When heated to decomposition it emits toxic fumes of SO_x and NO_x . See also SULFATES.

HFH500 HR: 3 p-HEXOXYBENZOIC ACID-3-(2'-METHYLPIPERI-DINO)PROPYL ESTER CAS: 63916 83.6 NIOSH: DH 1450000

CAS: 63916-83-6		NIOSH: DH 1450000
mf: C ₂₂ H ₃₅ NO ₃	mw: 361.58	

TOXICITY DATA:	CODEN:
scu-mus LD50:222 mg/kg	RCPRAN 15,143,54
ivn-mus LD50:23 mg/kg	RCPRAN 15,143,54

THR: Poison by subcutaneous and intravenous routes. When heated to decomposition it emits toxic fumes of NO_x . See also ESTERS.

HF1500HR: 1HEXYL ACETATECAS: 142-92-7NIOSH: AI 0875000mf: $C_8H_{16}O_2$ mw: 144.24PROP: Colorless liquid. D: 0.878, mp: -60.9° , bp: 171.5° ,insol in water, very sol in alc and ether.SYNS:ACETIC ACID HEXYL ESTERHEXYL ALCOHOL. ACETATEHEXYL ACETATEHEXYL ACETATE

1-HEXYL ACETATE TOXICITY DATA:

orl-rat LD50:42 g/kg

CODEN:

TXAPA9 28,313,74

Reported in EPA TSCA Inventory.

THR: Mildly toxic by ingestion. When heated to decomposition it emits acrid smoke and fumes. See also ESTERS.

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METHANE DICHLORIDE MDR000

ivn-rbt LD50:36 mg/kg

UNEAQ 5.305.66 AECTCV 12,355,83 d-qal LD50:1 g/kg

EPA Genetic Toxicology Program.

"HR: Poison by intraperitoneal and intravenous routes." foderately toxic by ingestion and subcutaneous routes. An experimental teratogen. Experimental reproductive effects. Mutagenic data. A minor tranquilizer. When heated) decomposition it emits very toxic fumes of HCl and .10,.

1D0500

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METHAMPHETAMINE HYDROCHLORIDE CAS: 826-10-8 NIOSH: SH 5250000

HR: 3

mf: $C_{10}H_{15}N \cdot ClH$ mw: 185.72

PROP: Crystals; bitter taste. Mp: 170-175°. Sol in H₂O, alc and chloroform; almost insol in ether.

SYNS:	
DIPEX	METH
1-DESOXYEPHEDRINE HYDRO-	I-N-METHYL-B-PHENYLISOPRO-
CHLORIDE	PYLAMINE HYDROCHLORIDE
-)-N-a-DIMETHYLPHENETHYL-	SPEED
AMINE HYDROCHLORIDE	SYNDROX
TOXICITY DATA:	CODEN:
vn-mus TDLo:15 mg/kg (9-11D	TJADAB 4,131,71
preg): TER	
ivn-rbt TDLo:6 mg/kg (12-15D	TJADAB 4,131,71
preg): TER	
pr-rat LD50:25 mg/kg	27ZQAG - ,346,72
cu-rat LD50:30 mg/kg	27ZQAG -,346,72
ipr-mus LD50:70 mg/kg	JPETAB 89,382,47
scu-mus LD50:180 mg/kg	27ZQAG -,346,72
vn-mus LD50:33 mg/kg	27ZQAG -,346,72
orl-dog LD50:10 mg/kg	27ZQAG -,346,72
ivn-dog LD50:2700 µg/kg	PSEBAA 118.557,65
scu-cat LD50:50 mg/kg	27ZQAG -,346,72

THR: Poison by ingestion, intravenous, intraperitoneal, and subcutaneous routes. An experimental teratogen. Experimental reproductive effects. A powerful central nervous system stimulant. Caution: Excessive use may lead to tolerance and habituation. When heated to decomposition it emits very toxic fumes of HCl and NOr. See also BENZE-DRINE.

MDQ750	HR: 3
METHANE	
CAS: 74-82-8	NIOSH: PA 1490000
DOT: 1971/1972	
mf: CH ₄ mw: 16.05	

PROP: Colorless, odorless, tasteless gas. Mp: -182.6°. Bp: -161.5°, lel: 5.3%, uel: 15%, fp: -183.2°. D: 0.554 (a) $0^{\circ}/4^{\circ}$ (air = 1) or 0.7168 g/L, autoign temp: 650°, vap d: 0.6, flash p: -368.6°F. Sol in water, alc and ether.

SYNS: METHANE, REFRIGERATE LIQUID FTRE DAMP MARSH GAS (DOT) METHANE, COMPRESSED (DOT) METHYL HYDRIDE

Reported in EPA TSCA Inventory.

DOT Classification: Flammable Gas; Label: Flammable Gas

THR: A simple asphyxiant. Verv dangerous fire and explosion hazard when exposed to heat or flame. Reacts violently with powerful oxidizers (e.g., bromine pentafluoride; chlorine trifluoride: chlorine: fluorine: iodine heptafluoride; dioxygenyl tetrafluoroborate; dioxygen difluoride; trioxygen difluoride; liquid oxygen; ClO₂; NF₃; OF₂). Incompatible with halogens or interhalogens; air (forms explosive mixtures). Explosive in the form of vapor when exposed to heat or flame. To fight fire, stop flow of gas, CO₂ or dry chemical. See also ARGON for a description of asphyxiants.

HR: 3 **MDQ800 METHANE BORONIC ANHYDRIDE-PYRIDINE** COMPLEX

mw: 120.95 mf: CH₃BO•C₅H₅N

THR: Ignites spontaneously in air. When heated to decomposition it emits toxic fumes of NO_x. See also ANHY-DRIDES, PYRIDINE, and BORON COMPOUNDS.

<i>MDR000</i> METHANE DI	CHLORIDE	*	HR: 3
CAS: 75-09-2			NIOSH: PA 8050000
DOT: 1593			
mf: CH ₂ Cl ₂	mw: 84.93		

PROP: Colorless, volatile liquid. Bp: 39.8°, lel: 15.5% in O_2 , uel: 66.4% in O_2 , fp: -96.7°, d: 1.326-@ 20°/4°, autoign temp: 1139°F, vap press: 380 mm @ 22°, vap d: 2.93.

SYNS: AEROTHENE MM METHYLENE CHLORIDE (ACGIH. CHLORURE de METHYLENE DOT METHYLENE DICHLORIDE (FRENCH) DCM DICHLOROMETHANE (DOT) NCI-C50102 FREON 30 METHYLENE BICHLORIDE SOLMETHINE TOXICITY DATA: CODEN: skn-rbt 810 mg/24H SEV

eye-rbt 162 mg MOD eye-rbt 10 mg MLD eye-rbt 17500 mg/m³/10M dni-hmn: fbr 5000 ppm/1H-C cyt-ham: ovr 5 g/L dni-ham: lng 5000 ppm/1H-C sce-ham: Ing 5000 ppm/1H-C ihl-rat TCLo:4500 ppm/24H (1-17D preg): REP ihl-mus TCLo: 1250 ppm/7H (6-15D preg): REP ihl-rat TCLo: 3500 ppm/6H/2Y-I: CAR ihi-mus TCLo: 2000 ppm/5H/2Y-C:CAR

METYLENU CHLOREK (POLISH) RCRA WASTE NUMBER U080 JETOAS 9,171,76 JETOAS 9,171,76 TXCYAC 6,173,76 TXCYAC 6.173,76 MUREAV 81,203,81

MUREAV 116,361,83 MUREAV 81,203,81 MUREAV 81,203,81 TXAPA9 52,29,80

TXAPA9 32,84,75

FAATDF 4,30,84

NTPTR* NTP-TR-306.86

MDR250 METHANESULFONIC ACID

ihl-rat TCLo: 500 ppm/6H/2Y: ETA	TXAPA9 48.A185.79
orl-hmn LDLo:357 mg/kg: PNS.CNS	34ZIAG390.69
ihl-hmn TCLo: 500 ppm/1Y-1: CNS.CVS	ABHYAE 43.1123.68
ihl-hmn TCLo: 500 ppm/8H: CNS	SCIEAS 176.295.72
orl-rat LD50:2136 mg/kg	PPGDS* JAN81
ihl-rat LC50:88000 mg/m ³ /30M	FAVUAI 7.35,75
ihl-mus LC50: 14400 ppm/7H	NIHBAZ 191,1,49
ipr-mus LD50:1500 mg/kg	TXAPA9 9.139.66
scu-mus LD50:6460 mg/kg	TXAPA9 4.354,62
orl-dog LDLo: 3000 mg/kg	QJPPAL 7.205.34
ihl-dog LCLo: 14108 ppm/7H	NIHBAZ 191,1,49
ipr-dog LDLo:950 mg/kg	TXAPA9 10.119,67
scu-dog LDLo: 2700 mg/kg	QJPPAL 7.205,34
ivn-dog LDLo: 200 mg/kg	QJPPAL 7.205,34
ihl-cat LCLo: 43400 mg/m ³ /4.5H	AHBAAM 116,131,36
orl-rab LDLo: 1900 mg/kg	HBTXAC 1.94.56
ihl-rbt LCLo: 10000 ppm/7H	JIHTAB 26.8.44
scu-rbt LDLo: 2700 mg/kg	QJPPAL 7.205,34
ihl-gpg LCLo:5000 ppm/2H	FLCRAP 1.197,67

1ARC Cancer Review: Human Inadequate Evidence IMEMDT 41.43,86; Animal Inadequate Evidence IMEMDT 20,449,79; Animal Sufficient Evidence IMEMDT 41,43,86. NTP Carcinogenesis Studies (inhalation); Clear Evidence: mouse, rat NTPTR* NTP-TR-306,86. Reported in EPA TSCA Inventory. EPA Genetic Toxicology Program. Community Right To Know List.

OSHA PEL: TWA 500 ppm; CL 1000 ppm; Pk 2000/5M/ 2H

ACGIH TLV: TWA 50 ppm, Suspected Carcinogen

NIOSH REL: (To Methylene Chloride) TWA 75 ppm; Pk 500 ppm/15M

DOT Classification: Poison B; Label: St. Andrews Cross

THR: Poison by intravenous route. Moderately toxic by ingestion, subcutaneous and intraperitoneal routes. Mildly toxic by inhalation. An experimental carcinogen and tumorigen. Human systemic effects by ingestion and inhalation: paresthesia, somnolence, altered sleep time, convulsions, euphoria, and change in cardiac rate. An experimental teratogen. Experimental reproductive effects. An eye and severe skin irritant. Human mutagenic data. It is flammable in the range of 12-19% in air but ignition is difficult. It will not form explosive mixtures with air at ordinary temperatures. Mixtures in air with methanol vapor are flammable. It will form explosive mixtures with an atmosphere having a high oxygen content; in liquid O₂; N₂O₄; K; Na; NaK. Explosive in the form of vapor when exposed to heat or flame. Reacts violently with Li; NaK; potassiumtert-butoxide; (KOH + n-methyl-n-nitrosourea). It can be decomposed by contact with hot surfaces and open flame. and then yield toxic fumes which are irritating and give warning of their presence. When heated to decomposition it emits highly toxic fumes of phosgene and Cl⁻. See also CHLORINATED HYDROCARBONS, ALIPHATIC,

MDR250 METHANESULFONIC ACID

CAS: 75-75-2

mf: CH_4O_3S mw: 96.11

PROP: Solid. D: 1.4812 (a 18°/4°, mp: 20°, bp: 167° (a 10 mm. Sol in water, alc and ether. Corrosive to iron, steel, brass, copper and lead.

SYN: wsq 1

TOXICITY DATA: orl-rat LDLo: 200 mg/kg ipr-rat LDLo: 50 mg/kg orl-qal LD50: 1000 mg/kg CODEN: KODAK* 21MAY71 KODAK* 21MAY71 JRPFA4 48,371,76

Reported in EPA TSCA Inventory.

THR: Poison by ingestion and intraperitoneal routes. May be corrosive to skin, eyes and mucous membranes. Explosive reaction with ethyl vinyl ether. Incompatible with hydrogen fluoride. When heated to decomposition it emits toxic fumes of SO_x . See also SULFONATES.

MSF

MDR750

CAS: 558-25-8

METHANESULFONYL FLUORIDE

NIOSH: PB 2975000

HR: 3

HR: 3

mf: CH₃FO₂S mw: 98.10

SYNS: fumette methanesulphonyl fluoride

TOXICITY DATA: CODEN: orl-rat LD50:2 mg/kg LAEC** 17JUN74 ihl-rat LCLo: 140 mg/m³ 31ZOAD 1,287,68 ipr-rat LD50:3 mg/kg NATUAS 173,33,54 28ZEAL 4,271,69 scu-rat LD50:3500 µg/kg scu-mus LDLo: 3500 µg/kg 31ZOAD 1,287,68 ivn-mus LD50:1 mg/kg IAEC** 17JUN74 scu-dog LDLo: 3500 µg/kg 31ZOAD 1,287,68 IAEC** 17JUN74 ivn-dog LD50:5620 μ g/kg 31ZOAD 1,287,68 scu-rbt LDLo: 3500 µg/kg ivn-rbt LD50:3370 µg/kg **IAEC** 17JUN74**

EPA Extremely Hazardous Substances List. Reported in EPA TSCA Inventory.

THR: Poison by ingestion, inhalation, intraperitoneal, intravenous, and subcutaneous routes. When heated to decomposition it emits very toxic fumes of F^- and SO_x . See also FLUORIDES and SULFONATES.

MDR775 METHANETELLUROL

CAS: 25284-83-7 mf: CH₄Te mw: 143.64

THR: A poison. Ignites spontaneously in air. Explodes on contact with oxygen at room temperature. When heated to decomposition it emits toxic fumes of Te. See also TEL-LURIUM COMPOUNDS.

HR: 3

NIOSH: PB 1140000

31 /

scu-mus LD50:1250 μg/kg	JPETAB 67.153.39
iv mus LD50:475 µg/kg	JPETAB 79.127.43
ip :pg LD50:10900 µg/kg	AIPTAK 144,416.63
scu-gpg LD50:4800 µg/kg	AIPTAK 144.416.63
ivn-gpg LD50:390 µg/kg	AIPTAK 144.416.63
01)wd LD50:5600 µg/kg	TXAPA9 21.315.72

E. A Extremely Hazardous Substances List.

THR: Poison by ingestion, intraperitoneal, intravenous and su cutaneous routes. When heated to decomposition it emits very toxic fumes of SO_r and NO_r. See also STRYCHNINE and SULFATES.

SMP400		HR: 2
S 5 557		
C S: 65928-58-7		NIOSH: RC 8899500
mf: $C_{20}H_{25}NO_2$	mw: 311.46	

S N: 17-α-CYANOMETHYL-17-β-HYDROXY-ESTRA-4.9(10)-DIEN-3-ONE

TOXICITY DATA:	CODEN:
scu-rat TDLo:8 mg/kg (1D	ATSUDG 4.248,80
reg): REP	
o mky TDLo:600 μg/kg (3D	FESTAS 40.688,83
pre): REP	
orl-mus LD50:4000 mg/kg	PHARAT 34.319.79
ij mus LDLo: 1 g/kg	EXCEDS 81,175,83
s -mus LDLo: 5000 mg/kg	PHARAT 34,319,79
orl-rbt LDLo: 1 g/kg	EXCEDS 81.175,83
ipr-rbt LDLo: 1500 mg/kg	EXCEDS 81,175,83

C anide and its compounds are on the Community Right 10 Know List. EPA Genetic Toxicology Program.

T'IR: Moderately toxic by ingestion, subcutaneous and int peritoneal routes. Experimental reproductive effects. When heated to decomposition it emits toxic fumes of NO_r and CN⁻. See also NITRILES.

SMP500		H R : 3
• YPHNIC ACID • • • • • • • • • • • • • • • • • • •	•	NIOSH: VH 3540000
DOT: 0219 : : C ₆ H ₃ N ₃ O ₈	mw: 245.12	

AOP: Hexagonal, yellow crystals; astringent taste. Mp: (dry) 175.5°. Very sol in alc, ether.

: 'NS :	
2,+-DIHYDROXY-1.3,5-TRINITRO-	2.4.6-TRINITRO-1.3-BENZENEDIOL
BENZENE	2,4,6-TRINITRORESORCINOL
DIHYDROXY-2.4.6-TRINITRO-	TRINTTRORESORCINOL (DOT)
BENZENE	TRINITRORESORCINOL, DRY
IYDROXY-2.4.6-TRINITROPHE-	(DOT)
NOL	TRINITRORESORCINOL, wetted
6-TRINITROBENZENE-1, 3-DIOL	with less than 20% water (DOT)

ported in EPA TSCA Inventory.

DOT Classification: Class A Explosive; Label: Explosive A

THR: Very explosive. Upon decomposition it emits toxic fumes of NOr. See also NITRO COMPOUNDS OF ARO-MATIC HYDROCARBONS and EXPLOSIVES, HIGH.

SM0000 STYRENE CAS: 100-42-5 DOT: 2055 mf: C_8H_8

NIOSH: WL 3675000

mw: 104.16

$C_6H_5CH=CH_7$

PROP: Colorless, refractive, oily liquid. Mp: -31°, bp: 146°, lel: 1.1%, uel: 6.1%, flash p: 88°F, d: 0.9074 @ $20^{\circ}/4^{\circ}$, autoign temp: 914°F, vap d: 3.6, fp: -33°, ulc: 40-50. Very sltly sol in water; misc in alc, ether.

SYNS: CINNAMENE CINNAMENOL DIAREX HF 77 ETHENYLBENZENE NCI-C02200 PHENETHYLENE PHENYLETHENE PHENYLETHYLENE STIROLO (ITALIAN) STYREEN (DUTCH) STYREN (CZECH)

TOXICITY DATA:

skn-hmn 500 mg nse skn-rbt 500 mg open MLD skn-rbt 100% MOD eye-rbt 18 mg mma-sat 1 µmol/plate mmo-smc 1 mmol/L mrc-smc 1 mmol/L dns-hmn:lym 100 µmol/L dni-hmn:hla 28 mmol/L cyt-hmn:lym 300 ppm/72H sce-hmn:lym 500 µmol/L dnd-mus-ipr 10 mmol/kg sce-mus-ihl 46400 µg/kg/4D-l sce-mus-ihl 125 ppm/4D-I hma-mus/smc 1 g/kg orl-rat TDLo: 8600 mg/kg (1-22D preg/21D post): REP ihl-rat TCLo: 300 ppm/7H (6-15D preg): TER ihl-rat TCLo: 1500 µg/m³/24H (1-22D preg): TER ihl-hmn LCLo: 10000 ppm/ 30M ihl-hmn TCLo:600 ppm: NOSE, EYE ihl-hmn TCLo: 20 µg/m³: EYE orl-rat LD50:5000 mg/kg ihl-rat LC50:24 g/m³/4H ipr-rat LD50:1220 mg/kg orl-mus LD50:316 mg/kg

ihl-mus LC50:21600 mg/m³/2H ipr-mus LD50:660 mg/kg

STYRENE MONOMER (ACGIH) STYRENE MONOMER, inhibited DOD STYROL (GERMAN) STYROLE STYROLENE STYRON STYROPOR VINYLBENZEN (CZECH) VINYLBENZENE VINYLBENZOL

> CODEN: INMEAF 17,199,48 UCDS** 12/13/63 AMIHAB 14,387,56 AJOPAA 29,1363,46 MUREAV 56,147,77 **BSIBAC 59,233,83 BSIBAC 59,233,83** CRNGDP 3,681,82 MUREAV 93.447,82 MUREAV 58.277,78 ATSUDG 7.286.84 CALEDO 21.9.83 TXAPA9 55.37,80 APTOD9 19,A34,80 MUREAV 40,317,76 NTOTDY 7,23,85

TXCYAC 11,335.78

GISAAA 39(11),65,74

29ZWAE - .77,68

AMIHAB 14,387,56

GISAAA 26(8),11,61 AMIHAB 14,387,56 GTPZAB 26(8),53,82 GTPZAB 26(8),53,82 NCILB* NCI-E-C-72-3252.73 GTPZAB 26(8),53,82 ARZNAD 19.617.69

HR: 3

ivn-mus LD50:90 mg/kg ihl-gpg LCLo:12 mg/m³/14H

ARZNAD 19.617.69 JIHTAB 24.295.42

IARC Cancer Review: Animal Sufficient Evidence IMEMDT 19,231,79; Human Inadequate Evidence IMEMDT 19,231,79. NCI Carcinogenesis Bioassay (gavage); Inadequate Studies: mouse, rat NCITR* NCI-CG-TR-170,79; (gavage). Reported in EPA TSCA Inventory. EPA Genetic Toxicology Program. Community Right To Know List.

OSHA PEL: TWA 100 ppm; CL 200: Pk 600/5M/3H

ACGIH TLV: TWA 50 ppm; STEL: 100 ppm (skin); BEI: mandelic acid in urine at end of shift 1 gram/L, styrene in mixed-exhaled air prior to shift 40 ppb, styrene in mixed-exhaled air during shift 18 ppm, styrene in blood end of shift 0.55 mg/L, styrene in blood prior to shift 0.02 mg/L

DFG MAK: 100 ppm (420 mg/m³)

NIOSH REL: (Styrene) TWA 50 ppm; CL 100 ppm

DOT Classification: Flammable Liquid; Label: Flammable Liquid: Flammable or Combustible Liquid: Label: Flammable Liquid

THR: Experimental poison by ingestion, inhalation and intravenous routes. Moderately toxic experimentally by intraperitoneal route. Mildly toxic to humans by inhalation. A suspected human carcinogen. An experimental carcinogen and teratogen. Human systemic effects by inhalation: eye and olfactory changes. It can cause irritation and violent itching of the eyes @ 200 ppm, lacrimation, and severe human eye injuries. Its toxic effects are usually transient and result in irritation and possible narcosis. Experimental reproductive effects. Human mutagenic data. A human skin irritant. An experimental skin and eye irritant.

The monomer has been involved in several industrial explosions. It is a storage hazard above 32°C. A very dangerous fire hazard when exposed to flame, heat or oxidants. Explosive in the form of vapor when exposed to heat or flame. Reacts with oxygen above 40°C to form a heatsensitive explosive peroxide. Violent or explosive polymerization may be initiated by alkali metal-graphite composites; butyllithium; dibenzoyl peroxide; other initiators (e.g., azoisobutyronitrile; di-tert-butyl peroxide). Reacts violently with chlorosulfonic acid; oleum; sulfuric acid; chlorine + iron(III) chloride (above 50°C). May ignite when heated with air + polymerizing polystyrene. Can react vigorously with oxidizing materials. To fight fire, use foam, CO₂, dry chemical. When heated to decomposition it emits acrid smoke and irritating fumes. For further information, see Vol. 6, No. 2 of DPIM Report.

SMQ500	HR: 2
STYRENE POLYMER	
CAS: 9003-53-6	NIOSH: WL 6475000
$mf: (C_8H_8)_n$	
DOT: 2211	

SYNS: A 3-80 ABCOLENE. ATACTIC POLYSTYRENE BACTOLATEX BAKELITE SMD 3500 BASE III BEXTRENE XL 750 **BICOLASTIC A 75** BUSTREN CADCO 0115 CARINEX GP COPAL Z COSDEN 550 DENKA QP3 DIAREX 43G DORVON DOW 860 DYLENE DYLITE F 40 ESBRITE ESCOREZ 7404 ESTYRENE G 20 ETHENYLBENZENE HOMO-POLYMER FOSTER GRANT 834 GEDEX HI-STYROL HOSTYREN S HT-F 76 IT 40

KB (POLYMER) KRASTEN 1.4 LACQREN 550 LUSTREX MX 5517-02 NBS 706 OWISPOL GE PICCOLASTIC POLIGOSTYRENE POLYSTROL D POLYSTYRENE POLYSTYRENE BEADS (DOT) POLYSTYRENE LATEX POLYSTYROL PRINTEL'S **REXOLITE 1422** RHODOLNE SHELL 300 STYRAFOIL STYRAGEL STYRENE POLYMERS STYROFOAM STYROLUX STYRON TOPOREX 855-51 TROLITUI. UBATOL U 2001 VESTYRON VINYLBENZENE POLYMER VINYL PRODUCTS R 3612

TOXICITY DATA: imp-rat TDLo: 19 mg/kg: ETA ihl-mus LC50:120 mg/m³/10M CODEN: CNREA8 15.333.55 APFRAD 35,461,77

IARC Cancer Review: Animal Limited Evidence IMEMDT 19,231,79. Reported in EPA TSCA Inventory.

DOT Classification: Other Regulated Material; Label: None

THR: An experimental tumorigen by implant. When heated to decomposition it emits acrid smoke and irritating fumes. See also POLYMERS, INSOLUBLE.

SMR000 HR: 1 **STYRENE POLYMER with 1.3-BUTADIENE** CAS: 9003-55-8 NIOSH: WL 6478000

SYNS:	
AFCOLAC B 101	D ST 5 0
ANDREZ	DURANIT
BASE 661	EDISTIR RB 268
1,3-BUTADIENE-STYRENE CO-	ETHENYLBENZENE POLYMER
POLYMER	with 1,3-BUTADIENE
BUTADIENE-STYRENE POLYMER	GOODRITE 1800X73
1.3-BUTADIENE-STYRENE	HISTYRENE S 6F
POLYMER	HYCAR LX 407
BUTADIENE-STYRENE RESIN	K 55E
BUTAKON 85-71	KOPOLYMER BUTADIEN
DIAREX 600	STYRENOVY (CZECH)
DIENOL S	KRO I
DOW 209	LITEX CA
DOW LATEX 612	LYTRON 5202

TBO250 1.1.2.2-TETRACHLOROETHYLENE

PROP: Liquid. D: 1.588 (a 20/4°, bp: 129-130°. Sol in water; misc in alc, ether.

SYNS:

NCI-C52459 RCRA WASTE NUMBER U208 TOXICITY DATA: CODEN: skn-rbt 500 mg/24H AMPMAR 35.593.74 eye-rbt 100 mg SEV AMPMAR 35.593,74 orl-mus TDLo: 129 g/kg/2Y-I: NTPTR* NTP-TR-CAR 237.82 orl-mus TD: 258 g/kg/2Y-I:CAR NTPTR* NTP-TR-237.82

IARC Cancer Review: Animal Limited Evidence IMEMDT 41.87.86. NTP Carcinogenesis Bioassay (gavage); Clear Evidence: mouse NTPTR* NTP-TR-237.82; (gavage); No Evidence: rat NTPTR* NTP-TR-237,82. Reported in EPA TSCA Inventory.

THR: An experimental carcinogen. A skin and severe eye irritant. Incompatible with dinitrogen tetraoxide; 2,4-dinitrophenyl disulfide: potassium: potassium hydroxide; nitrogen tetroxide; sodium; sodium potassium alloy. When heated to decomposition it emits very toxic fumes of Cl⁻. For further information, see Vol. 4, No. 3 of DPIM Report.

TBO250 HR: 3 1.1.2.2-TETRACHLOROETHYLENE 💥

NIOSH: KX 3850000

DOT: 1897 mf: C_2Cl_4 mw: 165.82

CAS: 127-18-4

PROP: Colorless liquid; chloroform-like odor. Mp: -23.35°, bp: 121.20°, d: 1.6311 @ 15°/4°, vap press: 15.8 mm @ 22°, vap d: 5.83.

SYNS:

3110	
ANKILOSTIN	PERCHLOROETHYLENE (ACGIH.
ANTISOL 1	DOT
CARBON BICHLORIDE	PERCLENE
CARBON DICHLORIDE	PERCLOROETILENE (ITALIAN)
CZTEROCHLOROETYLEN (POL-	PERCOSOLVE
ISH)	PERK
DIDAKENE	PERKLONE
DOW-PER	PERSEC
ENT 1,860	RCRA WASTE NUMBER U210
ETHYLENE TETRACHLORIDE	TETLEN
FEDAL-UN	TETRACAP
NCI-C04580	TETRACHLOORETHEEN (DUTCH)
NEMA	TETRACHLORAETHEN (GERMAN)
PERAWIN	TETRACHLOROETHENE
PERCHLOORETHYLEEN, PER	TETRACHLOROETHYLENE (DOT)
(DUTCH)	TETRACLOROETENE (ITALIAN)
PERCHLOR	TETRALENO
PERCHLORAETHYLEN, PER (GER-	TETRALEX
MAN)	TETRAVEC
PERCHLORETHYLENE	TETROGUER
PERCHLORETHYLENE, PER	TETROPIL
(FRENCH)	

skn-rbt 810 mg/24H SEV	JETOAS 9,171,76
eye-rbt 162 mg MLD	JETOAS 9,171,76
mmo-sat 50 µL plate	NIOSH* 5AUG77
mma-sat 200 µL/plate	NIOSH* 5AUG77
dns-hmn:ing 100 mg/L	NTIS** PB82-185075
otr-rat: emb 97 µmol/L	ITCSAF 14.290,78
ihl-rat TCLo: 1000 ppm/24H	APTOD9 19,A21,80
(14D pre/1-22D preg): TER	
ihl-rat TCLo: 900 ppm/7H	TJADAB 19.41A.79
(7-13D preg): REP	
ihl-mus TCLo: 300 ppm/7H	TXAPA9 32,84,75
(6-15D preg): TER	
orl-mus TDLo: 195 g/kg/50W-I:	NCITR* NCI-CG-TR-
CAR	13,77
ori-mus TD :240 g/kg/62W-I:	NCITR* NCI-CG-TR-
CAR	13,77
ihl-hmn TCLo:96 ppm/7H:	NTIS** PB257-185
PNS,EYE,CNS	
ihl-man TCLo: 280 ppm/2H:	AMIHBC 5,566,52
EYE,CNS	
ihl-man TCLo:600 ppm/10M:	AMIHBC 5,566,52
EYE,CNS	
ihl-man LDLo:2857 mg/kg:	MLDCAS 5,152,72
CNS,PUL	
orl-rat LD50:8850 mg/kg	NPIRI* 1,96,74
ihl-rat LCLo: 4000 ppm/4H	JOCMA7 4,262,62
orl-mus LD50:8100 mg/kg	NTIS** PB257-185
ihl-mus LC50: 5200 ppm/4H	APTOA6 9,303,53
ipr-mus LD50:4700 mg/kg	NTIS** PB257-185
orl-dog LDLo: 4000 mg/kg	AJHYA2 9,430,29
ipr-dog LD50:2100 mg/kg	TXAPA9 10,119,67
ivn-dog LDLo:85 mg/kg	QJPPAL 7,205,34
orl-cat LDLo:4000 mg/kg	AJHYA2 9,430,29
ori-rbt LDLo: 5000 mg/kg	АЛНҮА2 9,430,29
scu-rbt LDLo: 2200 mg/kg	QJPPAL 7,205,34

TOXICITY DATA:

IARC Cancer Review: Animal Limited Evidence IMEMDT 20,491,79. NCI Carcinogenesis Bioassay (gavage); Clear Evidence: mouse NCITR* NCI-CG-TR-13,77; (inhalation); Clear Evidence: mouse. rat NTPTR* NTP-TR-311,86; (gavage); Inadequate Studies: rat NCITR* NCI-CG-TR-13,77. Reported in EPA TSCA Inventory. EPA Genetic Toxicology Program. Community Right To Know List.

OSHA PEL: TWA 100 ppm; CL 200 ppm; Pk 300ppm/ 5M/3H

ACGIH TLV: TWA 50 ppm (skin); STEL 200 ppm DFG MAK: 50 ppm (345 mg/m³); BAT: blood 100 µg/dl NIOSH REL: (Tetrachloroethylene) Minimize workplace exposure.

DOT Classification: Poison B: Label: St. Andrews Cross: ORM-A; Label: None

THR: Experimental poison by intravenous route. Moderately toxic to humans by inhalation with the following effects: local anesthetic, conjunctiva irritation, general anesthesia, hallucinations, distorted perceptions, coma and pulmonary changes. Moderately experimentally toxic by ingestion, inhalation, intraperitoneal and subcutaneous routes. An experimental carcinogen and teratogen. Experimental reproductive effects. Human mutagenic data. An

CODEN:

eye and severe skin irritant. The liquid can cause injuries to the eves; however, with proper precautions it can be handled safely. The symptoms of acute intoxication from this material are the result of its effects upon the nervous system. Can cause dermatitis, particularly after repeated or prolonged contact with the skin. Irritates the gastrointestinal tract upon ingestion. It may be handled in the presence or absence of air, water, and light with any of the common construction materials at temperatures up to 140°C. This material is extremely stable and resists hydrolvsis. A common air contaminant. Reacts violently under the proper conditions with Ba; Be; Li; N₂O₄; metals; NaOH. When heated to decomposition it emits highly toxic fumes of Cl⁻. Used in commercial dry cleaning and as a degreasing solvent. See also CHLORINATED HYDROCARBONS, ALIPHATIC. For further information, see Perchloroethylene, Vol. 1, No. 2 of DPIM Report.

TBO255

TETRACHLOROETHYLENE CARBONATE CAS: 22432-68-4

mf: $C_3Cl_4O_3$ mw: 225.84

THR: Reacts with tributylamine to form the toxic phosgene gas. When heated to decomposition it emits toxic fumes of Cl^{-} .

TBQ275 TETRACHLOROETHYLE	<i>HR: 3</i>
CAS: $16650-10-5$ mf: C ₂ Cl ₄ O mw: 181.82	NIOSH: KI 8760000
SYNS: epoxyperchlorovinyl pceo	TETRACHLOROEPOXYETHANE
TOXICITY DATA:	CODEN:
otr-ham:emb 4300 µmol/L	JJIND8 69,531,82
skn-mus TDLo: 300 mg/kg/66W- I:CAR	
 scu-mus TDLo:20 mg/kg/70W-I ETA 	: CNREA8 43.159.83
THR: An experimental carcin genic data. When heated to fumes of Cl^{-} .	
TBQ300	HR: 3
2,3,4,5-TETRACHLOROH	EXATRIENE
	NIOSH: MP 5425500
mf: $C_6H_4Cl_4$ mw: 217.90	
TOXICITY DATA:	CODEN:
orl-rat LD50:370 mg/kg	85GMAT108.82
ihl-rat LCLo:670 mg/m ³ /2H	85GMAT108,82
or mus I DS0 200 matka	85CMAT 108 82

TOXICITY DATA:	CODEN:
orl-rat LD50:370 mg/kg	85GMAT108.82
orl-rat LD50:370 mg/kg ihl-rat LCL0:670 mg/m ³ /2H	85GMAT108,82
orl-mus LD50:290 mg/kg ihl-mus LCLo:190 mg/m ³ /2H	85GMAT108,82
ihl-mus LCLo: 190 mg/m ³ /2H	85GMAT 108,82

THR: Poison by inhalation and ingestion. When heated to decomposition it emits toxic fumes of Cl⁻. See also CHLO-RINATED HYDROCARBONS, ALIPHATIC.

TBO500

CAS: 87-87-6

mw: 247.88 mf: $C_6H_2Cl_4O_2$

SYNS: USAF DO-62

TOXICITY DATA:	CODEN:
Jnd-omi 100 µmoi/L	MUREAV 145,71,85
dnd-mam: ivm 50 mmol/L	MUREAV 145.71.85
orl-mus LD50:500 mg/kg	ARTODN 40.63.78
ipr-mus LD50:25 mg/kg	NTIS** AD277-689

Reported in EPA TSCA Inventory.

THR: Poison by intraperitoneal route. Moderately toxic by ingestion. Mutagenic data. When heated to decomposition it emits toxic fumes of Cl⁻.

TBQ750

HR: 2

HR: 3

HR: 3

NIOSH: NT 2600000

TETRACHLOROISOPHTHALONITRILE

CAS: 1897-45-6 mf: $C_8Cl_4N_2$ mw: 265.90

SYNS:	
BRAVO	EXOTHERM TERMIL
BRAVO 6F	FORTURF
BRAVO-W-75	NCI-C00102
CHLOROALONIL	NOPCOCIDE
CHLOROTHALONIL	SWEEP
CHLORTHALONIL (GERMAN)	TCIN
DAC 2797	m-TCPN
DACONIL	TERMIL
DACONIL 2787 FLOWABLE FUN- GICIDE	2.4.5.6-TETRACHLORO-3-CYANO- BENZONITRILE
DACOSOIL	m-TETRACHLOROPHTHALONI-
1.3-DICYANOTETRACHLOROBEN-	TRILE -
ZENE	TPN (pesucide)
EXOTHERM	

TOXICITY DATA:	CODEN:
ori-rat TDLo: 142 g/kg/80W-C:	NCITR* NCI-CG-TR-
CAR	41.78
orl-rat LD50:10 mg/kg	85ARAE 4.75.76
orl-mus LD50:6 g/kg	INHEAO 4.11,66
ipr-mus LD50:2500 mg/kg	INHEAO 4,11,66

IARC Cancer Review: Animal Limited Evidence IMEMDT 30,319,83. NCI Carcinogenesis Bioassay (feed); Clear Evidence: rat NCITR* NCI-CG-TR-41,78. Cyanide and its compounds are on the Community Right To Know List. Reported in EPA TSCA Inventory. EPA Genetic Toxicology Program.

THR: Moderately toxic by intraperitoneal route. Mildly toxic by ingestion. An experimental carcinogen. When heated to decomposition it emits very toxic fumes of Cl⁻, NO_r, and CN⁻. Used as a fungicide. See also NITRILES.

TBR000

TETRACHLORONAPHTHALENE

CAS: 1335-88-2 NIOSH: QK 3700000 mf: $C_{10}H_{a}Cl_{a}$ mw: 265.94

HR: 3

TETRACHLOROHYDROQUINONE NIOSH: MX 7700000

TGK750 **TOLUENE** CAS: 108-88-3 mf: C-H₉ mw: 92.15 DOT: 1294

NIOSH: XS 5250000

HR: 3

PROP: Coloriess liquid; benzol-like odor. Mp: -95 to -94.5° , bp: 110.4°, flash p: 40°F (CC), ulc: 75-80, lel: 1.27%, uel: 7%, d: 0.866 (a 20°/4°, autoign temp: 996°F, vap press: 36.7 mm (\tilde{a} 30°, vap d: 3.14. Insol in water: sol in acetone: misc in absolute alc, ether, chloroform.

SYNS:

ANTISAL la
METHACIDE
METHYLBENZENE
METHYLBENZOL
NCI-C07272
PHENYLMETHANE
RCRA WASTE NUMBER U220

TOLUEN (CZECH) TOLUOL TOLUOL (DOT) TOLUOLO (ITALIAN) TOLU-SOL

CODEN:

TOLUEEN (DUTCH)

TOXICITY DATA:

eve-hmn 300 ppm skn-rbt 435 mg MLD skn-rbt 500 MOD eye-rbt 870 µg MLD eye-rbt 2 mg/24H SEV eye-rbt 100 mg/30S rns MLD oms-grh-ihl 562 mg/L dns-rat: lvr 30 umol/L cyt-rat-ihl 5400 µg/m³/16W-I cyt-rat-scu 12 g/kg/12D-I mnt-mus-orl 200 mg/kg mnt-mus-ipr 433 µg/kg/24H ihl-rat TCLo: 1500 mg/m3/24H (1-8D preg): TER orl-mus TDLo:9 g/kg (6-15D preg): TER ihl-hmn TCLo: 200 ppm: BRN.CNS.BLD ihl-man TCLo:100 ppm: CNS orl-rat LD50:5000 mg/kg ihl-rat LCLo: 4000 ppm/4H ipr-rat LDLo:800 mg/kg ivn-rat LD50: 1960 mg/kg unr-rat LD50:6900 mg/kg ihl-mus LC50:5320 ppm/8H ipr-mus LD50:1120 µg/kg unr-mus LD50:2000 mg/kg skn-rbt LD50:12124 mg/kg ihl-gpg LCLo: 1600 ppm scu-frg LDLo:920 mg/kg

JIHTAB 25.282.43 UCDS** 7/23/70 FCTOD7 20.563.82 UCDS** 7/23/70 28ZPAK -.23.72 FCTOD7 20.573.82 MUREAV 113.467,83 SinJF# 26OCT82 GTPZAB 25(7),33,81 GTPZAB 17(3),24,73 MUREAV 147.294,85 ARTODN 58,106.85 **TXCYAC** 11,55,78 **TJADAB 19,41A,79** JAMAAP 123.1106,43 WEHRBJ 9.131.72 AMIHAB 19,403,59 AIHAAP 30,470.69 TXAPA9 1,156,59 MELAAD 54,486,63 GISAAA 45(12),64,80 JIHTAB 25,366,43 AGGHAR 18,109.60 GISAAA 45(12),64,80

AIHAAP 30,470,69 JIDHAN 10.261,28

AEPPAE 130,250,28

Community Right To Know List. Reported in EPA TSCA Inventory. EPA Genetic Toxicology Program.

OSHA PEL: TWA 200 ppm; CL 300; Pk 500/10M

- ACGIH TLV: TWA 100 ppm; STEL 150 ppm; BEI: toluene in venous blood end of shift 1 mg/L
- DFG MAK: 100 ppm (375 mg/m³); BAT: blood end of shift 340 µg/dl

NIOSH REL: (Toluene) TWA 100 ppm; CL 200 ppm/10M

DOT Classification: Flammable Liquid: Label: Flammable Liquid

THR: Poison by intraperitoneal route. Moderately toxic by intravenous, subcutaneous and possibly other routes. Mildly toxic by inhalation. An experimental teratogen, Human systemic effects by inhalation: CNS recording changes, hallucinations or distorted perceptions, motor activity changes, antipsychotic, psychophysiological test changes and bone marrow changes. Experimental reproductive effects. Mutagenic data. A human eye irritant. An experimental skin and severe eye irritant.

Toluene is derived from coal tar, and commercial grades usually contain small amounts of benzene as an impurity. Inhalation of 200 ppm of toluene for 8 hours may cause impairment of coordination and reaction time; with higher concentrations (up to 800 ppm) these effects are increased and are observed in a shorter time. In the few cases of acute toluene poisoning reported, the effect has been that of a narcotic, the workman passing through a stage of intoxication into one of coma. Recovery following removal from exposure has been the rule. An occasional report of chronic poisoning describes an anemia and leucopenia, with biopsy showing a bone marrow hypoplasia. These effects, however, are less common in people working with toluene, and they are not as severe. At 200-500 ppm, headache, nausea, eye irritation, loss of appetite, a bad taste, lassitude, impairment of coordination and reaction time are reported, but are not usually accompanied by any laboratory or physical findings of significance. With higher concentrations, the above complaints are increased and in addition, anemia, leukopenia and enlarged liver may be found in rare cases. A common air contaminant.

A very dangerous fire hazard when exposed to heat, flame or oxidizers. Explosive in the form of vapor when exposed to heat or flame. Explosive reaction with 1,3-dichloro-5.5-dimethyl-2.4-imidazolididione: dinitrogen tetraoxide: concentrated nitric acid: $H_2SO_4 + HNO_3$: N_2O_4 ; AgClO₄; BrF₃; UF₆. Forms an explosive mixture with tetranitromethane. Can react vigorously with oxidizing materials. To fight fire, use foam, CO₂, dry chemical. When heated to decomposition it emits acrid smoke and irritating fumes. For further information, see Vol. 7, No. 5 of *DPIM Report*.

TGL500 TOLUENEDIAMINE

NIOSH: XS 9445000

CAS: 25376-45-8mf: $C_7H_{10}N_2$ mw: 122.19 DOT: 1709

SYNS:

ar-METHYLBENZENEDIAMINE DIAMINOTOLUENE METHYLPHENYLENEDIAMINE TOLYLENEDIAMINE

Community Right To Know List. Reported in EPA TSCA Inventory.

HR: 3

CODEN:

TIM000 HR: 1 1.1.2-TRICHLORO-2,2-DIFLUOROETHANE CAS: 354-21-2 NIOSH: KI 1435000 mf: CHCl₃F₂ mw: 157.37

SYNS:

EI-DIFLUORO-1.2.2-TRICHLORO-	UCON FLUOROCARBON 122
ETHANE	

TOXICITY DATA:	CODEN:
orl-rat LDLo:7500 mg/kg	HXPHAU 20(Pt 1),-
	459.66
ihl-rat LCLo: 4000 ppm/4H	UCMH** 15NOV62

THR: Mildly toxic by ingestion and inhalation. When heated to decomposition it emits very toxic fumes of F⁻ and Cl⁻. See also CHLORINATED HYDROCARBONS. ALIPHATIC and FLUORIDES.

TIM500	HR: 1
TRICHLORO ESTERTIN	

NIOSH: WH 8240000

SYN: ESTERTRICHLOROSTANNANE

TOXICITY DATA:	CODEN:
unr-rat LD50:5500 mg/kg	TIUSAD 107.1.76

OSHA PEL: TWA 0.1 mg(Sn)/m³ ACGIH TLV: TWA 0.1 mg(Sn)/m³ (skin)

NIOSH REL: (Organotin Compounds) TWA 0.1 mg(Sn)/m³

THR: When heated to decomposition it emits toxic fumes of Cl⁻. See also TIN COMPOUNDS and ESTERS.

<i>TIM750</i> 1,1,1-TRICHLC	ROFTHANE	*	HR: 3
CAS: 71-55-6		NIOSH: K	J 2975000
DOT: 2831 mf: C ₂ H ₃ Cl ₃	mw: 133.40		

PROP: Colorless liquid. Bp: 74.1°, fp: -32.5°, flash p: none, d: 1.3376 @ 20°/4°, vap press: 100 mm @ 20.0°. Insol in water; sol in acetone, benzene, carbon tetrachloride. methanol, ether.

SYNS:	
AEROTHENE TT	RCRA WASTE NUMBER U226
CHLOROETENE	SOLVENT 111
CHLOROETHENE	STROBANE
CHLOROTHANE NU	a-T
CHLOROTHENE	L.I.I-TCE
CHLOROTHENE (INHIBITED)	1.1.1-TRICHLOORETHAAN
CHLOROTHENE NU	(DUTCH)
CHLOROTHENE VG	1.1.1-TRICHLORAETHAN (GER-
CHLORTEN	MAN)
INHIBISOL	TRICHLORO-1.1.1-ETHANE
METHYLCHLOROFORM	(FRENCH)
METHYL CHLOROFORM (ACGIH,	a-TRICHLOROETHANE
DOTI	1.1.1-TRICLOROETANO (ITALIAN)
METHYLTRICHLOROMETHANE	TRI-ETHANE
NCI-C04626	

TOXICITY DATA:

eve-man 450 ppm/8H skn-rbt 5 g/12D-1 MLD skn-rbt 500 mg/24H MOD eve-rbt 100 mg MLD eve-rbt 2 mg/24H SEV dnr-esc 500 mg/L otr-mus: emb 20 mg/L orl-rat TDLo:43 mg/kg (1-22D preg/21D post): TER ihl-rat TCLo: 2100 ppm/24H (14D pre/1-20D preg): TER ihl-man LCLo: 27 g/m³/10M ihl-man TCLo: 350 ppm: CNS orl-hmn TDLo:670 mg/kg:GIT ihl-hmn TCLo: 920 ppm/70M: EYE.CNS ihl-man TCLo: 200 ppm/4H:CNS orl-rat LD50: 10300 mg/kg ihl-rat LC50:18000 ppm/4H ipr-rat LD50:5100 mg/kg orl-mus LD50:11240 mg/kg thl-mus LC50: 3911 ppm/2H ipr-mus LD50:4700 mg/kg orl-dog LD50:750 mg/kg FMCHA2 -. C242,83 ipr-dog LD50:3100 mg/kg TXAPA9 10.119.67 ivn-dog LDLo:95 mg/kg HBTXAC 5,72,59 ihl-cat LCLo: 600 mg/m³/4H 85GMAT -.38.82 orl-rbt LD50:5660 mg/kg AIHAAP 19,353,58 skn-rbt LDLo: 1 g/kg 85GMAT -.38,82 scu-rbt LDLo: 500 mg/kg HBTXAC 5.72,59 orl-gpg LD50:9470 mg/kg AIHAAP 19.353,58

BJIMAG 28.286.71 AIHAAP 19,353,58 28ZPAK -.28.72 AIHAAP 19,353,58 28ZPAK -.28,72 PMRSDJ 1,195,81 CALEDO 28.85.85 TJADAB 29(2),25A.84 TOXID9 1.28.81 JOCMA7 8.358.66 WEHSAL 10,82.73 NTIS** PB257-185 AIHAAP 19.353.58 ATSUDG 5,96,82 NTIS** PB257-185 28ZPAK -.28,72 NTIS** PB257-185 NTIS** PB257-185 SAIGBL 13.226,71 TXAPA9 13.287.68

IARC Cancer Review: Animal Inadequate Evidence IMEMDT 20.515,79. NCI Carcinogenesis Bioassay (gavage); Inadequate Studies: mouse, rat NCITR* NCI-CG-TR-3,77. Community Right To Know List.- Reported in EPA TSCA Inventory. EPA Genetic Toxicology Program.

OSHA PEL: TWA 350 ppm

ACGIH TLV: TWA 350 ppm; STEL 450 ppm DFG MAK: 200 ppm (2080 mg/m³); BAT: blood 55 µg/dl NIOSH REL: (1,1,1-Trichloroethane) CL 350 ppm/15M

DOT Classification: ORM-A; Label: None; Poison B; Label: St Andrews Cross

THR: Poison by intravenous route. Moderately toxic by ingestion, inhalation, skin contact, subcutaneous and intraperitoneal routes. An experimental teratogen. Human systemic effects by ingestion and inhalation: conjunctiva irritation, hallucinations or distorted perceptions, motor activity changes, irritability, aggression, hypermotility, diarrhea, nausea or vomiting and other gastrointestinal changes. Experimental reproductive effects. Mutagenic data. A human skin irritant. An experimental skin and severe eye irritant. Narcotic in high concentrations. Causes a proarrhythmic activity which sensitizes the heart to epinephrine-induced arrhythmias. This sometimes will cause cardiac arrest, particularly when this material is massively inhaled as in drug abuse for euphoria.

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TIN000 1,1,2-TRICHLOROETHANE

Under the proper conditions it can undergo hazardous reactions with aluminum oxide + heavy metals; dinitrogen tetraoxide; inhibitors; metals (e.g., magnesium; aluminum; potassium; potassium-sodium alloy); sodium hydroxide; $N_{2}O_{4}$; oxygen. When heated to decomposition it emits toxic fumes of Cl⁻. Used as a cleaning solvent, a chemical intermediate to produce vinylidene chloride, and as a propellant in aerosol cans. See also CHLORINATED HYDROCAR-BONS, ALIPHATIC. For further information see methyl chloroform, Vol. 2, No. 5 of DPIM Report.

TINOOO

TIN000		HR: 3
1,1,2-TRICHL	OROETHANE	
CAS: 79-00-5		NIOSH: KJ 3150000
mf: $C_2H_3CI_3$	mw: 133.40	

PROP: Liquid; pleasant odor. Bp: 114°, fp: -35°, d: 1.4416 $(\hat{a} \ 20^{\circ}/4^{\circ}, \text{ vap press: } 40 \text{ mm} \ (\hat{a} \ 35.2^{\circ}, \text{ mm})$

SYNS:

ETHANE TRICHLORIDE	B-TRICHLOROETHANE
NC1-C04579	1.2.2-TRICHLOROETHANE
RCRA WASTE NUMBER U227	TROJCHLOROETAN(1.1.2) (POL-
β-Τ	(SH)
1.1.2-TRICHLORETHANE	VINYL TRICHLORIDE

TOXICITY DATA:

skn-rbt 500 mg open MLD skn-rbt 810 mg/24H SEV eye-rbt 162 mg MLD skn-gpg 1440 mg/15M otr-mus:emb 25 mg/L cyt-gpg-skn 2880 µg/kg dnd-mam: ivm 1 mmol/L orl-mus TDLo:532 mg/kg (14D male): REP ori-mus TDLo:76 g/kg/78W-I: CAR orl-mus TD :152 g/kg/78W-I: CAR orl-rat LD50:580 mg/kg ihl-rat LCLo: 500 ppm/8H orl-mus LD50:378 mg/kg ipr-mus LD50:494 mg/kg scu-mus LD50:227 mg/kg orl-dog LDLo: 500 mg/kg ipr-dog LD50:450 mg/kg ivn-dog LDLo:95 mg/kg ihl-cat LCLo: 13100 mg/m³/4.5H skn-rbt LD50:3730 mg/kg scu-rbt LDLo: 500 mg/kg

APTOA6 41.298.77 CALEDQ 28.85.85 APTOA6 41,298,77 TODED5 11,243,82 DCTODJ 8.333.85 NCITR* NCI-CG-TR-74,78 NCITR* NCI-CG-TR-74.78 AIHAAP 30,470.69 AIHAAP 30,470,69 DCTODJ 8,333,85 TXAPA9 9,139,66 JPETAB 123,224.58 AJHYA2 16.325.32 TXAPA9 10,119,67 QJPPAL 7.205,34 AHBAAM 116,131,36 AIHAAP 30,470,69

QJPPAL 7,205,34

CODEN:

UCDS** 6/28/72

JETOAS 9,171.76

JETOAS 9,171.76

IARC Cancer Review: Animal Limited Evidence IMEMDT 20,533,79. NCI Carcinogenesis Bioassay (gavage); No Evidence: rat NCITR* NCI-CG-TR-74.78; (gavage); Clear Evidence: mouse NCITR* NCI-CG-TR-74.78. Community Right To Know List. Reported in EPA TSCA Inventory.

OSHA PEL: TWA 10 ppm (skin) ACGIH TLV: TWA 10 ppm (skin) DFG MAK: 10 ppm (55 mg/m³)

HR: 3

THR: Poison by ingestion, intravenous and subcutaneous routes. Moderately toxic by inhalation, skin contact, and intraperitoneal routes. An experimental carcinogen. Experimental reproductive effects. Mutagenic data. An eve and severe skin irritant. Has narcotic properties and acts as a local irritant to the eyes, nose and lungs. It may also be injurious to the liver and kidneys. Incompatible with potassium. When heated to decomposition it emits toxic fumes of Cl⁻. A priority pollutant associated with EPA superfund sites. See also CHLORINATED HYDROCARBONS, AL-IPHATIC and other trichloroethane entries. For further information, see Vol. 5, No. 3 of DPIM Report.

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TIN500 TRICHLOROETHANOL NIOSH: KM 3850000

CAS: 115-20-8 mf: C₂H₃Cl₃O mw: 149.40

PROP: Liquid, Mp: 17.8°, bp: 150° @ 765 mm, d: 1.54

(a 25/4°, vap press: 1 mm (a 20°, vap d: 5.16,

SYNS:

TRICHLORETHANOL 2.2.2-TRICHLOROETHANOL TRICHLOROETHYL ALCOHOL

TOXICITY DATA:

mmo-asn 5 µL/plate/2H sin-asn 10240 µmol/L sce-hmn:lym 178 g/L ori-rat LDLo:500 mg/kg ipr-rat LDLo: 300 mg/kg orl-mus LDLo: 500 mg/kg ivn-mus LD50:201 mg/kg ivn-rbt LDLo: 50 mg/kg rec-rbt LDLo: 500 mg/kg

CODEN: CBINA8 30,9,80 MUREAV 155,105,85 TOERD9 3.63.81 CRAA7 17,258,38 JPETAB 63.453.38 CRAAA7 17,258,38 28ZPAK -,78,72 JPETAB 63.453,38 CRAA7 17,258,38

2.2.2 TRICHLOROETHYL ALCO-

HOL

EPA Genetic Toxicology Program. Reported in EPA TSCA Inventory.

THR: Poison by intravenous and intraperitoneal routes. Moderately toxic by ingestion and rectal routes. Human mutagenic data. Explosive reaction with concentrated sodium hydroxide solutions. When heated to decomposition it emits toxic fumes of Cl⁻. Used as an hypnotic and anesthetic. See also CHLORINATED HYDROCARBONS, ALIPHATIC.

TIN750

TRICHLOROETHENYLSILANE CAS: 75-94-5 DOT: 1305

NIOSH: VV 6125000

mw: 161.49 mf: C₂H₃Cl₃Si

PROP: Fuming liquid. Bp: 90.6°, d: 1.265 @ 25/25°, flash p: 16°F.

SYNS:

SILANE, VINYL TRICHLORO 1-150 TRICHLORO(VINYL)SILANE TRICHLOROVINYL SILICANE UNION CARBIDE A-150

VINYLSILICON TRICHLORIDE VINYL TRICHLOROSILANE (DOT) VINYL TRICHLOROSILANE. IN-HIBITED (DOT)

HR: 3

ivn-rat LD50:24 mg/kg	YACHDS 12(Suppl 6),969.84
orl-mus LD50:325 mg/kg	YACHDS 12(Suppl 6),969.84
scu-mus LD50:284 mg/kg	YACHDS 12(Suppl 6),969,84
ivn-mus LD50:28 mg/kg	YACHDS 12(Suppi 6),969.84
orl-dog LD50:405 mg/kg	YACHDS 12(Suppl 6),969,84
ivn-dog LD50:9200 µg/kg	YACHDS 12(Suppl 6),969,84
orl-rbt LD50:425 mg/kg	YACHDS 12(Suppl 6),969.84

THR: Poison by ingestion, subcutaneous and intravenous routes. When heated to decomposition it emits toxic fumes of NO_x and HCl.

<i>XGA725</i> XILOBAM		HR: 3
CAS: $50528-97-7$ mf: $C_{14}H_{19}N_3O$	mw: 245.36	NIOSH: YT 8850000
SYNS:		

N-(2,6-DIMETHYLPHENYL)-N'-(1-	MCN-3113
METHYL-2-PYRROLIDINYLI-	I-(I-METHYL-2-PYRROLIDINYLI-
DENE)UREA	DENE-3-(2.6-XYLYL)UREA
TOXICITY DATA:	CODEN:
ord-rat I DS0:830 mg/kg	AIPTAK 233 326 78

 orl-rat
 LD50:830 mg/kg
 AIPTAK
 233,326,78

 ipr-rat
 LD50:128 mg/kg
 AIPTAK
 233,326,78

 orl-mus
 LD50:320 mg/kg
 JMCMAR
 21,1044,78

 ipr-mus
 LD50:110 mg/kg
 AIPTAK
 233,326,78

THR: Poison by ingestion and intraperitoneal routes. When heated to decomposition it emits toxic fumes of NO_x .

XGS000	HR: 2
XYLENE T	
CAS: 1330-20-7	NIOSH: ZE 2100000
DOT: 1307	
mf: C_8H_{10} mw	: 106.18

PROP: A clear liquid. Bp: 138.5° , flash p: $100^{\circ}F$ (TOC), d: $0.864 @ 20^{\circ}/4^{\circ}$, vap press: $6.72 \text{ mm} @ 21^{\circ}$. Composition: as nonaromatics 0.07%, toluene 14%, ethyl benzene 19.27%, p-xylene 7.84%, m-xylene 65.01%, o-xylene 7.63%, C9 and aromatics 0.04% (TXAPA9 33,543,75).

SYNS:	
DIMETHYLBENZENE	VIOLET 3
KSYLEN (POLISH)	XILOLI (ITALIAN)
METHYL TOLUENE	XYLENEN (DUTCH)
NCI-C55232	XYLOL (DOT)
RCRA WASTE NUMBER U239	XYLOLE (GERMAN)
TOXICITY DATA: eye-hmn 200 ppm skn-rbt 100% MOD skn-rbt 500 mg/24H MOD eye-rbt 87 mg MLD	CODEN: JIHTAB 25,282,43 AMIHAB 14,387,56 28ZPAK24,72 AMIHAB 14,387,56

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eye-rbt 5 mg/24H SEV	28ZPAK24.72
cyt-smc 1 mmol/tube	HEREAY 33,457,47
ihl-rat TCLo: 1000 mg/m ³ /24H	TXCYAC 11,55,78
(9-14D preg): TER	
orl-mus TDLo:31 mg/kg (6-15D	JTEHD6 9,97,82
preg):REP	
ihl-mus TCLo: 2000 ppm/6H	TJADAB 28.22A.83
(6-12D preg): TER	
ihl-hmn TCLo: 200 ppm:	JIHTAB 25,282,43
NOSE.EYE.PUL	
ihl-man LCLo: 10000 ppm/6H	BMJOAE 3.442.70
orl-rat LD50:4300 mg/kg	AMIHAB 14.387.56
ihl-rat LC50: 5000 ppm/4H	NPIRI* 1.123,74
scu-rat LD50:1700 mg/kg	NPIRI* 1,123,74
ipr-mus LD50:1548 mg/kg	AGGHAR 18,109,60
ipr-gpg LDLo: 2000 mg/kg	AIHAAP 35,21,74
ipr-mam LDLo: 2000 mg/kg	AJHYA2 7,276,27

Reported in EPA TSCA Inventory. EPA Genetic Toxicology Program. Community Right To Know List.

OSHA PEL: TWA 100 ppm

- ACGIH TLV: TWA 100 ppm; STEL 150 ppm; BEI: methyl hippuric acids in urine end of shift 1.5 g/g creatinine
- DFG MAK: (all isomers) 100 ppm (440 mg/m³); BAT: blood end of shift 150 µg/dl, urine 2 g/l
- NIOSH REL: (Xylene) TWA 100 ppm; CL 200 ppm/ 10M
- DOT Classification: Flammable Liquid; Label: Flammable Liquid; Flammable or Combustible Liquid; Label: Flammable Liquid

THR: Moderately toxic by intraperitoneal and subcutaneous routes. Mildly toxic by ingestion and inhalation. An experimental teratogen. Human systemic effects by inhalation: olfactory changes, conjunctiva irritation and pulmonary changes. Experimental reproductive effects. Mutagenic data. A human eye irritant. An experimental skin and severe eye irritant. Some temporary corneal effects are noted, as well as some conjunctival irritation by instillation (adding drops to the eyes one at a time). Irritation can start at 200 ppm. A very dangerous fire hazard when exposed to heat or flame; can react with oxidizing materials. To fight fire, use foam, CO_2 , dry chemical. When heated to decomposition it emits acrid smoke and irritating fumes. See also other xylene entries. For further information, see Vol. 6, No. 4 of DPIM Report.

XHA000		HR: 3
m-XYLENE		
CAS: 108-38-3		NIOSH: ZE 2275000
DOT: 1307		
mf: C ₈ H ₁₀	mw: 106.18	

PROP: Colorless liquid. Mp: -47.9° , bp: 139°, lel: 1.1%, uel: 7.0%, flash p: 77°F, d: 0.864 @ 20/4°, vap press: 10 mm @ 28.3°, vap d: 3.66, autoign temp: 986°F. Insol in water; misc with alc, ether and some organic solvents.

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Chemical name, structure/formula, CAS and RTECS Nos	Synonyms, trade names, and conversion	Exposure limite (TWA	IDLH	Physical description	Chemical an prope		incompatibilities and reactivities	Measuremen method (See Table 1
and DOT ID and guide Nos.	factora	uniess noted otherwise)			MW, BP, SOL FI.P, IP, Sp.Gr, flammability	VP, FAZ UEL, LEL		
Acetaldehyde	Acetic aldehyde, Ethanal.	NIÓSH Ca	Ca [10.000	Coloriess liquid or gas (above	MW 44 1 BP 69 F	VP 740 mm FRZ 190°F	Strong oxidizers, acids, bases.	XAD 2*. Toluene
сн,сно	Ethyl aldehyde			69°F) with a pungent, fruity	Sol Miscible FLP 36 F	UEL 60%	acios, bases. alcohols, ammonia & amines, phenols,	GC/FID
75 07 0 AB1925000		OSHA 100 ppm (180 mg/m ¹) ST 150 ppm		odor	IP 10 22 eV		ketones, HCN, H,S [Note: Prolonged cont may cause formatio that may explode ar	(#2538) act with air h of peroxides
1089-26	1 ppm = 1 83 mg/m f	(270 mg/m ³)			Sµ Gr. 0.79 Class IA Flamma	ble Liquid	tainers; easily under polymerization]	
Acelic acid	Acetic acid (aqueous). Ethanoic acid.	NIOSH 10 ppm	1000 ppm	Coloriess liquid or crystals with	MW 60 1 BP. 244 F	VP. 11 mm FRZ 62 F	Strong exidizers (especially chromic	Char, HCOOH,
сн,соон	Glacial acetic acid (pure compound).	(25 mg/m ³) ST 15 ppm		a sour, viñegar like odor	Sol Miscible	UEL (200°F) 19 9%	acid, sodium peroxide & nitric acid).	
64 19 7 AF1225000	Methanecarboxiyic acid	(37 mg/m³)	ing/m³) (Note Pure c pound is a	Note Pure.com pound is a solid	IP 10 66 eV	LEL 40%	strong caustics [Note: Corrosive to	[#1603]
1842 29 (soin) 2790 60 (10 80% acid) 2789 29 (80% acid)	[Note: Can be found in concentrations of 5-8% in vinegar] 1 ppm = 2 50 mg/m ³	OSHA 10 ppm (25 mg/m³)		below 62°F Often used in an aqueous solution]	Sp.Gr. 1.05 Class II Combust	bie Liquid	` metals.)	
Acetic anhydride	Acetic acid anhydride,	NIOSH/OSHA C 5 ppm	1000 ppm	Colorless liquid	MW: 102 1 BP: 282°F	VP 4 mm FRZ 99°F	Water, alcohols,	Bub,
(CH,CO),O	Acetic oxide, Acetyl oxide, Ethunoic anhydride	(20 mg/m³)		with a strong, pungent, vinegar- like odor	Sol 12% FLP 120 F	UEL 10 3%.	strong oxidizers (especially chromic acid), amines	Reagent. Vis; III
108-24-7 AK1925000					IP 10 00 eV	IEL 27/6	strong caustics [Note: Corrosive to iron: steel & other	[#3506]
1715-39	1 ppm <mark>≍ 4 24 mg/m</mark> ⊁				Sp Gr 1 08 Class II Combusti	ble Liquid	metals. Reacts with water to form acetic acid]	
Acetone	Dimethyl ketone,	NIOSH 250 ppm	20.000 ppm	Colorless liquid with a fragrant,	MW 58 1 BP: 133 F	VP 180 mm	Oxidizers, acids	Char, CS ₂
сн,сосн,		(590 mg/m ³)	ppm	mint-like odor	Sol Miscible	FRZ. 140 F UEL 13% LEL 25%		GC/FID.
67 64 1 AL 3150000		OSHA 750 ppm (1800 mg/m²) ST 1000 ppm	TWA for "d acetate fi stayed or	rcement of the OSHA offers" in the cellulose ber industry was 1 9/5/89 until 9/1/90,		LEL 23%		(#1300. Ketones
1090 26	1 ppm = 2 42 mg/m ³	(2400 mg/m ³)		e OSHA STEL does	Sp Gr 0 79 Class IB Flammat	ale Liquid		

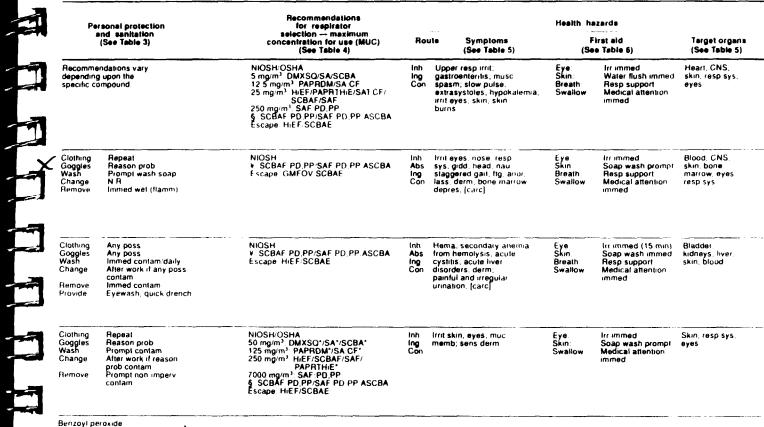
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	rsonal protection ind_sanitation	Recommendations for respirator selection — maximum			-	Health h	ezarde .	·
	(See Table 3)	concentration for use (MUC) (See Table 4)	Ro	ule	Symptoms (See Table 5)		irst sid e Table 6)	Target organ (See Table 5)
Clothing Goggles Wash Change Remove Provide	Repeat Any poss Prompt wet N R Immed wet (flamm) Eyewash	NIOSH ¥ SCBAF PD.PP/SAF PD.PP ASCBA Escape GMFOV/SCBAE	inh Ing	conj. eye, s	nose, throat writ; cough; CNS depres, ikin burns, derm, ed pulm edema, [carc]	Eye Skin Breath Swallow	Irr immed Water flush prompt Resp support Medical attention immed	Resp sys, skin. kidneys
Clothing Goggles Wash Change Remove Provide	Any poss >50%/Repeat 10 49% Any poss Immed contrum >50% Prompt 10 49% N R Immed non imperv contam >50% Prompt non imperv contam 10 49% Evewash (>5%/Ouck drench (>50%	NIOSH-OSHA 250 ppm SA CF1:PAPROV1 500 ppm CCRFOV-SCBAF:SAF GMFOV/PAPR1OV1 1000 ppm SAF-PD.PP § SCBAF PD.PP/SAF PD.PP ASCBA Escape GMFOV/SCBAE	Inh	throal bron; skin s	lac, irrit nose , phar edema, chronic burns eyes, skin, ens, dental erosion, skin, hyperkeratosis	Eye Skin Breath Swallow	fri immed Water flush immed Resp support Medical attention immed	Resp sys, skin eyes, teeth
Clothing Goggles Wash Change Remove	Reason prob Any poss Immed contam N R Immed non imperv contam Eyewash, quick drench	NIOSH/OSHA 125 ppm SA CF ¹ /PAPROV ⁴ 250 ppm CCRFOV/SCBAF/SAF/ GMFOV/PAPRTOV ⁴ 1000 ppm SAF PD.PP § SCBAF PD.PP/SAF PD.PP ASCBA Escape GMFOV/SCBAE	inh ing Con	opac, irrit; co	lac, corneal edema, photo, nasal, phar Jugh, dysp. bron, urns, vesic, sens	Eye Skin Breath Swallow	Irr immed Water flush immed Resp support Medical attention immed	Resp sys, eyes, skin
Clothing Goggles Wash Change Remove	Repeat Reason prob Prompt wet N R Immed wet (flamm)	NIOSH 1000 ppm CCROV*/PAPROV*/SA*/ SCBA* 6250 ppm SA CF* 12,500 ppm: SAF PD.PD 63, SCBAF PD.PP/SAF PD.PP 63, SCBAF PD.PP/SAF PD.PP ASCBA Escape GMFOV/SCBAE	inh Ing Con		es, nose, throat, dizz, derm	Eye Skin Breath Swallow	Irr immed Soap wash immed Resp support Medical attention immed	Rəsp sys, skin

Chemical name, structure/formula, CAS and RTECS Nos.,	Synonyme, trade names, and conversion	Exposure limits {TWA	IDLH	Physical description	Chemical an prope		incompatibilities and reactivities	Measuremen method (See Table 1				
and DOT ID and guide Nos.	lactora	uniess noted otherwise)			MW, BP, SOL FI.P, IP, Sp.Gr, Hammability	VP FRZ UEL, LEL						
Barium (soluble compounds as Ba) 1 Ba(NO ₃), 2 BaCi, 1 10022 31 8 CO9625000 2 10361 37 2 CO8750000 1446 42 (Barium nitrate)	t Barium nitrate, Barium dinitrate 2 Barium chloride, Barium dichloride Synonyms of other soluble compounds vary depend- ing upon the specific com pound	NiOSH OSHA 0 5 mg/m?	1100 mg/m³	Barrum nitrate & Barrum chloride are white odoriess solids	MW 2614 2083 BP Decompuses 72840 F Sot. 9/38% FI-P NA? IP ?? Sp Gr 3 24 3 86 Ba(NO), Nanca BaCI, Combustit	UEL NA? LEL NA? mbushble Solid	Acids, axidizers (Note: Contact of barium nutrate with combustible interna may cause fire.)	Filter, Water, AA, I NI {#7056}				
Benzene	Benzol, Phenyl hydride	NIOSH Ca	Ca (3000 ppm)	Coloriess to light yellow	MW 78 1 BP: 176 F	VP. 75 mm FRZ: 42 F	Strong oxidizers, many fluorides &	Char, CS ₂ ;				
C ₆ H ₆		See Appendix A 0 1 ppm			See Appendix A			liquid with an aromatic odor	Sol 0 07% FLP 12 F	UEL 79% LEL 13%	perchiorates	ĞČ/HD. III
71 43 2 CY1400000		ST i ppm OSHA		[Note: A solid below 42"F]	IP 924 eV			(#1500. Hydra				
		(1910.1028) 1 ppm	ACGIH A2				Note Measurements					
1114 27	1 µpm × 3 25 mg/m ¹	St'5 ppm			Sp Gr - 0 88 Class 18 Flammat	ble Liquid	made with a portable NIOSH #3700 (III)]	a GC using				
Benzidine	4.4 Bianiline 4.4 Biphenyldiamine;	NIOSH Ca	Ca	Gravish yellow, reddish gray, or	MW: 184 3 BP: 752*F	VP Low MLT 239 F	None reported	Filter Sigel				
NH,C,H,C,H,NH,	1.1 Siphenyl-4,4 diamine. 4.4 Diaminobiphenyl.	See Appendix A		white crystalline	Sol(54°F) 0 04%	UEL ?		Reagent HPLC/UVD				
92 87 5 DC9625000	p Diaminodiphenyl	OSHA [1910-1010] See Appendix B	10]	p 0	FIP?			111 111 [#5509]				
1865 53		ACGIH A1 [skin]			Sp Gr -1 25 Combustible Solid	, but difficult to b	nu fi					
Benzoyl peroxide	Benzoperoxide,	NIOSH/OSHA	7000	Coloriess to white	MW: 242.2 BP: ?	VP <1 mm	Combustible	Filter,				
(C ₆ H ₆ CO) ₂ O ₇	Dibenzoyl peroxide	5 mg/m ³	mg /m²	crystals or a granular powder with a faint	Sol. < 1% FLP. ?	MLT: 217 F UEL ? LEL ?	substances (wood, paper, etc.), acids, alkalis, alconols,	Diethyl ether, HPLC/UVD,				
94 36 0 DM8575000				benzaldehyde like odor			amines, ethers [Note: Containers may explode when heated	1 [#500a] [II				
2085/2086/2087 49 2088/2089/2090 49					Sp Gr(77 F): 1 33	lazedu jandi z -	Extremely explosion- to shock, heat, and fi nd burns very rapidly)					

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Benzoyl peroxide

Chemical name, structure/formula, CAS and RTECS Nos.,	Synonyms, trade names, and conversion	Exposure limits (TWA	IDLH	Physical description	Chemical an prope		Incompatibilities and reactivities	Measurement method (See Table 1)
and DOT ID and guide Nos.	factors	unless noted otherwise)			MW, BP, SOL FI.P. IP, Sp.Gr. Hammability	VP, FRZ UEL, LEL		
Вголипе	Molecular bromine	NIOSH OSHA 0 1 ppm	10 ppm	Dark reddish brown fuming liquid with	MW 159.8 BP: 139"F	VP 172 mm FRZ 19 F	Combustible organics (sawdust, wood,	Bub, none
Br _. .		(0.7 mg m.º) SI0 ∋ppm		suffocating. irritating fumes	Sol 4% FLP NA	UEL NA	collon, straw, etc.). aluminum: readily	IC OSHA
7726-95-6 EF9100000		(2 mg m ⁴)		-	IP 10 55 eV		oxidizable materials ammonia, hydrogen,	[#11)108]
1744 59					Sp.Gr. 3.12 Noncombustible Eigond but accelerates the		acetylene phosphorus potassium sodium [Note Corrodes iron, s	Sleet
	1 ppm + 6 64 mg.m ³	NIOSHOSHA			burning of cambu MW 252 8	VP 5 mm	stamless steel & cop	· · · · · · · · · · · · · · · · ·
Bromotorm CHBr	Methyl tribromide. Tribromomethane	0 5 ppm (5 mg m ³)	Unknown	Coloriess to yellow liquid with a chloroform like	BP 301°F Sol 0 1%	FRZ 47 F	Lithium, sodium. potassium, calcium.	Char. CS ₂ . GC/F1D.
75 25 2		(5 mg m3) [skin]		odor Note A solid	FIP NA IP 10 48 eV	LEL NA	aluminum, zinc, magnesium, strong	(II) (II) (II) (II) (II) (II) (II) (II)
PB560000				below 47 F)	IF 1048 EV		caustics, acetone [Note: Gradually	Haloge
					Sp Gr 2 89		decomposes, acquir ing yellow color, air & light accel-	Hydro carbons]
2515 59	1 ppm = 10 51 mg/m ³			. <u> </u>	Noncombustible L		erate decomposition	J
1,3 Butadiene	Biethylene Bivinyl	NIOSH Ca	Ca [20.000	Coloriess gas with a mild aromatic or gasoline-like odor [Note: A liquid below 24:5]	MW 54 1 BP. 24-F	VP ⇒1 atm FRZ: 164 F	Phenol, chlorine dioxide: copper	Char(2). CH ₂ Cl ₂ ,
СНСНСН-СН,	Butadiene Divinyl,	See Appendix A Reduce exposure			Sol Insoluble UEL 12.0% FLP NA (Gas) LEL 2.0% <0°F (Liquid) IP 9.07 eV		crotonaldehyde Note: May contain	GC/FID. III
106 99 0 E19275000	Erythrene. Vinylethylene	to lowest teasible concentration						[#1024]
1010-17 (inhibited)	1.ppm = 2.25 mg/m³	OSHA 1000 ppm (2200 mg.ml)	ACGIH A2, 10 ppm (22 mg/m ³)		Sp Gr 0.65(Liquid Flammable Gas Class IA Flammat	•	polymerization May form explosive j upon exposure to air	
2 Butanone	Ethyl methyl ketone,	NIOSH OSHA	3000 ppm	Coloriess liquid	MW: 72 1	VP 71 mm	Strong oxidizers	Ambersorb.
сн,сосн,сн,	MEK. Methyl acetons,	200 ppm (590 mg/m³)		with a moderately sharp, fragrant,	BP: 175 F Sol. 28%	FRZ 123 F UEL(200°F)	amines, ammonia, inorganic acids,	CS, GC/HD,
78 93 3 EL6475000	Methyl ethyl ketone	ŠT 300 [°] ppm (885 mg/m³)		mint- or acetone- like odor	FIP 16°F IP 954 eV	11 4% I FL (200 F) 1 4%	caustics, copper. isocyanates, pyridines	111 [#2500]
1193-12 32 -26	1 ppm = 3 00 mg/m ³				Sp Gr 0 81 Class IB Flammab	le Liquid		

•

Repeat Reason prob N R N R Prompt non impervicontam Eyewaish

Clothing

Goggles Wash Change Remove

Provide

2 Butanone

Irr immed Water wash immed Fresh air Medical attention immed

Eye Skin

Breath Swallow CNS lungs

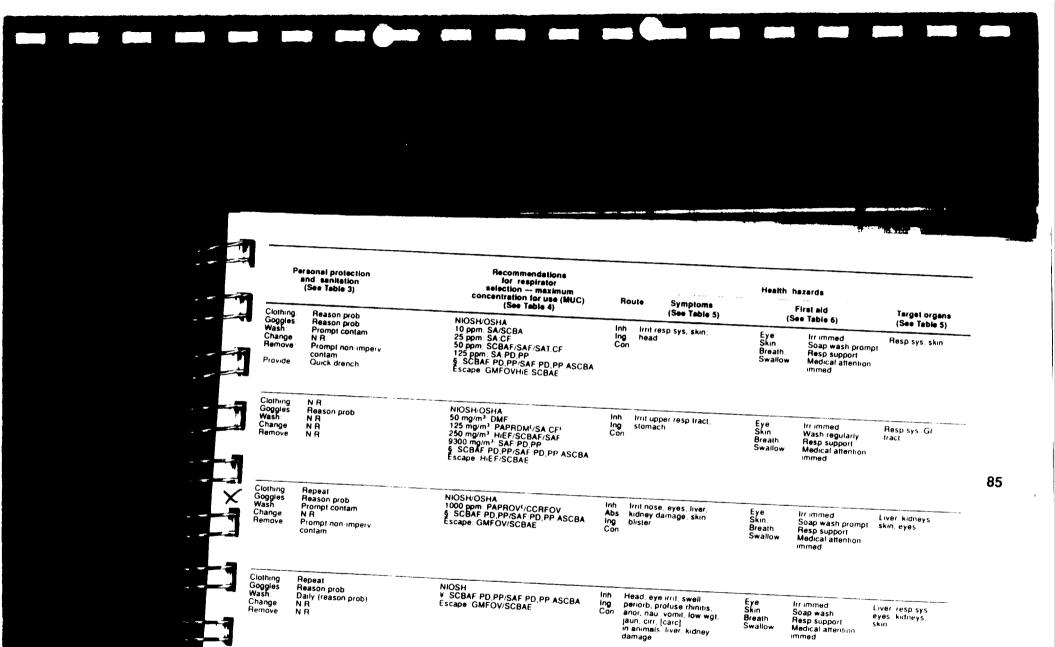
	sonal protection nd sanifation (See Table 3)	Recommendations for respirator selection maximum concentration for use (MUC) (See Table 4)	Rou	ite Symptoms (See Table 5)		izarda irat aid a Table 6)	Target organs (See Table 5)
Clothing Goggles Wash Change Remove Provide	Any poss Any poss Jimmed contam N R Immed contam Eyewash, quick dreamh	NIOSH:OSHA 2 5 ppm SA CF1:PAPRS: 5 ppm CCRFS: GMFS:/SCBAF SAF PAPRTS: 10 ppm SAF PD,PP \$ SCBAF PD,PP/SAF PD,PP ASCBA Escape GMFS:SCBAE	Inh Ing Con	Dizz, head fac, epis, cough, feeling of oppression pulm edema, pneu, abdom pain, diarr; measle like eruptions; severe burns eyes, skin	Eye Skin Breath Swallow	Irr immed Soap wash immed Resp support Medical attention immed	Resp sys eyes CNS
Clothing Goggles Wash Charige Remove	Repeat Reason prob Prompt contam N R Prompt iron implify contam	NIOSH OSHA 12 5 ppin SA CF ¹ PAPHOV ¹ 25 ppin CCRFOV GMFOV SCBAF SAF PAPRIOV ¹ 1000 ppin SAF PD PP § SCBAF PD PP SAF PD PP ASCBA Escape GMFOV SCBAE	inh Abs ing	Irnt eyes msp sys. CNS depr⊷sion, liver damage	Eye Skin Breath Swallow	Irr immed Soap wash prompt Resp support Medical attention immed	Skin, liver, kidneys, resp sys, CNS
Clothing Goggles Wash Change Remove	Prevent skin literang Reason prob Immed wet N R Immed wet (flamm)	NIOSH V SCBAF PD.PP/SAF PD.PP.ASCBA Escape: GMFS SCBAE	loh Con	Init eyes nose throat drow is head frostbite (carc)	Ey u Skin Breath	Irrammed Water flush immed Resp support	Eyes respisys CNS

linit eyes inose, head. dizz, vonut

NIOSH/OSHA Inh 1000 ppm PAPROV^E CCRFOV Ing 3000 ppm GMFOV/SA CF^T/SCBAF/SAF Con § SCBAF PD, PP/SAF PD, PP ASCBA Escape GMFOV/SCBAE

Chemical name, structure/formule, CAS and RTECS Nos.,	Synonyms, trade nemes, and conversion	Exposure limits (TWA	IDLH	Physical description	Chemical an prope		incompatibilities and reactivities	Measurement method (See Table 1)
and DOT ID and guide Nos.	factors	uniess noted otherwise)			MW, BP, SOL FI.P, IP, Sp.Gr, flammability	VP, FRZ UEL, LEL		
Dibutyl phosphate	Dibutyl acid o phosphate, di-n-Butyl hydrogen	NIOSH/OSHA	125 ppm	Pale amber, odorless liquid	MW 2102 BP 212"F	VP 1 mm (approx)	Strong oxidizers	Eiller, CH ₁ CN,
(C₄H ₄ O)₂(OH)PO	phosphate, Dibutyl phosphoric acid	(5 mg/m ³) ST 2 ppm			(Decomposes) Sol Insoluble	FRZ. ? UEL ?		GC/FPD III
107 66 4 189605000		(10 mg/m ²)			FIP? IP?	LEL ?		[#5017]
	1 ppm = 8 74 mg/m ³				Sp.Gr. 1.06 Combustible Liqui	d		
Dibutyiphthalate	DBP; Dibutyl 1,2 benzene	NIOSH/OSHA 5 mg/m ³	9300 mg/m3	Colorless to faint yellow, oily	MW 278 3 BP 644°F	VP <0.01 mm FR2 31 F	Nitrates, strong oxidizers, alkalis	Filter, CS,
C ₆ H ₄ (COOC ₄ H ₉) ₂	dicarboxylate; Di n butyl phthalate		.u9.un-	liquid with a slight, aromatic	Sol(77°F) 0 5%	UEL ? LEL(456 F)	& acids, liquid chlorine	GC/FID III
84 74 2 Tio875000					FLP: 315 F IP: 2	0.5%		[#5020]
	1 ppm = 11 57 mg/m³				Sp Gr 1 05 Class IIIB Combus	Stible Liquid		
Dichlorobenzene	o DCB; 1,2 Dichlorobenzene;			Coloriess to pale yellow liquid with	MW: 147 0 BP: 357"F	VP 1 mm FRZ 1 F	Strong oxidizers, aluminum, acids,	Char, CS,
°°H'CI'	ortho-Dichlorobenzene, o Dichlorobenzol	(300 mg/m ³)		a pleasant, aromatic odor	Sol 0.01% FLP 151°F	0.01% UEL 9.2% P.151°F LEL 2.2%	acid lumes, chlorides	
15 50 1 224500000					IP 9 06 eV			[#1003, Haloge nated
591 58	1 ppm = 6 11 mg/m³				Sp Gr 1 30 Class IIIA Combus	dible Eiquid		Hydro carbons)
Dichlorobenzene	p DCB; 1.4 Dichlorobenzene,		Ca (1000 ppm)	Colorless or white crystalline solid	MW: 147 0 BP: 345°F	VP(77°F) 0.4 mm	Oxidizers	Char, CS),
"H"CI,	para Dichlorobenzene; Dichlorocide	See Appendix A	[oo ppn]	with a mothball- like odor.	Sol: 0.008% FLP 150°F	MLT: 128 F UEL ?		GC/FID
06 46 7 Z4550000		OSHA 75 ppm (450 mg·m·)		(insecticide)	1P 8 98 eV	LEL		(#1003, Haloge nated
592 58		ST 110 ppm (675 mg/m³)	ST 110 ppm		Sp Gr 1 25 Combustible Solid. some effort to ignit			Hydro carbons)

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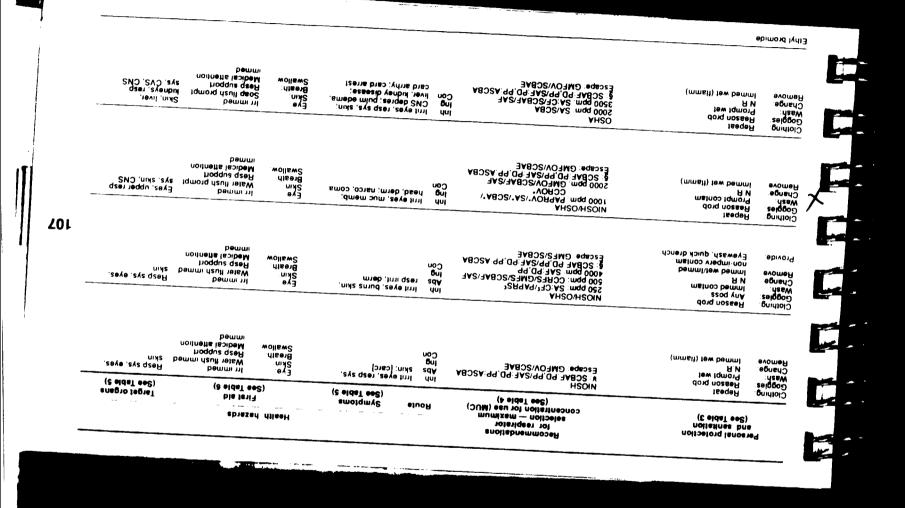
p Dichlorobenzene

Chemica structure/ CAS and R1	formule,	Synonyms, trade names, and conversion	Exposure Nmits (TWA	IDLH	Physical description	Chemical an propi	ortion	incompatibilities and reactivities	Messurement method (See Table 1)	
and DOT	and DOT ID and factors guide Nos.	factors	unless noted otherwise)			MW, BP, SOL FI.P, IP, Sp.Gr, flammability	VP, FRZ UEL, LEL			
3.3' Dichlorot (and its said	ts)	4,4 Diamino-3,3 dichloro- biphenyl; Dichlorobenzidine base;	NIOSH Ca See Appendix A	Ca	Gray to purple crystalline solid	MW: 253 1 BP: 788°F Sol: Almost	VP ? MLT 271 F UEL ?	None reported	Filter/ Si gel. Reagent.	
C ₆ H ₃ CiNH ₂ C ₆ 91-94-1	,H,CINH,	o.o' Dichlorobenzidine; 3.3' Dichlorobiphenyl 4.4' diamine;	OSHA (1910-1007) See Appendix B			Insoluble FLP? IP?	LEL ?		HPLČ/UVD 40 [#5509]	
DD0525000		3,3 · Dichloro·4,4 · biphenyl- diamine, ACG 3,3 · Dichloro 4,4 · diamino A2	ACGIH A2 (skin)							
Dichlorodifluo CCl ₂ F ₂	romethane	Difluorodichioromethane, Fluorocarbon 12, Freon*12, Halon*122, Propellani 12, Refrigerant 12	NIOSH/OSHA 1000 ppm (4950 mg/m ³)	50.000 ppm	Coloriess gas with an ether-like odor at extremely high	MW 120.9 BP: 22*F Sol(77*F)	VP >1 atm FRZ -252 F UEL NA	Chemically active metals such as sodium, potassium	Char(2) CH ₃ Cl ₂ GC/FID	
75 71 8 PA8200000					concentrations [Note: Shipped as a liquefied compressed gas]	0 03% Fi P·NA IP 11 75 eV	LEL NA	calcium, powdered aluminum, zinc & magnesium	 [#1018]	
1028 12	·	1 ppm - 5 03 mg/m ³				Nonflammable Ga	is			
1,3-Dichloro-5 dimethylhyd		Dactin, DDH, Halane	NIOSH/OSHA Ur 0.2 mg/m ³ ST C.4 mg/m ³	Unknown	nknown White powder with a chlorine-like odor.	MW: 197 0 BP: ? Sol: 0 2%	VP ? MLT 270°F UEL: ?	Water, strong acids, easily oxidized materials such as	None available	
C2H6CI2N2O2						FLP 346'F	LEL ?	ammonia salls & sulfides		
118 52-5 MU0700000										
						Sp.Gr. 1.5 Combustible Solid	1			
1,1-Dichloroet	hane	Asymmetrical dichloroethane; Ethylidene chloride:	NIOSH/OSHA 100 ppm	4000 ppm	Colorless, oily liquid with a	MW: 99.0 BP: 135*F	VP(77°F) 230 mm	Strong oxidizers, strong caustics	Char, CS	
снсі,сн,		1,1-Ethylidene dichioride	(400 mg/m ³)		chloroform-like odor	Sol. 0.6% FI.P(oc): 22"F	FRZ143°F	anding causiles	CS, GC/FID	
75-34 3 KI0175000						IP 11 06 eV	LEL 56%		[#1003, Haloge nated	
2362 27	62 27 1 ppm = 4.12 mg/m ³					Sp Gr. 1-18 Class IB Flammat	la Laura		Hydro carbons)	

Personal protection and sanitation (See Table 3)		Recommendations for respirator selection — maximum concentration for use (MUC) (See Table 4)		Route Symptoms (See Table 5)		Health hazarda		
						First aid (See Table 6)		
Clothing Goggles: Wash Change: Remove Provide:	Any poss Any poss Immed contam/daily After work if any poss contam Immed contam Eyewash, quick drench	NIOSH ¥. SCBAF:PD,PP/SAF:PD,PP:ASCBA Escape: HIEF/SCBAE	inh Abs Ing Con	Allergic skin reaction, sens, derm, head, dizz; causic burns; frequent urination, dysuria; hema; Gi upsets; upper resp infection; [carc]	Eye. Skin: Breath: Swallow.	Irr immed (15 min) Soap wash immed Resp support Medical attention immed	Bladder, liver, lung, skin, Gi tract	
Clothing: Goggles: Wash: Change: Remove:	Prevent wet or freezing Reason prob N.R. N R Immed wet (flamm)	NIOSH/OSHA 10,000 ppm: SA/SCBA 25,000 ppm: SA/CF 50,000 ppm: SCBAF/SAF §: SCBAF:PD, PP/SAF PD, PP:ASCBA Escape: GMFOV/SCBAE	Inh Con	Dizz, tremors, unconsciousness, card arrhy, card arrest	Eye: Skin: Breath:	Irr immed Water flush immed Resp support	CVS, PNS	
Clothing: Goggles Wash: Change Remove: Provide:	Repeat Any poss Prompt contam After work if reason prob contam Prompt non-imperv contam Eyewash	NIOSH/OSHA 2 mg/m ³ : SA/SCBA 5 mg/m ³ : SA/SCBAF/SAF § SCBAF: PD, PP/SAF: PD, PP: ASCBA § SCBAF: PD, PP/SAF: PD, PP: ASCBA Escape: GMFSHIE/SCBAE	inh Ing Con	linit eyes, muc memb, resp sys	Eye: Skin: Breath: Swallow.	Irr immed Soap wash prompt Resp support Medical attention immed	Resp sys, eye	
Clothing Soggles Vash: Change Temove:	Repeat Reason prob Immed wet N.R. Immed wet (flamm)	NIOSH/OSHA 1000 ppm: SA/SCBA 2500 ppm: SA/CF 4000 ppm: SCBAF/SAF §: SCBAF:PD,PP/SAF:PD,PP:ASCBA Escape: GMFOV/SCBAE	inh Ing Con	CNS depres; skin irril; liver, kidney damage	Eye: Skin: Breath: Swallow:	Irr immed Soap flush prompt Resp support Medical attention immed	Skin, liver, kidneys	

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Chemical name, structure/formula, CAS and RTECS Nos	Synonyma, trade names, and conversion factors	Exposure Hmits (TWA unless noted otherwise)	IDLH	Physical description	Chemical an prope	nd physical erties	incompatibilities and reactivities	Measurement method (See Table 1)
and DOT ID and guide Nos.					MW, BP, SOL FI.P, IP, Sp.Gr, flammability	VP, FRZ UEL, LEL		
Ethyl acrylate	Ethyl acrylate (inhibited), Ethyl ester of acrylic acid, Ethyl propenoate	NIOSH Ca Ca [2000 ppm] See Appendix A	Colorless liquid with an acrid	MW: 100 1 BP: 211"F	VP 29 mm FAZ: 96 F	Oxidizers, peroxides, polymerizers, strong	Char, CS	
CH,-CHCOOC,H,			12000 ppm3	odor	Sol 2% FLP 48 F	UEL 14%	alkalis, moisture, chiorosultonic acid	CS2. GC/FID
140 88 5 AT0700000		OSHA 5 ppm (20 mg·m ⁻¹) ST 25 ppm			IP. ?		Note Polymerizes readily unless an inhibitor such as hydroguingne is	[#1450. Esters I]
1917-27	1 ppm = 4.16 mg/m ³	(100 mg/m²) [skin]			Sp Gr 092 Class IB Flamma	ible Liquid	added [
Ethylamine	Aminoethane, Ethylamine (anhydrous),	NIOSH OSHA 10 ppm (18 mg/m³)	4000 ppm	Coloriess gas or water-white liquid (below 62°F) with an ammonia like odor [Note Shipped as a liquefied	MW: 45 1 BP: 62°F Sol. Miscible FLP: 1. F	VP >1 atm FRZ 114°F UEL 14 0% LEL 35%	Strong acids, strong oxidizers, copper, lin & zinc in presence of moisture, cellulose nitrate	Si gel. H.SO, GC/FID II(3) [#S144]
CH ₃ CH ₂ NH ₂	Monoethylamine							
75 04 7 KH2100000					ÌP 8 86 ⊎V			
10 3 6 68	1 ppm – 1 87 mg/m ³			compressed gas [Sp Gr. 0.69 (Liquid) Flammable Gas Class IA Flammable Liquid			
Ethyl benzene	Ethylbenzol. Phenylethane	NIOSH/OSHA 100 ppm (435 mg/m ³) ST 125 ppm	n ³) Sm	Coloriess liquid with an aromatic odor	MW 106 2 BP: 277 F Sol 0 01% FLP 55 F IP 8 76 eV	VP(79"F) 10 mm FRZ 139 F UEL 6 7% LEL 10%	Strong uxidizers	Char. CS2. GC/FID.
сн,сң,с,н,	i neny ieliane							
100 41 4 D A07000 00		(545 mg/m ³)						{#1501. Aromatic Hydro carbons]
1175-26	1 ppm = 4 41 mg/m+				Sp Gr 0 87 Class IB Flammat	ble Liquid		carbons
Ethyl bromide	Bromoethane	omoethane NIOSH See Appendix D OSHA		Coloriess to yellow liquid with an ether-like odor	MW: 109.0 BP: 101°F Sol. 0.9% FLP: <4°F	VP(70-F)	Chemically active metals such as sodium, potassium,	Char, 2 Propanol; GC/FID,
Сн,СН,Вг						400 mm FRZ 182°F		
74 96 4 KH6475000		200 ppm (890 mg/m ³) ST 250 ppm (1110 mg/m ³)		Note A gas above 101 F	IP 10 29 eV	UEL 80% LEL 68%	calcium, powdered aluminum, zinc & magnesium	 [#1011]
1891 58	1 ppm = 4 53 mg/m ³				Sp.Gr 1 46 Class IB Flammat			



	Chemical name, structure/lormula, CAS and RTECS Nos and DOT ID and guide Nos.	le, trade names,	Exposure limits (TWA uniess note(otherwise)	IDL) d	1 Physical description	properties		and reactivities	 Messurement method (See Table 1)
	Ethyl butyl ketone	Butyl ethyl ketone. 3 Heptanone	NIOSH/OSHA			MW, BP, SOL FI.P, IP, Sp.Gr. VP, FRZ flammability UEL, LEL			
	CH,CH,CO(CH,],CH 106 35 4 MJ\$250000	J TEPIANONE	50 ppm (230 mg/m ³)	3000 pp	om Coloriess liquid with a powerful fruity odor	MW 114 2 BP 298"F Sol 1% FI P(oc) 115 IP 9 02 eV	VP 4 mm FRZ 38 F	Oxidizers acetaldehyde, perchloric acid	Char. Methanol/ CS2 GC/FID
	Ethyl chloride CH ₁ CH Cl	1 ppm = 4 75 mg/m3 Chloroethane, Hydrochloric ether,	NIOSH	20.000		Sp Gr 0 82 Class II Comb	ustible Liquid		lli [#1301, Kelones II]
B	75 00 3 KH7525000	Monochloroethane. Muriatic ether	See Appendix D Handle with caution in the workplace.	20,000 ppm	Colorless gas or liquid (below 54°F) with a pungent, ether like odor	MW 64 5 BP: 54°F Sol: 0.6% FLP: NA (Gas)	VP >1 atm FRZ 218°F	Chemically active metals such as sodium, potassium,	Char(2) CS, GC/FID
	1037-27 Ethylene chlorohydrin	1 ppm - 2 68 mg.m3 drin 2 Chloroethanol, 2 Chloroethyl alcohol, Ethylene chlorhydrin	OSHA 1000 ppm (2600 mg/m³)		[Note: Shipped as a liquelied compressed gas]	IP 10 97 eV Sp Gr 0 92 (Liquid at 32 F) Flammable Gas		aluminuin, zinc & magnesium, oxidizers water or steam	111
	CH,CICH,OH		NIOSH OSHA C 1 ppm (3 mg/m ³) (skin)	10 ppm	Colorless liquid	Class IA Flammable Liquid		[Note Reacts with water to form hydrochloric acid]	
,	107 07 3 KK0875000				with a faint, ether-like odor	BP 262 F Soi Miscible FIP 140 F IP 10 90 eV	VP.5 mm FRZ: 90°F UEL 15 9% LEL 4 9%	Strong oxidizers, strong caustics, water or steam	Char(pet) 2 Propanol/ CS ₂ ; GC/FID
-	135 55	1 ppm = 3 35 mg/m3				Sp Gr 1 20			 [#2513]
	hylenediamine H ₂ CH ₂ CH ₂ NH ₂	1.2 Diaminoethane, 1.2 Ethanediamine:	NIOSH/OSHA	2000 ppm	Colorless, viscous	Class IIIA Combustible Liquid			
10	7 15 3 18575000	Ethylenediamine (anhydrous)	(25 mg/m ³)		ammonia like odor [Note: A solid	MW: 60, 1 BP: 241°F Sol Miscible Fi P: 93°F IP: 8 60 eV	VP: 11 mm FRZ: 47"F UEL: 14 4% LEL: 4 2%		XAD 2*, DMF, HPLC/UVD, III
160	04 29	1 ppm - 2 50 mg/m3				Sp Gr 0 91 Class IC Flammab			[#2540]

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Recommendations for respirator selection — maximum concentration for use (MUC) Health hazards Personal protection and sanitation (See Table 3) Symptoms First ald Target organs Route (See Table 4) (See Table 5) (See Table 6) (See Table 5) Clothing Goggles Wash Repeat Reason prob NIOSH/OSHA Inh freit eyes, muc memb; head; narco, coma; derm Eye. Skin: Breath. Irr immed Water flush Eyes, skin, resp NIOSH/OSHA 500 ppm. CCROV*/SA*/SCBA* 1000 ppm. PAPROV*/CCRFOV 1250 ppm. SA CF* 2500 ppm. GMFOV/SCBAF/SAF 3000 ppm. SAF PD.PP 5 SCBAF PD.PP.SAF PD.PP ASCBA Escape. GMFOV-SCBAE Ing Con sys Promot contam Resp support Change NR. Swallow Medical attention Remove Prompt non imperv immed contam

Ť	Clothing Goggles Wash Change Remove	Repeat Reason prob N R N R Immed wet (flamm)	OSHA 10.000 ppm SA*/SCBA* 20.000 ppm: SA CF*/SCBAF/SAF § SCBAF PD.PP/SAF PD.PP ASCBA Escape GMFOV/SCBAE	inh Abs Ing Con	Inco, inebriale, abdom cramps, card arrhy, card arrest, liver, kidney damage	Eye. Skin Breath Swallow	fir immed Water flush prompt Resp support Medical attention immed	Liver, kidneys, resp.sys, CVS	
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NIOSH/OSHA 10 ppm. SCBA*/SA* § SCBAF PD.PP/SAF PD.PP ASCBA Escape GMFOV/SCBAE Clothing Eye Skin Any poss Inh Irrit muc memb; nau, Irr immed Goggles Wash Any poss Abs vomit, verti, inco; Water flush immed Immed contam Ing Con numb; vis dist; head; Breath Resp support Change NR thirst, delirium, low BP, Swallow Medical attention Remove Immed non impervisionitam collapse, shock, coma immed Provide Evewash, quick drench

Clothing	Any poss	NIOSH/OSHA	Inh	Nasal irrit, primary irrit;	Eve	Irr immed	Resp sys, live
Goggies	Any poss	250 ppm: SA:CF ¹ /PAPRS ¹	Abs	sens derm, irrit resp sys.	Skin	Water flush immed	kidneys, skin
Wash	Immed contam daily	500 ppm: CCRFS/GMFS/SCBAF/SAF	Ina	asthma: liver, kidney	Breath	Resp support	
Change	After work if any poss	2000 ppm: SAF PD PP	Con	damage	Swallow	Medical attention	
5	contam	SCBAF PD.PP/SAF PD.PP ASCBA	00	contage	Children of	mmed	
Remove	Immed wet/Immed	Escape GMFS/SCBAE					
	non imperv contam						
Provide	Evewash (>5%), quick						
	drench						

Ethylenediamine

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Resp sys, liver kidneys, CNS, skin, CVS

Chemical name, structure/formula, CAS and RTECS Nos.,	Synonyms, trade names, and conversion	Exposure Nmits (TWA	IDLH	description properties d			properties		incompatibilities and reactivities	Measuremen method (See Table 1
and DOT ID and guide Nos.	factors	unless noted otherwise)						(200 1200 1		
Hexachloroethane	Carbon hexachloride. Perchloroethane	NIOSH Ca	Ca (300 ppm)	Coloriess crystals with a camphor-	MW: 236-7 BP: Sublimes	VP: 0.2 mm MLT: 368 F	Alkalis, metals such as zinc, cadmium.	Char, CS.		
CI,CCCI,	- ercinoroemano	See Appendix A	[996 55.1]	like odor.	Sol(72°F) 0 005%	(Sublimes) UEL NA	aluminum, hot iron & mercury	CS; GC/FID		
67-72 1 K14025000		(10 mg/m ³) {skin} OSHA 1 ppm			FI P. NA IP 11 22 8V	LEL NA	,	(#1003 Haloge nated Hydro		
9037 53		(10 mg m) [skin]			Sp.Gr. 2.09 Noncombustible	Solid		carbons		
Hexachloronaphthalene	Halowax® 1014	NIOSH/OSHA 0 2 mg/m ³	2 mg/m³	White to light- vellow solid with	MW: 334 9 BP: 650 730"F	VP <1 mm ML1 279 F	Strong oxidizers	Filter,		
C, JH, CL		(skin)		an aromatic odor	Sol Insoluble	UEL NA		Hexane. GC/ECD, II(2)		
1335 87 1 QJ7350000					IP ?			[#\$100]		
					Sp.Gr. 1.78 Noncambustible Solid					
n Hexane	Hexane,	NIOSH/OSHA	5000 ppm	Coloriess liquid	MW: 86 2	VP(77"F)	Strong oxidizers	Char.		
Сн,[Сн,],Сн,	Hexyl hydride, Normal-hexane	50 ppm (180 mg/m³)		with a gasoline- like odor.	8P: 156°F Sol. 0.002% FLP - 7°F	150 mm FRZ 219 F UEL: 7.5%		CS, GC/FID.		
110 54 3 MN9275000					IP 10 18 ⊌V	LEL 11%		[#1500. Hydro carbons]		
1208 27	1 ppm = 3.58 mg/m3				Sp Gr: 0.66 Class IB Flammal	ble Liquid				
2-Hexanone	Butyl methyl ketone, MBK.	NIOSH	5000 ppm	Coloriess liquid	MW: 100.2	VP(77°F)	Strong oxidizers	Char,		
CH3CO[CH2]3CH3	Methyl butyl ketone,	1 ppm (4 mg/m³)		with an acetone- like odor.	BP: 262°F Sol: 2% FLP: 77°F	4 mm FRZ 71 F UEL 8%		CS ₂ ; GC/FID; III		
591-78-6 MP1400000	Methyl n-butyl ketone	OSHA 5 ppm (20 mg/in³)	n		IP 9:34 eV	LEL?		III [#1300, Ketones I		
	1.ppm = 4.17 mg/m ³				Sp Gr. 0 81 Class IC Flammal	ble Liquid				

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8	rsonal protection and sanitation (See Table 3)	Recommendations for respirator selection — maximum concentration for use (MUC) (See Table 4)	Ro	ute Symptoms (See Table 5)		azards First aid na Table 6)	Target organa (See Table 5)
Clothing Goggles Wash Change Remove	Repeat N A Prompt contam-daily After work if reason prob contam Prompt non imperv contam	NIOSH V SCBAF PD,PP/SAF PD,PP ASCBA Escape GMFOV/SCBAE	Inh Abs Ing Con		Eye Skin: Breath Swallow	fir immed Soap wash immed Resp support Medical attention immed	Eyes
Clothing Goggles Wash Change	Any poss moll Repeat lig sol Reason prob- lumes Any poss moll Reason prob-lig Prompt contain daily Atter work if reason prob-contain	NIOSH OSHA 2 ing in' sa' SCBA' § SCBAF PD PP'SAF PD PP ASCBA Escape: GMEOV SCBAE	Inti Abs Ing Con	· •	Eye Skin Breath Swallow	fir immed Soap wash prompt Resp support Medical attention immed	liver skir.
Remove Clothing Goggles Wash Change Remove	Immed wet mott Prompt non imperv contain lig Repeat Reason prob Prompt contain N R Inimed wet (flamin)	NIOSH OSHA 500 ppm SA':SCBA' 1250 ppm SA CF' 2500 ppm SAT CF':SCBAF:SAF 5000 ppm SAF PD.PP § SCBAF PD.PP/SAF PD.PP ASCBA Escape GMFOV SCBAF	Inh Ing Con	Li head, nau, head, numb extremities, musc weak, irrit eyes, nose, derm, chemical pneu, gidd	Eye Skin Breath Swallow	Irr immed Soap wash immed Resp support Medical attention immed	Skin ayes rasp sys
Clothing Goggles Wash. Change Remove	Reason prob Reason prob Prompt contam N R Immed wet (flamm)	NIOSH 10 ppin SA SCBA 25 ppin SA CF 50 ppin SCBAF/SAF-SAI CF 2000 ppin SAF-PD.PP § SCBAF-PD.PP'SAF-PD.PP ASCBA E-scape GMFOV-SCBAE	inh Abs Ing Con	hrit eyes, nose, peri neur weak, pares, derm, head, drow	Eye Skin Breath Swallow	Irr immed Soap wash in sed Resp support Medical attention immed	CNS skin, resp sys

Chemical name, structure/formula, CAS and RTECS Nos	Synonyms, trade names, and conversion	Exposure Smits (TWA	IDLH	Physical description	Chemical an propi		incompatibilities and reactivities	Measuremeni method (See Table 1)
and DOT ID and guide Nos.	factors	uniess noted otherwise)			MW, BP, SOL FI.P, IP, Sp.Gr, Hammability	VP, FRZ UEL, LEL		
Methyl chloroform	Chlorothene, 1,1,1 Trichloroethane,	NIOSH C 350 ppm	1000 ppm	Coloriess liquid with a mild.	MW 1334 BP 165°F	VP 100 mm FRZ 23 F	Strong caustics, strong oxidizers,	Char, CS-
CH ₁ CCI, 71 55 6	1.1.1 Trichloroethane (stabilized)			chlaroform like adar	Sol 0.4% FLP None	UEL 12.5% LEL 7.5%	chemically active metals such as zinc	CS, GC/FID III
KJ2975000		OSHA 350 ppm (1900 mg/m²)			IP 11.00 eV Sp.Gr. 1.34 Noncombustible	Liquid	aluminum, magnesium powders, sodium & potassium, water (Note: Reacts slowly	(#1003) Haloge nated Hydro
2831 74	1.ppm = 5.55 mg/m [⊥]	ST 450 ppm (2450 mg/m ³)			however the vapr burn.	or will	with water to form hydrochloric acid]	cárbons]
Methylcyclohexane	Cyclohexylmethane. Hexahydrotoluerie	NIOSH/OSHA 400 ppm	10.000 ppm	Colortess liquid with a faint.	MW 98.2 BP 214 F	VP(72 F) 43 mm	Strong oxidizers	Char CS,
CH₁C₅H₁,		(1600 mg·m ¹)	pp	benzene like odor	Sol Insuluble FLP 25"F	FRZ 196 H		GC/FID. III
108 87 2 GV6125000					IP 985 eV	LEL 12%		(#1500, Hydro carbons)
2296 27	1 ppm - 4 08 mg/m³				Sp.Gr. 0.77 Class IB Flammai	ble Liquid		
Methylcyclohexanol	Hexahydrocresol, Hexahydromethylphenol	NIOSH/OSHA	10.000 ppm	Straw-colored liquid with a	MW: 114 2 BP: 311-356"F	VP(86°F) 2 mm	Strong oxidizers	Char,
CH,C,H,JOH	nexanyorometry prenor	(235 mg/m ³)		weak odor like coconut oil	Sol 4% FIP 154°F	FRZ 58 F		CH ₂ Cl ₂ GC/FID II(4)
25639 42 3 GW0175000					1P 9	LEL?		[#\$374]
2617-26	1 ppm - 4 75 mg/m³				Sp Gr - 0 92 Class IIIA Combu	slible Liquid		
Methylcyclohexanone	2 Methylcyclohexanone	NIOSH/OSHA	2500 ppm	Colorless liquid	MW: 112.2 BP: 325°F	VP: 1 mm	Strong oxidizers	Porapak,
CH'C'H'O		30 ppm (230 mg/m³) ST 75 ppm		with a weak peppermint like odor.	Sol. Insoluble	(approx) FRZ: 7°F UEL: ?		Acetone, GC/FID; III
583-60-8 GW1750000		(345 mg/m ³ (skin)			10 7 19 7	LEL?		[#2521]
2297 26	1 ppm - 4 66 mg/m ³				Sp.Gr. 0.93 Class II Combusti	ble Liquid		

	Personal protection and sanitation (See Table 3)		Recommendations for respirator selection — maximum concentration for use (MUC) (See Table 4)	Ro	ute Symptome (See Table 5)		Health hazarda First aid (See Table 6)		
Gog Was Chai	içləs: sh:	Repeat Reason prob Prompt wet N R Prompt non imperv wet	NIOSH/OSHA 1000 ppm: SA'/SCBA* § SCBAF PD, PP/SAF PD, PP ASCBA Escape: GMFOV/SCBAE	Inh Ing Con	Head, lass, CNS depres, poor equi; irrit eyes; derm; card arrhy	Eye. Skin: Breath. Swallow:	Irr immed Soap wash prompt Resp support Medical attention immed	Skin, CNS, CVS eyes	
Clott Gogg Wasi Char Rem	ih: inge:	Repeat Reason prob Prompt wet N.R. Immed wet (fiamm)	NIOSH/OSHA 4000 ppm: SA/SCBA 10,000 ppm: SA/SCBA § SCBAF PD.PP/SAF:PD.PP'ASCBA Escape: GMFOV/SCBAE	Inh Ing Con	Li head, drow; skin, nose, throat irrit	Eye: Skin. Breath: Swallow:	Irr immed Soap wash prompt Resp support Medical attention immed	Resp sys, skin	
Cloth Gogg Wash Chan Remi	gles. h. nge:	Repeat Reason prob Prompt contam N R Prompt non imperv contam	NIOSH/OSHA 500 ppm: SA'/SCBA' 1250 ppm: SA CF' 2500 ppm: SBAF SAF 10,000 ppm: SAF.PD,PP § SCBAF PD,PP/SAF.PD,PP,ASCBA Escape. GMFOV/SCBAE	lah Abs Ing Con	Head, irrit eyes, upper resp sys; in animals narco, liver, kidney damage	Eye: Skin: Breath Swallow	Irr immed Soap wash prompt Resp support Medical attention immed	Resp sys, skin, eyes; in animals CNS, liver, kidneys	
Cloth Gogg Wash Chan Remo	gleš: h: nge:	Repeal Reason prob Prompt contam N R Prompt non imperv contam	NIOSH/OSHA 500 ppm. SA*/SCBA* 1250 ppm. SA CF* 2500 ppm. SCBAF/SAF § SCBAF.PD,PP/SAF.PD,PP.ASCBA Escape. GMFOV/SCBAE	inh Abs ing Con	in animais narco; irrit eyes, muc memb; derm	Eye. Skin: Breeth: Swallow	Irr immed Scap wash prompt Resp support Medical attention immed	in animais, resp sys, liver, kidneys, skin	

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o Methylcyclohexanone

Chemical name, structure/formula, CAS and RTECS Nos	Synonyms, trade names, and conversion	Exposure timits (TWA	IDLH	Physical description	Chemical an prop		incompatibilities and reactivities	Measurement method (See Table 1)
and DOT ID and guide Nos.	factors	unièss noted otherwise)			MW, BP, SOL FLP, IP, Sp.Gr, Itemmability	VP, FAZ UEL, LEL		
Methylene bisphenyl isocyanate	4,4: Diphenylmethane diisocyanate; MDI	NIOSH 0.05 mg/m² (0.005 ppm)	100 mg/m3	While to light yellow, odorless liakes.	MW 250 3 BP: 342"F Sol: 0.2%	VP(104 'F) 0 001 mm MLT 99'F	Strong atkalis, acids, alcohol	Bub, Acetylate, HPLC/UVD,
CH ₂ (C ₆ H ₄ NCO) ₂ 101-68-8	Methylene bis{4 phenyl isocyanate; Methylene di piphenylene	C 0 2 mg/m ³ (0 020 ppm) (10 min)		[Note: A liquid above 99: F]	FI P(DC) 396 F IP 2	UEL ? LEL ?		 #5521]
NQ9350000 2489 53	ester of isocyanic acid	OSHA C 0 2 mg/m²) (0:02 ppm)			Sp Gr(122 F) 1 Combustible Soli			
Methylane chloride CH,CI,	Dichloromethane, Methylene dichloride	NIOSH Ca See Appendix A	Ca [5000 ppm]	Colorless liquid with a chloroform- like odor.	MW: 84.9 BP: 104"F Sol. 2%	VP 350 mm FRZ: 139"F UEL: 22%	Strong Oxidizers, caustics, chemically active metals such as	Char(2). CS ₂ : GC/FID.
75 09 2 PA8050000		Reduce exposure to lowest feasible concentration		(Note: A gas above 104 F]		LEL. 14%	atuminum, magnesium powders, potassium & sodium, concentrated	
1593 74	1.ppm = 3.53 mg/m ³	OSHA 500 ppm C 1000 ppm 2000 ppm (5 mm a	ACGIH A2: 50 ppm (175 mg/m ³) hax peak in ar)	Sp. Gr. 1-33 Combustible Liquid		nitric acið	
Methyl formate	Methyl ester of formic acid,	NIOSH OSHA	5000 ppm		MW 60 1	VP. 476 mm	Strong oxidizers	Carbo B(2).
нсоосн,	Methyl methanoate	100 ppm (250 mg/m ³)		with a pleasant odor [Note: A gas above	BP 89°F Sol 30% FLP 2°F	FRZ: -148"F UEL: 23% LEL: 4.5%	Note Reacts slowly with water to form	Ethyl acetate;
107 31 3 LQ8925000	(300 mg/m²) ST 150 ppm (375 mg/m³)			1P 10 82 eV	LEL 4 5%	methanol & formic acid)	GC/FID; II(5) {#S291}	
1243-26 1 ppm = 2.50 mg/m ³					Sp.Gr. 0.98 Class IA Flammable Liquid			
5-Methyl-3-heptanone	Aniyi ethyi ketone, Ethyi amvi ketone	NIOSH/OSHA	3000 ppm	Coloriess liquid	MW: 128 2 BP: 315*F	VP:2 mm	Strong oxidizers	Char, Methanol/
сн,сн,с ојсн ,),сн,		23 ppm (130 mg/m³)		with a pungent odor	Sol. Insoluble FLP: 138°F	FAZ: 70"F UEL.? LEL.?		CS, GC/FID,
541 85 5 MJ7350000					IP.?	LEL		III. [#1301, Ketones
2271 26	1 ppm = 5 33 mg/m³				Sp Gr 0 82 Class II Combusti	bie Liquid		113

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rsonal protection and sanitation (See Table 3)	Recommendations for respirator						
		I			First aid		
Any poss Prompt contam After work if reason prob contam Prompt non imperv contam	NIOSH 2 mg/m ³ SA*/SCBA* 5 mg/m ³ SA CF* 10 mg/m ³ SCBAF/SAF 100 mg/m ³ SCBAF/PD,PP 5 SCBAF PD,PP/SAF PD,PP ASCBA § SCBAF PD,PP/SAF PD,PP ASCBA	Ini Ini Ci	nh Irriteyes, nose, throat;	Eye	Irr immed Soap wash immed Resp support		
	Escape GMFOVHIE/SCBAE				immed immed		
Repeat Reason prob Prompt wet N R	NIOSH V. SCBAF PD.PP/SAF.PD.PP ASCBA Escape GMFOV/SCBAE	Inh	I head limbs such	Eye	Irr immed		
Prompt non imperviwet	UNIT OVIGUEAE	Con		Skin. Breath Swallow	irr immed Soap wash prompt Resp support Medical attention immed	Skin, CVS, eyes CNS	
lepeal							
leason prob frompt wet IR nmed wet (flamm)	NIOSH/OSHA 1000 ppm: SA*/SCBA* 2500 ppm: SA*/SCF 5000 ppm: SCBAF/SAF §: SCBAF PD,PP/SAF:PD,PP ASCBA Escape: GMFOV/SCBAE	Ing	Opression due	Eye. Skin: Breath. Swallow	Irr immed Soap wash immed Resp support Medical attention immed	Eyes, resp sys, CNS	
eson prob	NIOSH/OSHA 250 ppm: SA*/SCBA*	Inh I	Irrit eyes, muc memb:				
e All the second	625 ppm: SA:CF*/PAPROV* 1000 ppm: CCRFOV 1250 ppm: GMFOV/SCBAF/SAF 3000 ppm: SAF:PD, pp § SCBAF PD, PP/SAF:PD, PP.ASCBA Escape: GMFOV/SCBAE	ing F Con	read, narco, coma; derm	Breath		Eyes, skin, resp sys, CNS	
	Reason prob Any poss Prompt contam After work if reason prob contam Prompt non imperv Contam Repeat Reason prob Prompt non imperv wet Prompt non imperv wet Repeat Repeat Reason prob Prompt non imperv wet Repeat R	Resentation Tor respirator (See Table 3) selection maximum concentration for use (MUC) (See Table 4) Reason prob NKOSH Any poss 2 mg/m ³ SA'/SCBA* Prompt contam 5 mg/m ³ SCBAF/SAF prob contam 10 mg/m ³ SCBAF/SAF prob contam 10 mg/m ³ SCBAF/PD, PP prob contam 10 mg/m ³ SCBAF/PD, PP prompt non imperv § SCBAF PD, PP/SAF.PD, PP ASCBA Repeat NIOSH Prompt non imperv wet V SCBAF PD, PP/SAF.PD, PP ASCBA Repeat NIOSH/OSHA Prompt non imperv wet 1000 ppm: SA'/SCBA* Prompt wet 2500 ppm: SA'/SCBA* Prompt wet 5 SCBAF PD, PP/SAF, PD, PP ASCBA Escape: GMFOV/SCBAE Escape: GMFOV/SCBAE Prompt wet 5000 ppm: SA'/SCBA* Prompt wet 500 ppm SA'/SCBA*	Restriction Tor respirator (See Table 3) selection - maximum concentration for use (MUC) (See Table 4) Reason prob NKOSH Any poss 2 mg/m ³ SA'/SCBA* Prompt contam 2 mg/m ³ SA'/SCBA* prob contam 10 mg/m ³ SCBAF/SAF prob contam 10 mg/m ³ SCBAF/SAF prompt non imperv § SCBAF PD, PP/SAF.PD, PP contam 100 mg/m ³ SCBAF, PD, PP / ASCBA Repeat NIOSH Prompt non imperv wet V.SCBAF PD, PP/SAF, PD, PP ASCBA Prompt non imperv wet V.SCBAF PD, PP/SAF, PD, PP ASCBA Prompt non imperv wet V.SCBAF PD, PP/SAF, PD, PP ASCBA Prompt non imperv wet SCBAF PD, PP/SAF, PD, PP ASCBA Prompt non imperv wet NOSH/OSHA Repeat NIOSH/OSHA Repeat NIOSH/OSHA Repeat NIOSH/OSHA Prompt non imperv wet 1000 ppm: SA'/SCBA* Repeat NIOSH/OSHA Repeat NIOSH/OSHA Repeat NIOSH/OSHA Repeat NIOSH/OSHA Repeat NIOSH/OSHA Repoint wet 250 ppm: SA'/SCBA*	Research Nicshi Nicshi	Internation Internation Internation Internation Health Gee Table 3) Concentration for use (MUC) Route Symptoms Reason prob NOSH Inh Irrit eyes, nose, throat; Eye Any poss NOSH Inh Irrit eyes, nose, throat; Eye Any poss Sign prob Sign prob Sign prob Sign prob Atter work if reason 5 mg/m² SA (F) PD, PP ASCBA Inh Irrit eyes, nose, throat; Eye Prompt contam 5 mg/m² SA (F) PD, PP ASCBA Con chest pain, dysp, asthma Breath Prompt non imperv § SCBAF PD, PP/SAF PD, PP ASCBA Inh Fig. weak, sleepiness, slima Eye Repeat NIOSH ¥ SCBAF PD, PP/SAF PD, PP ASCBA Inh Fig. weak, sleepiness, skin, interest Eye NB Breant Escape GMFOV/SCBAE Inh Fig. weak, sleepiness, skin, interest Eye Prompt wet 2500 ppm: SA'/SCBA* Inh Eye, nose irrit, chest Eye Prompt wet 2500 ppm: SA'/SCBA* Inh Eye, nose irrit, chest Eye R 2500 ppm: SA/SCBA* Inh Eye,	Recommendations disanitation (See Table 3) Recommendations tor respirator selection - meatimum concentration for use (MUC) (See Table 4) Route (See Table 5) Health hazerds (See Table 6) Reason pub Any poss Arise work if reason prob contam NKOSH 2 mg/ml SA/SCBA* Inh 10 mmed Inh 10 mmed	

Chemical name, structure/formule, CAS and RTECS Nos.,	Synonyms, trade names, and conversion	Exposure limits (TWA	IDLH	Physical description	Chemical an propi		incompatibilities and reactivities	Measurement method (See Table 1)
and DOT ID and guide Nos.	factors	uniess noted otherwise)			MW, BP, SOL FI.P, IP, Sp.Gr, flammability	VP, FRZ UEL, LEL		. ,
Styrene	Ethenyi benzene, Phenyiethylene,	NIOSH/OSHA 50 ppm	5000 ррт	Coloriess to vellow, oily	MW: 104 2 BP: 293*F	VP: 5 mm FRZ: 23 /F	Oxidizers, catalysts for vinyi polymers,	Char, CS
C ₆ H ₅ CH-CH ₂	Styrene monomer, Styrol.	(215 mg/m ³) ST 100 ppm		liquid with a sweet, floral	Sol Slight	UEL 7.0%	peroxides, strong acids, aluminum	CS2 GC/FID
100 42 5 WL 3675000	Vinyl benzene	(425 mg/m³)		odor	IP. 8 40 eV		chloride [Note: May polymerize if contaminated or subjected to heat	(#1501, Aromatic Hydro- carbonst
2055 27 (inhibited)	1 ppm - 4 33 mg/m³				Sp Gr 0 91 Class IC Flamma	ble Liquid	Usually contains an such as tert-butylcat	inhibitor
Sultur dioxide	Sulturous acid anhydride. Sulturous oxide,	NIOSH/OSHA 2 ppm	100 ppm	Colorless gas with a characteristic,	MW 64 1 BP: 14*F	VP >1 atm FRZ 104 F	Powdered and alkali metals such as	Filters(2), NaHCO
SO ₂ 7446 09 5 WS 45500 00	Sulfur oxide	(5 mg/m³) ST 5 ppm (10 mg/m³)		irritating, pungent odor (Note: A liquid below 14°F. Shipped as a	Sol. 10% FIP NA IP 12 30 ⊎V	UEL NA LEL NA	sodium & potassium, water, ammonia, aluminum [Note: Reacts with water to form	Na₂CO₃, IC, Itt [#6004]
10/9 16	1 ppm = 2 66 mg/m³			liquefied compressed gas }	Nonflammable Gas		sulfuric acid.)	
Sulfuric acid	Battery acid, Hydrogen sulfate,	NIOSH/OSHA 1 mg/m3	80 mg/m ³	Coloriess to dark brown, oily,	MW: 98 1 BP: 554 F	VP(295 F)	Organic materials, chlorates, carbides,	Si gel, NaHCO,/
H ₂ SO ₄	Oil of vitriol, Sulfuric acid (aqueous)	•		adarless liquid (Note: Pure	Sol Miscible	FRZ 51 F	fulminates, water, powdered metals	Na ₂ CO ₃ . IC;
7664 93 9 WS5600000				compound is a solid below 51 F	IP ? LEL NA		[Note: Reacts violently with water	HI [#7903.
1830–39 (51-95% acid) 1831–39 (fuming) 1832–39 (spent)	1 ppm = 4 08 mg/m³			Often used in an aqueous solution j	Sp Gr 1 84 (96-9 Noncombustible L but capable of ign finely divided com	liquid. Iting	with evolution of heat. Corrosive to metals] is	Inorganic Acids]
Sullur monochloride	Sulfur chloride, Sulfur subchloride,	NIOSH/OSHA C 1 ppm	10 ppm	Light amber to velice-red, oily	MW 135.0 BP: 280°F	VP 7 mm FAZ: 107°F	Peroxides, oxides of phosphorous,	None available
S ₂ Cl ₂	Thiosulturous dichloride	(6 mg/m²)		liquid with a pungent,	Sol. Decom	UEL ?	organics, water (Note Decomposes	
10025 67 9 WS4300000				nauseating, irritating odor	FLP: 245°F IP: 9:40 eV		violently in water to form hydrochloric acid, sulfur dioxide, sulfur, sulfite, thiosulf	ate
1828 39	1 ppm = 5 61 mg/m ³				Sp Gr 1 68 Class IIIB Combus	stible Liquid	and hydrogen sulfide Corrosive to metals)	

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	rsonal protection and sanitation	Recommendations for respirator selection — maximum				Health h	azarda	
	(See Table 3)	concentration for use (MUC) (See Table 4)	Ro	ute	Symptoms (See Table 5)	 (Se	Target organ (See Table 5	
Clothing Goggles: Wash Change Remove	Repeat Reason prob Prompt contam N R Immed wet (flamm)	NIOSH/OSHA 500 ppm CCROV//SA*/SCBA* 1000 ppm CCROV/PAPROV* 1250 ppm SA CF* 2500 ppm GMFOV/SCBAF/SAF 5000 ppm SAF PD,PP § SCBAF PD,PP SAF PD,PP ASCBA Escape GMFOV/SCBAE	Inh Ing Con	weak,	es, nose, drow, unsteady gait; defatting derm	Eye Skin: Breath Swallow	Irr immed Water flush Resp support Medical attention immed	CNS, resp sys. eyes, skin
Clothing Goggles Wash Change Remove Provide	Prevent skin freezing Any poss N R N R Immed wet Eyewash	NIOSH/OSHA 20 ppm CCRS*/SA*/SCBA* 50 ppm PAPRS*/SA CF* 100 ppm CCRFS/GMFS/PAPRTS*/ SCBAF/SAF/SATCF* § SCBAF PD,PP/SAF PD,PP ASCBA Escape GMFS/SCBAE	Inh Can	rhin, c reflex i	es, nose, throat, hoking, cough; bronchoconstriction, kin burns	Eye. Skin Breath	lrr immed Water flush immed Resp support	Resp sys, skin eyes
Clothing Goggles Wash Change Remove Provide	Any poss >1% Repeat <1% Any poss Immed contain N R Immed non imperv contain >1% Eyewash, quick drench	NIOSH/OSHA 25 mg/m ³ : PAPRAGHIE ⁵ /SA CF ⁴ 50 mg/m ³ : CCRFAGHIE/SCBAF/SAF/ GMFAGHIE 80 mg/m ³ : SAF-PD,PP §: SCBAF PD,PP/SAF PD,PP ASCBA Escape: GMFAGHIE/SCBAE	Inh Ing Con	pulm e emphy dental	ose, throat irrit, dema, bron; , con; stomatis, erosion, trachbronc, ye burns, derm	Eye Skin Breath Swallow	fri immed Water flush immed Aesp support Medicat attention immed	Resp sys, eyes skin, teeth
Clothing Goggles Wash Change Remove Provide	Any poss Any poss Immed contam N R Immed non impery contam Eyewash, quick drench	NIOSH/OSHA 10 ppm PAPRS ¹ /CCRFS/GMFS/ SCBAF/SAF § SCBAF PD.PP/SAF.PD.PP ASCBA Escape GMFS/SCBAE	inh Con Irig		ough, burn eyes, ulm ederna	Eye. Skin Breath Swallow	Irr immed Water flush immed Resp support Medical attention immed	Resp.sys.skin. eyes

Chemical name, structure/formula, CAS and RTECS Nos	Synonyms, trade names, and conversion	Exposure limits (TWA	IDLH	Physical description	Chemical an prope		incompatibilities and reactivities	Measurement method (See Table 1)		
and DOT ID and guide Nos.	factors	uniess noted otherwise)			MW, BP, SOL FI.P, IP, Sp.Gr, flammability	VP, FRZ UEL, LEL		(,		
1,1,2.2 Tetrachloroethane	Acetylene tetrachloride. Symmetrical tetrachloro- ethane	NIOSH Ca See Appendix A	Ca [150 ppm]	Coloriess to pale- yellow liquid with a pungent,	MW: 167.9 BP: 296°F Sol: 0.3%	VP(86"F): 9 mm FRZ: -33 F	Chemically active metals, strong caustics, fuming	Char(pel); CS ₂ ; GC/FID;		
CHCI2CHCI2	01110-14	1 ppm (7 mg/m ³)		chloroform like	FLP. NA IP. 11.10 eV		sulfuric acid [Note: Degrades	10119]		
79-34 5 Ki8575000	[skin] OSHA 1 pρm			0001			slowly when exposed to air j	[#ioia]		
1702 55	1 ppm = 7 00 mg/m ³	(7 mg/m²) [skin]			Sp Gr(77°F), 1.59 Noncombustible L) Liquid				
Tetrachloroethylene	Perchloreihylene, Perchloroeihylene,	NIOSH Ca	Ca (500 ppm)	Coloriess liquid with a mild.	MW-165.8 BP: 250°F	VP. 14 mm FBZ: -2*F	Strong oxidizers; chemically-active	Char;		
Cl ₂ C=CCl ₂	Perk Perk Tetrachlorethylene	See Appendix A Minimize workplace	• • •	chloroform-like odor	Sol(77°F): 0.02%	UEL: NA	metals such as lithium, bervilium &	CS2 GC/FID		
127 18 4 KX3850000	letternoren yrene	exposure concentra limit number of		0001	0.025 FEP:NA IP:9.32.eV	LEL NA	barium, caustic soda, sodium hydroxide;			
KV3030000		workers exposed OSHA			IF 932 84		potash	nated Hydro-		
1897 74	1 ppm = 6 89 mg/m³	25 ppm (170 mg/m ³)			Sp Gr: 1.62 Noncombustible L	rdniq		carbonsj		
Tetrachioronaphthalene	Halowax*, Nibren wax,	NIOSH/OSHA 2 mg/m ³	Unknown	Coloriess to pale- yellow solid with an aromatic odor	MW: 265.9 BP: 593-680°F	VP. <1 mm FRZ: 360°F	Strong oxidizers	Filter/Bub; none;		
C ₁₀ H ₄ Cl ₄	Seekay wax	[skin]			Sol: Insoluble FI.P(oc): 410°F	UEL: ?	EL: ?	GC/FID;		
1335 88 2 QK3700000					IP. ?			[#\$130]		
					Sp.Gr: 1.59-1.65 Combustible Solid	t				
Tetraethyl lead (as Pb)	Lead tetraethyl, TEL	NIOSH/OSHA 0.075 mg/m ³	40 mg/m)	Colorless liquid (unless dyed red,	MW: 323.5 BP: 228°F	VP: 0.2 mm FRZ: -202°F	Strong oxidizers, sulfuryl chloride,	XAD-2, Pentane		
Pb(C2H2)4	155	[skin]		orange, or blue) with a pleasant,	(Decomposes) Sol: Insoluble	UEL: ?	rust, potassium permanganate	GC/PID;		
78 00 2 TP4550000				sweet odor. [Note: Main usage is in anti-knock additives for	FLP: 200°F IP: 11.10 eV	LELIUM	Note: Decomposes slowly at room temperature and	[#2533]		
1649 56	1 ppm = 13 45 mg/m ³			gasoline.]	Sp Gr 1 65 Class IIIB Combustible Liquid		higher tempera- tures.]			

Per	sonal protection	Recommendations for respirator			Health hazarda				
	nd sankation See Table 3}	selection — maximum concentration for use (NUC) (See Table 4)	Rou	ite Symptoms (See Table 5)	F (80	Target organ: (See Table 5)			
Clothing: Goggles: Wash: Change: Remove: Provide:	Any poss Any poss Immed contam N.R. Immed non-imperv contam Eyewash, quick drench	NIOSH ¥: SCBAF PD,PP/SAF:PD,PP:ASCBA Escape: GMFOV/SCBAE	inh Abs Ing Con	Nau, vomit, abdom pain; tremor fingers; jaun, enlarged tend liver; derm; monocy, kidney damage	Eye: Skin: Breath: Swallow:	Irr immed Soap wash prompt Resp support Medical attention immed	Liver, kidneys, CNS		
Clothing: Goggles: Wash: Change: Remove:	Repeat Reason prob Prompt contam N R. Prompt non-imperv contam	NIOSH V: SCBAF PD,PP/SAF PD,PP:ASCBA Escape: GMFOV/SCBAE	inh ing Con	Irrit eyes, nose, throat; nau; flush face, neck; verti, dizz, inco; head, som; skin eryt; liver damage, [carc]	Eye Skin: Breath: Swallow:	Irr immed Soap wash prompt Resp support Medical attention immed	Liver, kidneys, eyes, upper res; sys, CNS		
Clothing: Goggles: Nash.	Any poss moll/Repeat liq sol Any poss moll/Reason prob liq-sol Prompi contam	NIOSH/OSHA 20 mg/m ³ : SCBAF/SAF §: SCBAF:PD.PP/SAF:PD.PP:ASCBA Escape: GMFOVHIE/SCBAE	Inh Abs Ing Con	Acne-form derm; head, ftg. anor, verti; jaun, liver inj	Eye: Skin: Breath: Swallow:	Irr immed Soap wash immed Resp support Medical attention immed	Liver, skin		

Inh Abs

Ing Con

Insom, lass, anxiety;

disorientation, halu,

coma; eye irrit

tremor, hyper-reflexia, spastic; bradycardia, hypotension, hypothermia, pallor, nau, anor, low-wgt;

psychosis, mania, convuls,

NIOSH/OSHA 0.75 mg/m³- SA/SCBA 1.875 mg/m³- SA/SCBA 3.75 mg/m³- SA/SCFA 3.75 mg/m³- SA/BD,PP 40 mg/m³- SA/BD,PP 5, SCBAF-PD,PP/SAF-PD,PP:ASCBA Escape: GMFOV/SCBAE

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Irr immed

immed

Soap wash immed Resp support Medical attention

CNS, CVS,

kioneys, eyes

Eye: Skin: Breath: Swallow:

Tetraethyl lead (as Pb)

After work if any poss

Any poss >0 1% Reason prob

After work if any poss contam >0.1%

(>0.1%) Quick drench (>0.1%)

Immed non-imperv contam

Immed non-imperv contam molt/ Prompt non imperv contam sol

contam

N.R.

Change

Remove

Clothing. Goggles: Wash:

Change

Remove

Provide

	Chamical name, structure/formula, CAS and RTECS Nos.,	Synonyms, trade names, and conversion	Exposure Nmits (TWA	IDLH	Physical description	Chemical and physical properties		Incompatibilities and reactivities	Measurement method (See Table 1)
	and DOT ID and guide Nos.	lactors	uniess noted otherwise)			MW, BP, SOL FI.P, IP, Sp.Gr, flammability	VP, FAZ UEL, LEL		
	Tin (organic compounds as Sn)	Synonyms vary depending upon the specific organic compound.	NIOSH/OSHA 0.1 mg/m³ [skin]	Unknown	Appearance and odor vary depending upon the specific organic compound.	Properties vary depending upon the specific organic compour	nd.	Strong oxidizers	Filter/ XAD-2, HPLC; AA; III [#5504]
	Tilanium dioxide TiO2	Ruble, Titanium oxide, Titanium peroxide	NIOSH Ca See Appendix A	Ca (N.E.)	White, odorless powder.	MW: 79.9 BP: 4532 to 5432'F	VP. 0 mm (approx) MLT. 3326 to	None reported	Filter, Acid; AA,
1	13463-67-7 XR2275000		OSHA 10 mg/m³			Sol Insoluble FI P NA IP NA	3362 F UEL NA LEL NA		II(3) [#\$385]
,						Sp.Gr. 4 26 Noncombustible	Solid		
	Toluene C ₆ H ₅ CH ₃ 108-88-3 XS5250000	Methyl benzene, Methyl benzol, Phenyl methane, Toluol	NIOSH/OSHA 100 ppm (375 mg/m ³) ST 150 ppm (560 mg/m ³)	2000 ppm	Coloriess liquid with a sweet, pungent, benzene- like odor	MW 92.1 BP: 232*F Sal(61*F): 0 05% FI P: 40*F IP: 8 82 eV	VP(65°F) 20 mm FAZ: 139°F UEL: 7.1% LEL: 1.2%	Strong öxidizers	Char, CS ₂ : GC/FID, III [#1500, Hydro- carbons]
	1294 27	1 ppm = 3.83 mg/m³				Sp Gr: 0 87 Class IB Flamma	ble Liquid		
	Toluene-2,4- diisocyanale	TDI; 2.4 TDI,	NIOSH Ca	Ca [10 ppm]	Coloriess to pale- yellow solid or	MW: 174.2 BP: 484*F	VP(77*F). 0.01 mm	Strong oxidizers, water, acids, bases	Coated glass
	CH ₃ C ₆ H ₃ (NCO) ₂	2,4-Toluene diisocyanate	See Appendix A 0.005 ppm (0.04 mg/m ³)		liquid (above 71°F) with a sharp, pungent	Sol: Insoluble FI.P: 260*F IP: ?	MLT: 71°F UEL 9.5% LEL 0.9%	& amines may cause foam and spatter; alcohols	wool: Methanol, HPLC/UVD,
	584-84-9 CZ6300000		ST 0.02 ppm (0.15 mg/m ³) OSHA		odor.			Note: Reacts slowly with water to form carbon dioxide and	(#2535)
	2078 57	0.005 ppm (0.04 mg/m ³) 1 ppm = 7.24 mg/m ³ ST 0.02 ppm (0.15 mg/m ³				Sp.Gr: 1 22 Class IIIB Combustible Liquid		polyureas.]	

and the second second

Recommendations Health hazards Personal protection for respirator selection -- maximum concentration for use (MUC) (See Table 4) and sanitation (See Table 3) Route Symptoms First aid Target organa (See Table 5) (See Table 5) (See Table 6) NIOSH/OSHA 1 mg/m³: CCROVDM/SA/SCBA 2 5 mg/m³: SA:CF/PAPROVDM 5 mg/m³: CCRFOVHE/SCBAF/SAF/ **Recommendations vary** Eye: Skin: CNS, eyes, liver. inh Head, verti; irrit eyes; irr immed psycho-neurologic dist; sore throat, cough, abdom pain, vomit; urine depending upon the Abs Water flush immed urinary tract, specific compound. Ing Con Breath Resp support skin, blood Medical attention Swallow GMFOVHIE/PAPRTOVHIE/ retention paresis, focal immed SAT CF anes, skin burns; pruritus; 200 mg/m³: SAF.PD.PP § SCBAF PD.PP/SAF-PD.PP ASCBA Escape: GMFOVHIE/SCBAE in animals: hemolysis, hepatic nec Clothing: Goggles: Wash: N.R. NIOSH Inh Slight lung fib. [carc] Breath: Resp support Lunas NR ¥ SCBAF:PD/PP/SAF.PD,PP:ASCBA NR Escape: HiEF/SCBAE NA Change: NR Remove Clothing: NIOSH/OSHA Repeat Fig. weak; conf. euph, Eye: Skin: CNS, liver, inh Irr immed Goggles: Wash: Reason prob 1000 ppm: CCROV*/SA*/PAPROV*/ SCBA* Abs dizz, head; dilated pupils, Scap wash prompt kidneys, skin lac; ner, musc fig, insom; Breath: Prompt wet Ing lac; ner, mus Con pares; derm Resp support 2000 ppm: SA:CF*/SCBAF/SAF/GMFOV §: SCBAF:PD,PP/SAF:PD,PP:ASCBA Escape: GMFOV/SCBAE Change: NR Swallow Medical attention Remove Immed wet (flamm) immed Clothing NIOSH Repeat Eye: Skin: Inb frrit nose, throat; choke, frr immed Resp sys, skin Goggies: Wash V: SCBAF:PD.PP/SAF:PD.PP:ASCBA Any poss Prompt contam ing Con paroxysmal cough; chest pain, retster soreness; nau, Soap wash immed Escape: GMFOV/SCBAE Breath: Resp support Change After work if reason Medical attention vomit, abdom pain; bron Swallow:

spasm, pulm edema; dysp,

asthma, conj. lac. derm,

skin sens; [carc]

immed

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Toluene-2,4-diisocyanate

Remove

Provide:

prob contam

contam

Prompt non-imperv

Eyewash, quick drench

Exposure limits (TWA Physical description Chemical name. Synonyms, trade names, IDLH Chemical and physical properties incompatibilities Measurement structure/formule method and CAS and FITECS Nos. and conversion reactivities (See Table 1) and DOT ID and factors unless noted MW. BP. SOL guide Nos. otherwise) FI.P. IP. Sp.Gr. VP. FRZ UEL, LEL NIOSH Ca [100 ppm] o Toluidine o-Aminotoluene, Coloriess to pale-MW: 107 2 VP. 0.3 mm Strong oxidizers, Si gel; Ethanol; BP: 392°F FRZ: 6°F UEL: ? LEL ? Aminotoluene, Ca yellow liquid with nitric acid Sol. 2% FLP: 185°F CH3C6H NH2 1-Methyl-2-aminobenzene, See Appendix A an aromatic GC/FID; o Methylandine, 2 ppm aniline like odor 10 95-53-4 XU2975000 IP: 7 44 eV 2 Methylaniline. (9 mg/m³) Iskini 1#2002. ortho Toluidine Aromatic OSHA ACGIH Amines A2 5 ppm (22 mg/m³) Sp Gr 1 01 1708 55 1 ppm 4 46 mg/m³ (skin) Class IIIA Combustible Liquid NIOSH/OSHA MW: 266.3 VP(351°F). 127 mm Butyl phosphate, TBP. 125 ppm Coloriess to pale-Tributyl phosphate None reported Filter. BP 552°F 0 2 ppm (2.5 mg/m³) yellow, odorless Diethyl (CH,[CH,],O),PO FRZ 112"F Tributyl ester of liquid (Decomposes) ether phosphoric acid. Sol: 0 6% UEL ? GC'FPD. 126 73 8 TC7700000 FI P(oc) 295"F In Bulyl phosphate LEL ? 11(3) [#S208] 216 Sp Gr 0 98 1 ppm = 11 07 mg/m³ Class IIIB Compustible Liquid 1.1.2 Trichloroethane beta-Trichlorgethane, NIOSH Colorless liquid MW. 133 4 VP: 19 mm Strong oxidizers Char, Ca CS, GC/FID, Vinyl trichloride [500 ppm] with a sweet, 8P: 237"F FRZ -34"F & caustics; Ca CHCI,CH,CI See Appendix A chloroform-like Sol: 0 4% UEL NA chemically active 10 ppm LEL NA odor metals such as 10 79 00 5 (45 mg/m³) IP. 11.00 eV j#1003, aluminum, magnesium KJ3150000 (skin) OSHA powders, sodium & Haloge nated potassium 10 ppm (45 mg/m³) Hydro Sp Gr 1 44 carbons 2831 74 1 ppm = 5 55 mg/m³ Noncompustible Liquid İskinl Trichloroethylene Ethylene trichloride NIOSH **Colorless liquid** MW: 131.4 VP: 58 mm Strong caustics & alkalis, chemically Char, Ca FRZ: 99"F UEL(77*F): Triclene, [1000 ppm] (unless dyed blue) BP: 189"F CS2, GC/FID, Ca CICH=CCI, Trichloroethene See Appendix A with a chloroform-Sol(77'F): active metals such 25 ppm like odor 0.1% 10.5% as barium, lithium. 111 FI.P: 90°F 79-01-6 LEL(77"F). [#1022] sodium, magnesium, KX4550000 OSHA IP 945 eV 8% titanium & beryllium 50 ppm (270 mg/m³) ST 200 ppm Sp.Gr. 1.46 Class IC Flammable Liquid, but burns with difficulty 1710 74 1 ppm = 5 46 mg/m3 (1080 mg/m³)

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Per	rsonal protection	Recommendations for respirator	Health hazards						
and sanitation (See Table 3)		selection — maximum concentration for use (MUC) (See Table 4)		Route Symptoms (See Table 5)		First ald Table 6)	Target organa (See Table 5)		
Clothing: Goggles: Wash: Change. Remove Provide	Any poss Any poss Immed contam N R Immed non imperv contain Eyewash, quick drench	NIOSH V: SCBAF:PD,PP/SAF:PD,PP:ASCBA Escape: GMFOV/SCBAE	Inh Abs Ing Con	Anoxia, head, cyan; weak, dizz, drow; micro hematuria, eye burns, derm, [carc]	Eye. Skin: Breath: Swallow	Irr immed Soap wash immed Resp support Medical attention immed	Blood, kidneys, liver, CVS, skin, eyes		
Clothing Goggles: Wash: Change Remove	Repeat Reason prob Prompt wet N R Prompt non imperviwet	NIOSH/OSHA 2 ppm: SA/SCBA 5 ppm: SA CF 10 ppm: SCBAF/SAF 125 ppm: SAF.PD.PP § SCBAF PD.PP/SAF PD.PP ASCBA Escape: GMFOVHIE.SCBAE	Inh Con Ing	Eyes, resp. skiñ irrit; head, nau	Eye. Skin: Breath: Swallow	Irr immed Soap wash prompt Resp support Medical attention immed	Resp sys, skin. eyes		
Clothing: Goggles Wash: Change. Remove	Repeat Reason prob Prompt contam N R Prompt non imperv contam	NIOSH V SCBAF PD.PP/SAF:PD.PP:ASCBA Escape: GMFOV/SCBAE	inh Abs Ing Con	Irrit nose, eyes; CNS depres, Irver, kidney damage, {carc}	Eye. Skin. Breath. Swallow	Irr immed Soap wash prompt Resp support Medical attention immed	CNS, eyes, nose. liver, kidneys		
Clothing Goggles Wash: Change Remove	Repeat Reason prob Prompt wet N R Prompt non umperv wet	NIOSH ¥: SCBAF PD.PP/SAF.PD.PP.ASCBA Escape: GMFOV/SCBAE	Inh Ing Con	Head, verti, vis dist, tremors, som, nau, vomit, irrit eyes, derm; card arrhy pares; [carc]	Eye Skin Breath: Swallow	frr immed Soap wash prompt Resp support Medical attention immed	Resp sys, hearl, liver, kidneys, CNS, skin		

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Chemical name, structure/formula, CAS and RTECS Nos.	Synonyms, trade names, and conversion	Exposure timits (TWA	IDLH	Physical description	Chemical and prope	id physical artiss	incompatibilities and reactivities	Measurement method (See Table 1)
and DOT ID and guide Nos.	factors	uniess noted otherwise)	_		MW, BP, SOL FI.P, IP, Sp.Gr, flammability	VP, FRZ UEL, LEL	1995111112	(399 10010 .)
Xylenes (or, mr, p isomers) C _b H ₄ (CH ₁).	o: 1,2-Dimethylbenzene; o-Xylol; m. 1,3-Dimethylbenzene; m. Xylol;	NIOSH/OSHA 100 ppm (435 mg/m³) ST 150 ppm (655 mg/m³)	1000 ppm	Coloriess liquids with an aromatic odor [Note: Pure	MW 106 2 BP: 292/269/ 2811 F Sal Insoluble	VP 7/9/9 mm FRZ 13 54.50 f	Strong oxidizers	Char, CS, GC/ED, III [#1501, Aromatic Hydro carbons]
1330-20-7 ZE2100000	p 1.4 Dimethylbenzene. p Xylaj			p xylene is a solid below 56°F }	FIP 63/84/ 81‴F IP 856/856/844 SpGr:088/086/0	UEL 70/70/0 LEL 11/10/11 44.eV 5/0.85		
1307 27	1 ppm - 4 41 mg/m³				Class I8 Flammat Class IC Flammat			ι.
Xylidine	Aminodimethylbenzene, Aminoxylene,	NIOSH/OSHA 2 ppm	150 ppm	Pale yellow to brown liquid with	MW 121 2 BP: 415 439 F	VP <1 mm FRZ ?	Strong axidizers, hypochlorite bleaches	- Sigel
(CH,)2C,H3NH2	Aminoxylene, Dimethylaminobenzene, Dimethylaniline,	2 ppm (10 mg/m³) (skin)		a weak, aromatic. amine-like odor	501 Slight FIP 206°F	UEL ? LEL 10%	Nypochionte Dieacnes	GC/FID,
1300-73-8 2E8575000	Xylidine isomers, Xylidine (mixed o , m , p)	lound		dimie me www	IP ?			[#2002]
1711 55	1 ppm - 5 04 mg/m³				Sp Gr -0.98 Class IIIB Combus	istible Liquid		
YItrium compounds (as Y) Y	Metal: Yttrium metal Synonyms of other com- pounds vary depending upon the specific com- pound.	NiOSH/OSHA 1 mg/m3	NE	Metal. Dark gray to black solid.	MW: 88 9 BP: 5301:F Sol ? FLP: NA	VP: 0 mm (approx) MLT 2732 F UEL: NA	Oxidizers	Filter. Acid, ICP; III
r 7440-65-5 (Metal) ZG2980000 (Metal)					IP NA	LEL NA		[#7300 Elements]
					Sp.Gr. 4.47 Noncombustible Si	šolid in bulk form		
Zinc chloride fume	None	NIOSH/OSHA	4800 mg/m³	White particulate dispersed in air.	MW: 136.3 VP: 0 mm BP: 1350'F (approx)		Potassium	None available
ZnCi,	ST 2 mg/m ³	ST 2 mg/m ³	an a	disharsen m em	BP: 1350'F (approx) Sol(77"F): MLT: 554"F 432% UEL NA	8v3il30ie		
7646 85 7 ZH1400000					FLP: NA IP: NA	LEL NA		
					Sp Gr(77"F), 2.91 Noncombustible Sc			

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Personal protection and sanilation (See Table 3)		Recommendations for respirator selection — maximum concentration for use (MUC) (See Table 4)		ute Symptoms (See Table 5)	Health hazards First aid (See Table 6)		Target organs (See Table 5)
Clothing Goggles Wash Change Remove	Repeat Reason prob Prompt contain N R Immed wet (flamm)	NIOSH OSHA 1000 ppm CCROV* PAPROV* SA* SCBA* § SCBAF PD PP SAF PD PP ASCBA Escape GMEOV SCBAE	inh Abs Ing Con	Dizz, excitement, drow, inco, staggering gai, irrit eyes, nose, throat; com-ai vacuolization; anor nau, vomit, abdoin pain, derm	Eye Skin Breath Swallow	Irr immed Soap wash prompt Resp support Medic at attention immed	CNS eyes, GI fract blood liver kidneys skin
Clothing Goggles Wash Change Remove Provide	Any poss Any poss Initied contain N H Immed non impervicantion Eyewash quick drench	NIOSH OSHA 20 ppm CCROV:SA SCBA 50 ppm SA CF PAPROV 100 ppm CCRFOV PAPRTOV GMFOV SCBAF/SAF 150 ppm SA PD.PP § SCBAF PD.PP SAF PD PP ASCBA Escape GMFOV/SCBAE	Inh Abs Ing Con	Anoxia cyan lung liver, kidney damage	Eye Skiri Breath Swallow	Irr immed Soap wash immed Resp support Medik at attention immed	Bluod lungs liver kidneys CVS
Clothing Goggles Wash Change Remove Provide	N A Any poss N A N A N A t yewash	NIOSH OSHA 5 mg/m ³ DM 10 mg/m ³ DMXSO SA.SCBA 25 mg/m ³ PAPRDM.SA.CF 50 mg/m ³ HIEF/PAPRTHIE SAT.CF SCBAF/SAF 500 mg/m ³ SA.PD.PP § SCBAF PD.PP/SAF PD.PP § SCBAF PD.PP/SAF PD.PP SCBAF PD.PP/SAF PD.PP	Inh Ing Con	Irrit eyes. In animals pulm irrit, eye inj. possible liver damage	Eye Skin Breath Swallow	Irr immed Soap wash prompt Resp support Medical attention immed	Eyes lungs
Clothing Goggles Wash Change Hemove	N R N R N R N R N R	NIOSH OSHA 10 nig m ³ DMFu' SA' SCBA' 25 nig m ³ PAPRDMFu' SA CF' 50 mg/m ³ HEF/PAPRTHIE' SCBAF SAF 2000 mg/m ³ SAF PD PP § SCBAF PD PP/SAF PD PP ASCBA Escape HiEL SCBAF	Con	Conj. urit nose, throat, cough, copious sputum, dysp, chest pain, pulm edema, broncopneu, pulm fib, cor pulmonate, fever, cyan, tachypnea, burn skin, rind skin, eyes	Breath	Resp support	Resp sys skin eyes

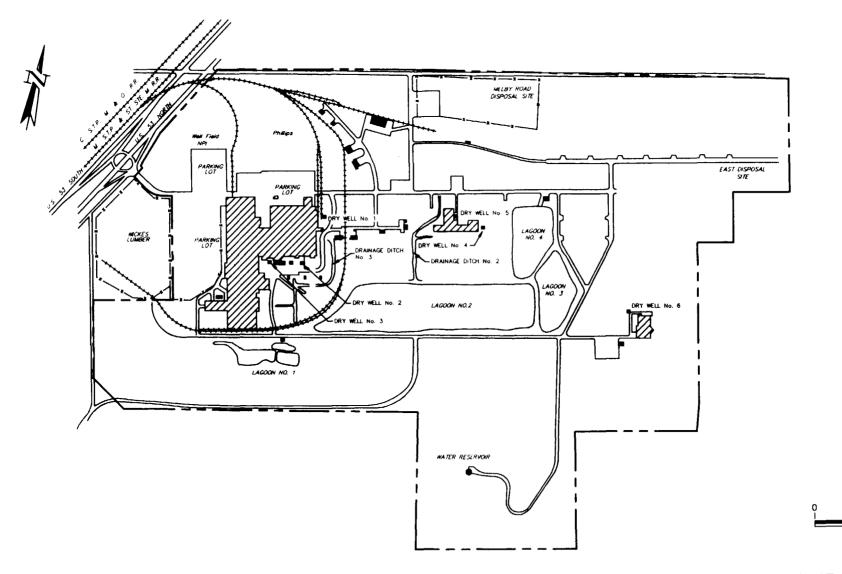
Zinc chloride fume

ATTACHMENT E

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SITE MAP

LLV3760



eder associates

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FIGURE 1

SITE PLAN NATIONAL PRESTO INDUSTRIES, INC. SITE EAU CLAIRE, WISCONSIN

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500'