# SAFETY DATA SHEET



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# **PECORA P-225 PRIMER**

#### 1. PRODUCT IDENTIFICATION

## **IDENTIFICATION of the SUBSTANCE or PREPARATION**

TRADE NAME (AS LABELED):	PECORA P-225 Primer
PRODUCT DESCRIPTION:	Primer
CHEMICAL NAME/CLASS:	Aliphatic Solvent Polymer Mixture
SYNONYMS:	None

#### COMPANY/UNDERTAKING IDENTIFICATION:

SUPPLIER/MANUFACTURER'S NAME:	Pecora Corporation
ADDRESS:	165 Wambold Road, Harleysville, PA 19438
EMERGENCY PHONE:	800-424-9300 (CHEMTREC, 24-hours)
BUSINESS PHONE:	215-723-6051 (Mon–Fri, 8 AM–5 PM ET)

PREPARATION DATE:	April 2009
REVISION DATE:	May 10, 2017

This product is sold for commercial use. This SDS has been developed to address safety concerns of those individuals working with bulk quantities of this material, as well as those of potential users of this product in industrial/occupational settings. ALL United States Occupational Safety and Health Administration Standard (29 CFR 1910.1200), U.S. State equivalent Standards, Canadian WHMIS 2015 and the Global Harmonization required information is included in appropriate sections based on the Global Harmonization Standard format. This product has been classified in accordance with the hazard criteria of the countries listed above and the SDS contains all the information required by the Canadian WHMIS 2015 [HPR-GHIS], the Global Harmonization Standard and OSHA 1910.120.

## 2. HAZARD IDENTIFICATION

GLOBAL HARMONIZATION LABELING AND CLASSIFICATION: Classified in accordance with Global Harmonization Standard under U.S. OSHA Hazard Communication Standard, Canadian WHMIS HPR-GHS 2015.

Classification: Flammable Liquid Cat. 3, Acute Inhalation Toxicity Cat. 3, Acute Dermal Toxicity Cat. 4, Acute Oral Toxicity Cat. 4, Eye Irritation Cat. 2A, Skin Irritation Cat. 2, STOT (Inhalation-Respiratory Irritation) Cat. 3, Respiratory Sensitization Cat. 1, Skin Sensitization Cat. 1

Signal Word: Danger

Hazard Statement Codes: H226, H331, H312, H303, H315, H319, H335, H334, H317

Precautionary Statement Codes: P210, P233, P240, P241, P242, P243, P261, P264, P271, P272, P280, P284, P303 + P361 + P353, P370 + P378, P304 + P340, P342 + P311, P321, P302 + P352, P333 + P313, P362 + P364, P305 + P351 + P338, P321, P403 + P235 + P233, P405, P501

Hazard Symbols/Pictograms: GHS02, GHS07, GHS08







## **EMERGENCY OVERVIEW:**

Physical Description: This product is a colorless, highly flammable liquid with a sweet, aromatic odor.

Health Hazards: DANGER! Flammable liquid. This product may cause respiratory, skin and eye irritation. Eye irritation may severe, depending on duration and concentration of exposure. Harmful or fatal if swallowed. May cause toxic systemic effects by skin absorption. Exposure may cause adverse central nervous system effects. Can cause skin and respiratory sensitization and allergic reaction in susceptible individuals. Contains compound that is a suspect teratogen and carcinogen.

Flammability Hazard: This product is flammable and can ignite if exposed to high temperature or direct flame.

<u>Reactivity Hazard</u>: This product may have some sensitivity to water. Heating of closed containers may cause violent rupture. Closed containers may develop pressure and rupture on prolonged exposure to heat or if contaminated with water.

Environmental Hazard: This product has not been tested for environmental impact. All release to the environment should be avoided.

# HAZARDOUS MATERIALS IDENTIFICATION SYSTEM (HMIS®)

Health	2*	See Section 16 for definitions of rat			
Flammability	3	0 = Minimal 1 = Slight	3 = Serious 4 = Severe		
Physical Hazard	1	1 = Slight 2 = Moderate	* = Chronic		

HMIS® is a registered trademark of the National Paint and Coatings Association.

<u>CANADIAN WHMIS (HPR-GHS) 2015 CLASSIFICATION AND SYMBOLS</u>: See Section 16 for in Classification and Symbols under HPR-GHS 2015.

<u>U.S. OSHA REGULATORY STATUS</u>: This material has a classification under the Global Harmonization Standard, as applied under OSHA regulations, as given earlier in this Section.

## 3. COMPOSITION AND INFORMATION ON INGREDIENTS

Chemical Name	CAS#	W/W%	LABEL ELEMENTS GHS Classification under U.S. OSHA Hazard Communication Standard & Canadian WHMIS (HPR-GHS) 2015 Hazard Statement Codes	
Xylene	1330-20-7	40-60	HARMONISED CLASSIFICATION - ANNEX VI OF REGULATION (EC) NO 1272/2008 (CLP REGULATION)  Classification: Flammable Liquid Cat. 2, Acute Dermal Toxicity Cat. 4, Acute Inhalation Toxicity Cat. 4, Skin Irritation Cat. 2  Hazard Statement Codes: H225, H312 + H332, H315  ADDITIONAL SELF-CLASSIFICATION  Classification: Acute Oral Toxicity Cat. 5, Eye Irritation Cat. 2A  Hazard Statement Codes: H303, H319	
Dicyclohexylmethane-4,4'- Diisocyanate	5124-30-1	10-20	HARMONISED CLASSIFICATION - ANNEX VI OF REGULATION (EC) NO 1272/2008 (CLP REGULATION) Classification: Acute Inhalation Toxicity Cat. 3, Skin Irritation Cat. 2, Eye Irritation Cat. 2A, Skin Sensitization Cat. 1B, Respiratory Sensitization Cat. 1B, STOT (Inhalation-Respiratory Irritation) SE Cat. 3 Hazard Statement Codes: H331, H315, H319, H317, H334, H335	
Proprietary Butylamine Cross-Linker		10-20	NOTIFIED CLASSIFICATION Classification: Acute Dermal Toxicity Cat. 4, Skin Corrosion Cat. 1B, Skin Sensitization Cat. 1A, Aquatic Acute Toxicity Cat. 2, Aquatic Chronic Cat. 2 Hazard Statement Codes: H312, H314, H317, H401, H411	
Proprietary Propoxylated Polymer	Proprietary Propoxylated Polymer		Classification: Not Applicable	
Proprietary Polyether Diol		2-5	Classification: Not Applicable	
Proprietary Amine Cross-Linker		0.1-0.9	NOTIFIED CLASSIFICATION Classification: Acute Dermal Toxicity Cat. 4, Skin Corrosion Cat. 1B, Skin Sensitization Cat. 1A, Aquatic Acute Toxicity Cat. 2, Aquatic Chronic Cat. 2 Hazard Statement Codes: H312, H314, H317, H401, H411	
Other trace components Balance Classification: Not Applicable		Balance	Classification: Not Applicable	
The specific chemical identity and/or e	exact percentage (co	oncentration) o	f composition has been withheld as a trade secret.	

# 4. FIRST-AID MEASURES

<u>PROTECTION OF FIRST AID RESPONDERS</u>: Rescuers should not attempt to retrieve victims of exposure to this material without adequate personal protective equipment. Rescuers should be taken for medical attention, if necessary.

<u>DESCRIPTION OF FIRST AID MEASURES</u>: Remove victim(s) to fresh air, as quickly as possible. Only trained personnel should administer supplemental oxygen and/or cardio-pulmonary resuscitation, if necessary. Remove and isolate contaminated clothing and shoes. Seek immediate medical attention. Take copy of label and SDS to physician or other health professional with victim(s).

Inhalation: If aerosols of this material are inhaled, remove victim to fresh air. If necessary, use artificial respiration to support vital functions.

Skin Exposure: If the material contaminates the skin, immediately begin decontamination with running water. Minimum flushing is for 20 minutes. Do not interrupt flushing. Remove exposed or contaminated clothing, taking care not to contaminate eyes. Victim must seek immediate medical attention.

Eye Exposure: If this product enters the eyes, open victim's eyes while under gently running water. Use sufficient force to open eyelids. Have victim "roll" eyes. Minimum flushing is for 20 minutes. Do not interrupt flushing.

<u>Ingestion</u>: If this material is swallowed, CALL PHYSICIAN OR POISON CONTROL CENTER FOR MOST CURRENT INFORMATION. DO NOT INDUCE VOMITING, unless directly by medical personnel. Have victim rinse mouth with water or give several cupfuls of water, if conscious. Never induce vomiting or give diluents (milk or water) to someone who is <u>unconscious</u>, having convulsions, or <u>unable to swallow</u>. If vomiting occurs, lean patient forward or place on left side (head-down position, if possible) to maintain an open airway and prevent aspiration.

<u>MEDICAL CONDITIONS AGGRAVATED BY EXPOSURE</u>: Acute or chronic respiratory conditions, skin and central nervous system conditions or skin problems may be aggravated by exposure to this product.

<u>INDICATION OF IMMEDIATE MEDICAL ATTENTION AND SPECIAL TREATMENT IF NEEDED</u>: Treat symptoms and eliminate exposure.

# 5. FIRE-FIGHTING MEASURES

<u>FLASH POINT</u>: 23.3°C (74°F)

AUTOIGNITION: Not known for product. For Xylenes: 464°C (867°F)

FLAMMABLE LIMITS IN AIR: Not known for product.

For Xylenes: LEL: 1.0% UEL: 7.0%

**EXTINGUISHING MEDIA:** 

<u>Suitable Extinguishing Media</u>: Use materials appropriate for surrounding materials. May react vigorously with water at high temperatures. Closed containers may rupture violently when heated.

Unsuitable Extinguishing Media: Water.

# PROTECTION OF FIREFIGHTERS:

Special Hazards Arising From the Product: This is a highly flammable liquid. Not sensitive to mechanical impact under normal conditions. Vapors may form explosive mixtures in air. Vapors are heavier than air and can accumulate in confined spaces creating a toxicity and explosion hazard. Vapors can travel long distances and flashback to ignition source. Closed containers may develop pressure and rupture in event of fire or if contaminated with water.

Special Protective Actions for Fire-Fighters: Incipient fire responders should wear eye protection. Structural firefighters must wear Self-Contained Breathing Apparatus and full protective equipment. Move containers from fire area if it can be done without risk to personnel. If possible, prevent runoff water from entering storm drains, bodies of water, or other environmentally sensitive areas.

NFPA RATING
FLAMMABILITY

3
1
INSTABILITY

OTHER
See Section 16 for
Definitions of Ratings

# 6. ACCIDENTAL RELEASE MEASURES

<u>PERSONAL PRECAUTIONS AND EMERGENCY PROCEDURES</u>: An accidental release can result in a fire. Uncontrolled releases should be responded to by trained personnel using pre-planned procedures. Proper protective equipment should be used. Eliminate any possible sources of ignition, and provide maximum explosion-proof ventilation.

# 6. ACCIDENTAL RELEASE MEASURES (Continued)

PERSONAL PRECAUTIONS AND EMERGENCY PROCEDURES: An accidental release may result in a fire. Uncontrolled releases should be responded to by trained personnel using pre-planned procedures. Proper protective equipment should be used. Eliminate any possible sources of ignition, and provide maximum explosion-proof ventilation. Use only non-sparking tools and equipment during the response. The atmosphere must at least 19.5 percent Oxygen before non-emergency personnel can be allowed in the area without Self-Contained Breathing Apparatus and fire protection. Avoid contact with water.

<u>PERSONAL PROTECTIVE EQUIPMENT</u>: Responders should wear the level of protection appropriate to the type of chemical released, the amount of the material spilled, and the location where the incident has occurred.

<u>Small Spills</u>: For releases of 1 drum or less, Level D Protective Equipment (gloves, chemical resistant apron, boots, and eye protection) should be worn.

<u>Large Spills</u>: Minimum Personal Protective Equipment should be rubber gloves, rubber boots, face shield, and Tyvek suit. Minimum level of personal protective equipment for releases in which the level of oxygen is less than 19.5% or is unknown must be **Level B: triple-gloves** (rubber gloves and nitrile gloves over latex gloves), chemical resistant suit, fire-retardant clothing and boots, hard hat, and Self-Contained Breathing Apparatus.

# METHODS FOR CLEAN-UP AND CONTAINMENT:

All Spills: Access to the spill area should be restricted. Spread should be limited by gently covering the spill with polypads. Absorb spilled liquid with clay, sand, polypads, or other suitable inert absorbent materials. All contaminated absorbents and other materials should be placed in an appropriate container and seal. Do not mix with wastes from other materials. Dispose of in accordance with applicable Federal, State, and local procedures (see Section 13, Disposal Considerations). Dispose of recovered material and report spill per regulatory requirements. Remove all residue before decontamination of spill area. Clean spill area with soap and copious amounts of water. Monitor area for combustible vapor levels and confirm levels are below exposure limits given in Section 8 (Exposure Controls-Personal Protection), if applicable, and that levels are below applicable LELs (see Section 5 – Fire Fighting Measures) before non-response personnel are allowed into the spill area. Purge equipment with inert gas prior to reuse.

ENVIRONMENTAL PRECAUTIONS: Minimize use of water to prevent environmental contamination. Prevent spill or rinsate from contaminating storm drains, sewers, soil or groundwater. Place all spill residues in a suitable container and seal. Do not discharge effluent containing this product into streams, ponds, estuaries, oceans or other waters unless in accordance with the requirements of a National Pollutant Discharge Elimination System (NPDES) permit and the permitting authority has been notified in writing prior to discharge. Do not discharge effluent containing this product to sewer systems without previously notifying the local sewage treatment plant authority. For guidance, contact your State Water Board or Regional Office of the EPA.

<u>OTHER INFORMATION</u>: U.S. regulations may require reporting of spills of this material that reach surface waters if a sheen is formed. If necessary, the toll-free phone number for the US Coast Guard National Response Center is 1-800-424-8802.

<u>REFERENCE TO OTHER SECTIONS</u>: See information in Section 8 (Exposure Controls – Personal Protection) and Section 13 (Disposal Considerations) for additional information.

# 7. HANDLING and STORAGE

PRECAUTIONS FOR SAFE HANDLING: As with all chemicals, avoid getting this product ON YOU or IN YOU. Wash thoroughly after handling this product. Do not eat or drink while handling this material. Avoid contact with eyes, skin, and clothing. Avoid breathing fumes, vapors or mist. Do not taste or swallow. Use only with adequate ventilation. Wash hands after handling this product. Contaminated clothing needs to be laundered prior to reuse. Keep away from heat and flame. In the event of a spill, follow practices indicated in Section 6: ACCIDENTAL RELEASE MEASURES. Empty containers may contain residual product; therefore, empty containers should be handled with care. Decontaminate empty containers by filling with water or a solution of ammonium hydroxide (0-10%), detergent (2-5%), isopropanol (0-20%: may create a fire or vapor hazard in some situations, e.g. confined spaces; if so, do not use), water (balance of solution). Heat and CO2 gas are released when isocyanates reacts with water or solution. Let stand uncovered or loosely covered for at least 24 hours. Decontaminate (using above solution) and clean isocyanate handling equipment after use. Stand upwind of all opening, pouring and mixing operations. Keeping work areas clean is essential. Use work surfaces that can be easily decontaminated. Maintain good personal hygiene.

CONDITIONS FOR SAFE STORAGE: Keep container tightly closed when not in use. Store containers in a cool, dry location, away from direct sunlight, sources of intense heat, or where freezing is possible. Material should be stored in secondary containers or in a diked area, as appropriate. Inspect all incoming containers before storage, to ensure containers are properly labeled and not damaged. Containers should be separated from oxidizing materials by a minimum distance of 20 ft. or by a barrier of non-combustible material at least 5 ft. high having a fire-resistance rating of at least 0.5 hours. Storage areas should be made of fire resistant materials. **Local Fire Departments should be notified of the storage of this product on site. Storage and processing areas of this product should be identified with a NFPA 704 placard (diamond) large enough to be seen from a distance.** Post warning and "NO SMOKING" signs in storage and use areas, as appropriate. Refer to NFPA 30, *Flammable and Combustible Liquids Code*, for additional information on storage. Have appropriate extinguishing equipment in the storage area (such as sprinkler systems or portable fire extinguishers). Inspect all incoming containers before storage to ensure containers are properly labeled and not damaged. Empty containers may contain residual product; therefore, empty containers should be handled with care. Store container below 27°C (80°F) to avoid possible reactions related to heat and overpressure of containers.

PRODUCT USE: This product is used as a primer. Follow all industry standards for use of this product.

## 8. EXPOSURE CONTROLS - PERSONAL PROTECTION

#### EXPOSURE LIMITS/CONTROL PARAMETERS:

Ventilation and Engineering Controls: Use with adequate, explosion proof ventilation to ensure exposure levels are maintained below the limits provided above.

Occupational/Workplace Exposure Limits/Guidelines:

<u>Chemical Name</u>	al Name CAS#		<u>Value</u>
Proprietary Butylamine Cross-Linker		NE	NE
Dicyclohexylmethane-4,4'-diisocyanate 5124-30-1		ACGIH TLV TWA NIOSH REL STEL DFG MAK	0.054 mg/m³ 0.01 mg/m³ (ceiling) Danger of Skin Sensitization
Proprietary Polyether Diol		AIHA WEEL TWA	10 mg/m <sup>3</sup>
Proprietary Amine Cross-Linker Exposure limits given are for diethylenetriamine		ACGIH TLV TWA NIOSH REL TWA DFG MAK	4.2 mg/m3 (skin) 4 mg/m3 (skin) Danger of sensitization of the Skin
Proprietary Propoxylated Polymer		NE	NE
Xylene 1330-20-7		ACGIH TLV TWA ACGIH TLV STEL OSHA PEL TWA OSHA PEL STEL NIOSH REL TWA NIOSH REL STEL DFG MAK TWA DFG MAK PEAK	100 ppm 150 ppm 100 ppm 150 ppm (vacated 1989 PEL) 100 ppm 150 150 100 (skin) 4•MAK 15 minute average value, 1-hr interval 4 per shift

NE = Not Established. See Section 16 for Definitions of Terms Used.

Biological Exposure Indices (BEIs): Currently, the following BEI's have been established for the Xylene component.

CHEMICAL: DETERMINANT	SAMPLING TIME	BEI
Xylenes  ◆ Methylhippuric Acid in Urine	• End of Shift	• 1.5 g/g Creatinine

PERSONAL PROTECTIVE EQUIPMENT (PPE): The following information on appropriate Personal Protective Equipment is provided to assist employers in complying with OSHA regulations found in 29 CFR Subpart I (beginning at 1910.132, including the Respiratory Protection Standard (29 CFR 1910.134), Eye Protection Standard 29 CFR 1910.13, the Hand Protection Standard 29 CFR 1910.138, and the Foot Protection Standard 29 CFR 1910.136), equivalent standards of Canada (including the Canadian CSA Respiratory Standard Z94.4-93-02, the CSA Eye Protection Standard Z94.3-M1982, Industrial Eye and Face Protectors and the Canadian CSA Foot Protection Standard Z195-M1984, Protective Footwear). Please reference applicable regulations and standards

Eye/Face Protection: Use approved safety goggles or safety glasses. If necessary, refer to appropriate regulations.

Skin Protection: Wear chemical impervious gloves (e.g., Nitrile or Neoprene). Use triple gloves for spill response. If necessary, refer to appropriate regulations.

Body Protection: Use body protection appropriate for task (e.g., lab coat, coveralls, Tyvek suit). If necessary, refer to the OSHA Technical Manual (Section VII: Personal Protective Equipment) or appropriate Standards of Canada. If a hazard of injury to the feet exists due to falling objects, rolling objects, where objects may pierce the soles of the feet or where employee's feet may be exposed to electrical hazards, use foot protection, as described in appropriate regulations.

Respiratory Protection: If mists or sprays from this product are created during use, use appropriate respiratory protection. If necessary, use only respiratory protection authorized in appropriate regulations. Oxygen levels below 19.5% are considered IDLH by OSHA. In such atmospheres, use of a full-facepiece pressure/demand SCBA or a full facepiece, supplied air respirator with auxiliary self-contained air supply is required under appropriate regulations. The following are NIOSH respiratory equipment guidelines are presented for additional assistance in respiratory protective equipment selection.

## **DICYCLOMETHANE-4,4'-DIISOCYANATE**

RESPIRATORY PROTECTION **CONCENTRATION** Up to 0.1 ppm: Supplied-Air Respirator (SAR)

Up to 0.25 ppm: SAR operated in a continuous-flow mode.

Full-facepiece Self-Contained Breathing Apparatus (SCBA); or full-facepiece SAR. Up to 0.5 ppm:

Up to 1 ppm: Positive pressure, full-facepiece SAR.

Emergency or Planned Entry into Unknown Concentrations or IDLH Conditions: Positive pressure, full-facepiece SCBA; or positive pressure, full-facepiece SAR

with an auxiliary positive pressure SCBA.

Gas mask with organic vapor canister; or escape-type SCBA. Escape:

**XYLENES** 

RESPIRATORY PROTECTION

CONCENTRATION

Up to 900 ppm: Any Chemical Cartridge Respirator with organic vapor cartridge(s), or any Powered, Air-Purifying Respirator (PAPR) with organic

vapor cartridge(s), or any Supplied-Air Respirator (SAR), or any Self-Contained Breathing Apparatus (SCBA) with a full facepiece.

Emergency or Planned Entry into Unknown Concentrations or IDLH Conditions: Any SCBA that has a full facepiece and is operated in a pressure-demand or other

positive-pressure mode, or any SAR that has a full facepiece and is operated in a pressure-demand or other positive-pressure mode in

combination with an auxiliary SCBA operated in pressure-demand or other positive-pressure mode.

Any Air-Purifying, Full-Facepiece Respirator (gas mask) with a chin-style, front- or back-mounted organic vapor canister, or any Escape:

## 9. PHYSICAL and CHEMICAL PROPERTIES

FORM: Somewhat viscous liquid.

MOLECULAR WEIGHT: Mixture.

ODOR: Sweet, aromatic.

VAPOR DENSITY: (air = 1) > 1

COLOR: Clear, colorless.

MOLECULAR FORMULA: Mixture.

ODOR THRESHOLD: For Xylenes: 20 ppm (detection); 40 ppm (recognition).

BOILING POINT: Not available.

# 9. PHYSICAL and CHEMICAL PROPERTIES (Continued)

SPECIFIC GRAVITY (water = 1): 0.94 pH: Not available.

SOLUBILITY IN WATER: Not available. EVAPORATION RATE (nBuAc = 1): Not available.

<u>VAPOR PRESSURE</u>: Not available. <u>VOC</u>: 625 g/L

FLASH POINT: 23.3°C (74°F) PERCENT VOLATILE BY VOLUME: ~70%

COEFFICIENT WATER/OIL DISTRIBUTION: Not available. VISCOSITY: Not available.

<u>HOW TO DETECT THIS SUBSTANCE (WARNING PROPERTIES)</u>: The odor of this product may act as warning properties in the event of an accidental release. The odor of Xylenes in this product may be a good warning property as its TLV is more than 10 times the odor threshold.

# 10. STABILITY and REACTIVITY

CHEMICAL STABILITY: Stable under normal circumstances of use and handling.

CONDITIONS TO AVOID: Avoid contact with incompatible chemicals and exposure to extreme temperatures.

<u>INCOMPATIBLE MATERIALS</u>: This product is not compatible with oxidizers, water, alcohols, acids, bases, amines, amides, phenols, mercaptans, urethanes, ureas and surface active materials, metal compounds (e.g. organotin catalysts) and 1,3-dichloro-5,5-dimethyl-2,4-imidazolidindione (dichlorohydrantoin). Due to the isocyanate material in this product, it may attack copper and copper alloys, such as brass and bronze, tin and zinc.

<u>HAZARDOUS DECOMPOSITION PRODUCTS</u>: <u>Combustion</u>: Thermal decomposition of this product can generate carbon, nitrogen and silicon and tin oxides, hydrogen, reactive hydrocarbons, low molecular aldehydes, propylene glycol, furan, dioxalane and formaldehyde. <u>Hydrolysis</u>: Carbon dioxide, methane and non-toxic, insoluble polyureas.

<u>POSSIBILITY OF HAZARDOUS REACTIONS</u>: This product may undergo uncontrolled exothermic polymerization upon contact with incompatible materials, especially strong bases, such as triethylamine and sodium hydroxide, trialkyl phosphines, potassium acetate, many metal compounds soluble in organic media, or if heated. The resulting pressure build-up could rupture closed containers.

## 11. TOXICOLOGICAL INFORMATION

<u>POTENTIAL HEALTH EFFECTS</u>: The most significant routes of occupational exposure are inhalation and contact with skin and eyes. The symptoms of exposure to this product are as follows:

Contact with Skin or Eyes: Depending on the duration of skin contact, skin exposure can cause reddening, discomfort or irritation. Prolonged contact can cause corrosive effects with severe inflammation, redness, rash, swelling, blistering, tissue death and permanent scarring. Isocyanates, in general, can cause skin discoloration (staining) and hardening of the skin after repeated exposure. The Dicyclohexylmethane-4,4'-Diisocyanate is a very strong sensitizing agent in humans; other components are also considered as skin sensitizers. Skin sensitization may occur after only one contact with. Brief contact with the liquid or vapors from this product and the eyes can cause irritation, reddening and watering. Permanent eye injury, including blindness, could result from direct contact with the liquid.

Skin Absorption: Prolonged skin contact may cause adverse systemic toxicity by skin absorption as described under ingestion or inhalation.

<u>Ingestion</u>: If the product is swallowed, it can irritate the mouth, throat, and other tissues of the gastro-intestinal system and may cause nausea, vomiting, and diarrhea as well as adverse effects on the central nervous system. Symptoms can include dizziness, vomiting and incoordination. Ingestion of large amounts may be harmful and cause systemic toxicity. Aspiration into the lungs after ingestion can pose a serious hazard of chemical and pulmonary edema. Ingestion of large amount may be fatal.

Inhalation: Inhalation of vapors, mists, or sprays of this product can moderately to can irritate the tissues of the nose, mouth, throat, and upper respiratory system. Symptoms of exposure may include coughing, sneezing, and difficulty breathing. Coughing with chest pain or tightness may also occur, frequently at night. These symptoms may occur during exposure or may be delayed several hours. Inhalation of high concentrations of this product (as may occur in a poorly ventilated area) may be fatal. Inhalation can also lead to adverse central nervous system effects, including dizziness, incoordination, nausea and vomiting. High aerosol concentrations could cause inflammation of the lungs (chemical pneumonitis), chemical bronchitis with severe asthma-like wheezing, severe coughing spasms and accumulation of fluid in the lungs (pulmonary edema), which could prove fatal. Symptoms of pulmonary edema may not appear until several hours after exposure and are aggravated by physical exertion. Respiratory sensitization and lung damage may be permanent.

<u>Injection</u>: Accidental injection of this product (e.g. puncture with a contaminated object) may cause burning, redness, and swelling in addition to the wound.

Target Organs: Acute: Skin, eyes, respiratory systems. Chronic: Skin, respiratory and central nervous systems.

<u>TOXICITY DATA</u>: There are currently no toxicity data available for this product; the following toxicology data are available for components greater than 1% in concentration. Due to the large amount of data, only human data, LD50 Oral-Rat or Mouse, LD50 Skin-Rat or Mouse, LC50 Inhalation-Rat or Mouse and skin irritation data are provided in this SDS. Contact Pecora for more information.

#### PROPRIETARY BUTYLAMINE CROSS-LINKER:

 $LD_{50}$  (Oral-Rat) 13,500  $\mu L/kg$ : Gastrointestinal: changes in structure or function of salivary glands, hypermotility, diarrhea, other changes

LD<sub>50</sub> (Skin-Rabbit) 16 mL/kg: Lungs, Thorax, or Respiration: other changes; Liver: other changes; Skin and Appendages: dermatitis, other (after systemic exposure)

#### DICYCLOHEXYLMETHANE-4,4'-DIISOCYANATE:

Standard Draize Test (Skin-Rabbit) 500 µL/24 hours: Moderate

Standard Draize Test (Eye-Rabbit) 100 µL: Mild

Standard Draize Test (Eye-Rabbit) 100 µL/24 hours: Severe

LD<sub>50</sub> (Oral-Rat) 9900 mg/kg: Behavioral: food intake (animal); Gastrointestinal: hypermotility, diarrhea; Liver: other changes

# PROPRIETARY POLYETHER DIOL:

 $\begin{array}{l} LD_{50} \; (Oral\text{-Rat}) \sim 4000 \; mg/kg \\ LD_{50} \; (Skin\text{-Rabbit}) \sim 2000 \; mg/kg \end{array}$ 

LC<sub>50</sub> (Inhalation-Rat) ~ 200 mg/L/1 hour

XYLENE:

Standard Draize Test (Eye-Human) 200 ppm

#### XYLENE (continued):

LDLo (Oral-Human) 50 mg/kg

LCLo (Inhalation-Human) 10,000 ppm/6 hours: Behavioral: general anesthetic; Lungs, Thorax, or Respiration: cyanosis; Blood: other changes

TCLo (Inhalation-Human) 200 ppm: Sense Organs and Special Senses (Olfaction): effect, not otherwise specified; Sense Organs and Special Senses (Eye): conjunctive irritation; Lungs, Thorax, or Respiration: other changes

Standard Draize Test (Skin-Rabbit) 100%: Moderate

Standard Draize Test (Skin-Rabbit) 500 mg/24 hours: Moderate

Standard Draize Test (Eye-Rabbit) 87 mg/L: Mild Standard Draize Test (Eye-Rabbit) 5 mg/24 hours: Severe

Open Irritation Test (Skin-Rat) 60 uL/8 hours: Mild

LC<sub>50</sub> (Inhalation-Rat) 5000 ppm/4 hours

LD<sub>50</sub> (Oral-Rat) 4300 mg/kg: Liver: other changes; Kidney/Ureter/Bladder: other changes

LD<sub>50</sub> (Oral-Mouse) 2119 mg/kg

LD<sub>50</sub> (Skin-Rabbit) > 1700 mg/kg

# 11. TOXICOLOGICAL INFORMATION (Continued)

<u>CARCINOGENIC POTENTIAL</u>: The following table summarizes the carcinogenicity listing for the components of this product. "NO" indicates that the substance is not considered to be or suspected to be a carcinogen by the listed agency, see section 16 for definitions of other ratings.

CHEMICAL	IARC	EPA	NTP	NIOSH	ACGIH	OSHA	PROP 65
Proprietary Amine Cross-Linker	No	No	No	No	No	No	No
Proprietary Butylamine Cross-Linker	No	No	No	No	No	No	No
Dicyclohexylmethane-4,4'-Diisocyanate	No	No	No	No	No	No	No
Proprietary Polyether Diol	No	No	No	No	No	No	No
Proprietary Propoxylated Polymer	No	No	No	No	No	No	No
Xylene	3	I	No	No	A4	No	Yes

IARC-3: Unclassifiable as to Carcinogenicity in Humans. EPA-I (Data are Inadequate for and Assessment of Human Carcinogenic Potential). ACGIH TLV-A4: Not Classifiable as a Human Carcinogen.

<u>IRRITANCY OF PRODUCT</u>: This product is irritating by all routes of exposure. Depending on concentration and duration of exposure, eye exposure may be severe.

SENSITIZATION TO THE PRODUCT: This product may cause skin and respiratory sensitivity and allergic reaction in susceptible individuals. In general, diisocyanates are well known to cause respiratory sensitization and, therefore, Dicyclohexylmethane-4,4'-Diisocyanate is considered a respiratory sensitizer. Diisocyanate respiratory sensitization is usually caused by a very large exposure, or by multiple exposure. Although varying periods of exposure (1 day to years) may elapse before sensitization occurs, it develops more often during the first few months of exposure. Sensitized individuals react to very low levels of airborne diisocyanates that have no effect on non-sensitized people. At first, the symptoms may appear to be a cold or mild hay fever. However, severe asthmatic symptoms can develop and include wheezing, tightness of the chest, shortness of breath, difficulty breathing and/or coughing. Fever, chills, general feelings of discomfort, headache, and fatigue can also occur. Symptoms may occur immediately upon exposure (within an hour), several hours after exposure or both, and/or at night. Typically, the asthma improves with removal from exposure (e.g. weekends or vacations) and returns, in some cases, in the form of an "acute attack", on renewed exposure. Sensitized people who continue to be exposed to diisocyanates may develop symptoms sooner after each exposure. The number and severity of symptoms may increase. Cross-sensitization between different diisocyanates may occur. Exposure to diisocyanates is likely to aggravate individuals with existing respiratory disease, such as chronic bronchitis and emphysema. Respiratory sensitization may be permanent.

Skin sensitization may occur after only a few days working with products containing Dicyclohexylmethane-4,4'-Diisocyanate. Onset of symptoms is usually delayed. Symptoms include a rash on the hands, arms, neck, face, chest or abdomen, even when contact occurs with a small amount of product. Other effects such as coughing, a burning sensation in the throat or redness and swelling of the eyes might also occur.

<u>TOXICOLOGICAL SYNERGISTIC PRODUCTS</u>: There have been several studies in humans and animals on the interaction of Xylenes with drugs, alcohol and other solvents. Xylene has a high potential to interact with other compounds because it increases metabolic enzymes in the liver and decreases metabolic enzymes in the lungs. In general, exposure to related solvents, such as benzene, toluene and ethanol (alcohol) slows the rate of clearance of Xylenes from the body, thus enhancing its toxic effects.

<u>REPRODUCTIVE TOXICITY INFORMATION</u>: This product has not been tested for reproductive toxicity. The following information is available for some components.

<u>Mutagenicity</u>: Either no information is available for components, or negative results from testing have been obtained.

Embryotoxicity/Teratogenicity: Xylene (mixed isomers) are considered fetotoxic in humans, based on observations of reduced fetal weight, delayed ossification and persistent behavioral effects in animal studies in the absence of maternal toxicity. Other developmental effects have been observed in animal studies in the presence of maternal toxicity. Several human population studies have suggested a link between exposure to organic solvents (including xylene) and increased occurrence of miscarriages or birth defects in children. However, in the majority of cases, there was exposure to a variety of solvents at the same time, exposure were ill-defined, and the number of cases examined was small.

Reproductive Toxicity: No information is available.

# 12. ECOLOGICAL INFORMATION

## ALL WORK PRACTICES MUST BE AIMED AT ELIMINATING ENVIRONMENTAL CONTAMINATION.

MOBILITY: This product has not been tested for mobility in soil. The following information is available for components.

XYLENE: Several experimental Koc values for this compound have been reported depending upon the pH and organic carbon content of the soil. Batch experiments conducted with five low organic carbon content (0.04-1.12%), field contaminated soils (3 silty clay and two sandy loams) yielded Koc values ranging from 39-365. This compound in Norwegian forest soil at pH 5.6 and organic carbon content of 0.2 percent has a reported experimental Koc of 129; in Norwegian agricultural soil at pH 7.4 and organic carbon content of 2.2 percent has a reported experimental Koc of 158; in Norwegian forest soil at pH 4.2 and organic carbon content of 3.7 percent has a reported experimental Koc of 289. Based on a recommended classification scheme and the experimentally determined Koc values, this material is expected to have moderate to high mobility in soils. Xylene isomers have been observed to pass through soil at a dune-infiltration site on the Rhine River and to leach into groundwater under a rapid infiltration site.

<u>PERSISTENCE AND BIODEGRADABILITY</u>: This product has not been tested for persistence or biodegradability. The following information is available for components.

XYLENE: Based upon an experimental vapor pressure of 7.99 mm Hg at 25°C, this compound is expected to exist entirely in the vapor phase in the ambient atmosphere. Vapor-phase material is degraded in the atmosphere by reaction with photochemically-produced hydroxyl radicals with an estimated atmospheric lifetime of about 1-2 days. This compound is expected to have moderate to high mobility in soils based upon experimental Koc values obtained with a variety of soils at differing pH values and organic carbon content. Volatilization from moist soil surfaces is expected based on an experimental Henry's Law constant of 7.0X10-3 atm-cu m/mole. Biodegradation is an important environmental fate process for this compound. In water, this compound is expected to adsorb somewhat to sediment or particulate matter based on its measured Koc values. This compound is expected to volatilize from water surfaces given its experimental Henry's Law constant. Estimated half-lives for a model river and model lake are 3 and 99 hours, respectively. Log Kow = 3.5-68.

## 12. ECOLOGICAL INFORMATION (Continued)

<u>BIO-ACCUMULATION POTENTIAL</u>: This product has not been tested for bio-accumulation potential. The following values are available for components.

XYLENE: An experimental BCF value of 20 was measured for all isomers in eels exposed to petroleum for 10 days. According to a classification scheme, this BCF value suggests that bioconcentration in aquatic organisms is low.

<u>ECOTOXICITY</u>: This product has not been tested for aquatic or animal toxicity. All release to terrestrial, atmospheric and aquatic environments should be avoided. The following aquatic toxicity data are available for some components. Only select data are given due to the large amount of data available. Contact Pecora for more information.

DICYCLOHEXYLMETHANE-4,4'-DIISOCYANATE:

LC<sub>50</sub> (Brachydanio rerio Zebra fish) 96 hours = 1.2 mg/L; 91.6% a.i./ /Conditions of bioassay not specified in source examined

EC<sub>50</sub> Activated sludge (test for inhibition of oxygen consumption) 3 hours = 19 mg/L;

XYLENE:

LC50 (rainbow trout) 96 hours = 13.5 mg/L (conditions of bioassay not specified, no specific

isomer)

XYLENE (continued):

 $LD_{50}$  (goldfish) 24 hours = 13 mg/L (conditions of bioassay not specified, no specific isomer)  $LC_{50}$  (fathead minnow) 1 hour = 42 mg/L at 18-22°C, in a static bioassay (No specific isomer)  $LC_{50}$  (fathead minnow) 24-96 hours = 46 mg/L at 18-22°C, in a static bioassay (No specific isomer)

LC<sub>50</sub> (Carassius auratus goldfish) 96 hours = 16.9 ppm (conditions of bioassay not specified, no

specific isomer)

OTHER ADVERSE EFFECTS: This material is not expected to have any ozone depletion potential.

<u>ENVIRONMENTAL EXPOSURE CONTROLS</u>: Controls should be engineered to prevent release to the environment, including procedures to prevent spills, atmospheric release and release to waterways.

#### 13. DISPOSAL CONSIDERATIONS

<u>PREPARING WASTES FOR DISPOSAL</u>: As supplied, this product would be a hazardous waste as defined by U.S. federal regulation (40 CFR 261) if discarded or disposed. It has the characteristic of Ignitibility. State and local regulations may differ from federal regulations. The generator of the waste is responsible for proper waste determination and management.

<u>U.S. EPA WASTE NUMBER</u>: Wastes of this product should be test to see if they meet the criteria of D001 (Ignitability characteristic).

## 14. TRANSPORTATION INFORMATION

<u>U.S. DEPARTMENT OF TRANSPORTATION</u>: This product is classified as Dangerous Goods, per U.S. DOT regulations, under 49 CFR 172.101.

UN Identification Number: UN 1866

Proper Shipping Name: Resin solution, flammable

Hazard Class Number and Description: 3 (Flammable)

Packing Group: PG II

DOT Label(s) Required: Class 3 (Flammable)

North American Emergency Response Guidebook Number (2016): 127

Marine Pollutant: The components of this product not classified by the DOT as a Marine Pollutant (as defined by 49 CFR 172.101.

TRANSPORT CANADA TRANSPORTATION OF DANGEROUS GOODS REGULATIONS: This product is classified as Dangerous

Goods, per regulations of Transport Canada.

UN Identification Number: UN 1866

Proper Shipping Name: Resin solution, flammable

Hazard Class Number and Description: 3 (Flammable)

Packing Group: PG II

Hazard Shipping Label(s) Required: Class 3 (Flammable)

Special Provisions:83Excepted Quantities:E2Explosive Limit & Limited Quantity Index:5 LERAP Index:NonePassenger Carrying Ship Index:NonePassenger Carrying Road Or Rail Vehicle Index:5 L

INTERNATIONAL AIR TRANSPORT ASSOCIATION SHIPPING INFORMATION (IATA): This product is classified as dangerous

goods, per the International Air Transport Association.

UN Identification Number: UN 1866

Proper Shipping Name: Resin solution, flammable

Hazard Class or Division: 3 (Flammable)
Hazard Label(s) Required: Class 3 (Flammable)

Packing Group:

Excepted Quantities:

Passenger and Cargo Aircraft Packing Instruction:

Passenger and Cargo Aircraft Maximum Net Quantity per Pkg.:

Passenger and Cargo Aircraft Limited Quantity Packing Instruction:

Y341

Passenger and Cargo Aircraft Limited Quantity Maximum Net Quantity per Pkg.:

1 L

Cargo Aircraft Only Packing Instruction:
364
Cargo Aircraft Only Maximum Net Quantity per Pkg.:
60 L
Special Provisions:
A3
ERG Code:
3L

# 14. TRANSPORTATION INFORMATION (Continued)

INTERNATIONAL MARITIME ORGANIZATION SHIPPING INFORMATION (IMO): This product is classified as dangerous goods,

per the International Maritime Organization.

UN No.: 1866

Proper Shipping Name: Resin solution, flammable

Hazard Class Number: 3 (Flammable)
Labels: Class 3 (Flammable)

Packing Group:IISpecial Provisions:NoneLimited Quantities:5 LExcepted Quantities:E2

Packing: Instructions: P001; Provisions: PP1
IBCs: Instructions: IBC02; Provisions: None
Tanks: Instructions: T4; Provisions: T1, TP8

EmS: F-E, S-E Stowage Category: Category B.

Segregation:

Marine Pollutant: No component of this product is designated by the IMO to be a Marine Pollutant.

# 15. REGULATORY INFORMATION

## **U.S. REGULATIONS:**

<u>U.S. SARA Reporting Requirements</u>: The following components of this product are subject to the reporting requirements of Sections 302, 304, and 313 of Title III of the Superfund Amendments and Reauthorization Act.

CHEMICAL	SECTION 302 EHS (TPQ)	SECTION 304 RQ	SECTION 313 TRI (threshold)	
	(40 CFR 355, Appendix A)	(40 CFR Table 302.4)	(40 CFR 372.65)	
Dicyclohexylmethane-4,4'-Diisocyanate	No	No	Yes	
Xylene	No	No	Yes	

U.S. SARA Hazard Categories (Section 311/312, 40 CFR 370-21): ACUTE: Yes; CHRONIC: Yes; FIRE: Yes; REACTIVE: No; SUDDEN RELEASE: No

<u>U.S. TSCA Inventory Status</u>: All components of this product are in compliance with the inventory listing requirements of the U.S. Toxic Substances Control Act (TSCA) Chemical Substance Inventory.

U.S. CERCLA Reportable Quantity (RQ): Xylene = 100 lb (45.4 kg)

<u>U.S. Clean Air Act (CA 112r) Threshold Quantity (TQ)</u>: The Xylene component is listed as a Hazardous Air Pollutant (HAP) generally known or suspected to cause serious health problems. The Clean Air Act, as amended in 1990, directs EPA to set standards requiring major sources to sharply reduce routine emissions of toxic pollutants. EPA is required to establish and phase in specific performance based standards for all air emission sources that emit one or more of the listed pollutants. Xylenes are included on this list.

<u>U.S. Clean Water Act Requirements</u>: Xylene (mixed) is designated as a hazardous substance under section 311(b)(2)(A) of the Federal Water Pollution Control Act and further regulated by the Clean Water Act Amendments of 1977 and 1978. These regulations apply to discharges of this substance. This designation includes any isomers and hydrates, as well as any solutions and mixtures containing this substance.

California Safe Drinking Water And Toxic Enforcement Act (Proposition 65): No component is on the California Proposition 65 lists.

## CANADIAN REGULATIONS:

Canadian DSL/NDSL inventory status: The components of this product are on the DSL Inventory.

Canadian Environmental Protection Act (CEPA) Priorities Substances Lists: The Xylene component is on the CEPA Priority Substances 1 list, not considered as "TOXIC" under Section 64 of CEPA.

Canadian WHMIS (HPR-GHS) 2015 Classification and Symbols: See Section 16 in Classification and Symbols under HPR-GHS 2015.

## **MEXICAN REGULATIONS:**

Mexican Workplace Regulations (NOM-018-STPS-2000): This product is not classified as hazardous.

## 16. OTHER INFORMATION

U.S. ANSI STANDARD LABELING (Precautionary Statements): DANGER! FLAMMABLE LIQUID. MAY BE TOXIC IF INHALED. MAY CAUSE EYE, SKIN AND RESPIRATORY IRRITATION; EYE IRRITATION MAY BE SEVERE. VAPORS MAY CAUSE CENTRAL NERVOUS SYSTEM EFFECTS. CHRONIC INHALATION EXPOSURE MAY CAUSE DAMAGE TO LUNGS. CONTAINS COMPOUND THAT IS A SUSPECT CARCINOGEN AND REPRODUCTIVE TOXIN. MAY CAUSE SKIN AND RESPIRATORY SENSITIZATION. Avoid contact with eyes, skin, and clothing. Avoid breathing mist, vapors or fume. Do not taste or swallow. Wash thoroughly after handling. Keep container tightly closed. Use only with adequate ventilation. Keep away from heat and flame. Wear gloves, eye protection, respiratory protection, and appropriate body protection. FIRST-AID: In case of contact, immediately flush skin and eyes with plenty of water. Remove contaminated clothing and shoes. Get medical attention if irritation develops or persists. If inhaled, remove to fresh air. If not breathing, give artificial respiration. If breathing is difficult, give oxygen. If swallowed, do not induce vomiting. Get medical attention. IN CASE OF FIRE: Use water fog, foam, dry chemical, or CO2. IN CASE OF SPILL: Absorb spilled product with polypads or other suitable absorbing material. Place all spill residue in an appropriate container and seal. Dispose of in accordance with U.S. Federal, State, and local hazardous waste disposal regulations and those of Canada.

# **16. OTHER INFORMATION (Continued)**

GLOBAL HARMONIZATION LABELING AND CLASSIFICATION: Classified in accordance with Global Harmonization Standard under U.S. OSHA Hazard Communication Standard, Canadian WHMIS HPR-GHS 2015.

<u>Classification</u>: Flammable Liquid Category 3, Acute Inhalation Toxicity Category 3, Acute Dermal Toxicity Category 4, Acute Oral Toxicity Category 5, Eye Irritation Category 2A, Skin Irritation Category 2, Specific Target Organ Toxicity (Inhalation-Respiratory Irritation) Category 3, Respiratory Sensitization Category 1, Skin Sensitization Category 1

Signal Word: Danger

<u>Hazard Statements</u>: H226: Flammable liquid and vapor. H331: Toxic if inhaled. H312: Harmful in contact with skin. H303: May be harmful if swallowed. H315: Causes skin irritation. H319: Causes serious eye irritation. H335: May cause respiratory irritation. H334: May cause allergy or asthma symptoms or breathing difficulties if inhaled. H317: May cause an allergic skin reaction.

#### **Precautionary Statements:**

Prevention: P210: Keep away from heat/sparks/open flames/hot surfaces. — No smoking. P233: Keep container tightly closed. P240: Ground/bond container and receiving equipment. P241: Use explosion-proof electrical/ventilating/lighting/equipment. P242: Use only non-sparking tools. P243: Take precautionary measures against static discharge. P261: Avoid breathing mists, sprays, fume. P264: Wash contaminated tissues after handling. P270: Do not eat, drink or smoke when using this product. P271: Use only outdoors or in a well-ventilated area. P272: Contaminated work clothing should not be allowed out of the workplace. P280: Wear protective gloves, clothing, eye protection and face protection. P284: Wear respiratory protection.

Response: P303 + P361 + P353: IF ON SKIN (or hair): Remove/Take off immediately all contaminated clothing. Rinse skin with water/shower. P370 + P378: In case of fire: Use materials appropriate for surrounding fire for extinction.

Response (continued): P342 + P311: If experiencing respiratory symptoms: Call a POISON CENTER or doctor/physician. P302 + P352: IF ON SKIN: Wash with plenty of soap and water. P333 + P313: If skin irritation or rash occurs: Get medical advice/attention. P362 + P364: Take off contaminated clothing and wash it before reuse. P305 + P351 + P338: IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. P321: Specific treatment (remove from exposure and treat symptoms).

Storage: P403 + P233 + P235: Store in a well-ventilated place. Keep container tightly closed. Keep cool. P405: Store locked up.

Disposal: P501: Dispose of contents/containers in accordance with all local, regional, national and international regulations.

Hazard Symbols/Pictograms: GHS02, GHS07, GHS08

### DISCLAIMER OF EXPRESSED AND IMPLIED WARRANTIES

The information presented in this Safety Data Sheet is presented in good faith based on data believed to be accurate as of the date this Safety Data Sheet was prepared. HOWEVER, NO WARRANTY OF MERCHANTABILITY, FITNESS FOR ANY PARTICULAR PURPOSE, OR ANY OTHER WARRANTY IS EXPRESSED OR IS TO BE IMPLIED REGARDING THE ACCURACY OR COMPLETENESS OF THE INFORMATION PROVIDED ABOVE, THE RESULTS TO BE OBTAINED FROM THE USE OF THIS INFORMATION OR THE PRODUCT, THE SAFETY OF THIS PRODUCT, OR THE HAZARDS RELATED TO ITS USE. In no case shall the descriptions, information, data or designs provided be considered a part of our terms and conditions of sale.

All materials may present hazards and should be used with caution. Because many factors may affect processing or application/use, we recommend that you make tests to determine the suitability of a product for your particular purpose prior to use. No responsibility is assumed for any damage or injury resulting from abnormal use or from any failure to adhere to recommended practices or applicable federal, state, or local laws or regulations. The information provided above, and the product, are furnished on the condition that the person receiving them shall make their own determination as to the suitability of the product for their particular purpose and on the condition that they assume the risk of their use. In addition, no authorization is given nor implied to practice any patented invention without a license

REFERENCES AND DATA SOURCES: Contact the supplier for information.

METHODS OF EVALUATING INFORMATION FOR THE PURPOSE OF CLASSIFICATION: Bridging principles were used to classify this product.

<u>REVISION DETAILS</u>: September 2012: Up-date and revise entire SDS to include current GHS requirements; change in formulation. May 2017: Update of SDS to current format and regulations. May 2017: Review and up-date for current format and regulations Adjustment of formula.

DATE OF PRINTING

May 11, 2017

## **DEFINITIONS OF TERMS**

A large number of abbreviations and acronyms appear on a SDS. Some of these, which are commonly used, include the following:

#### KEY ACRONYMS:

CHEMTREC: Chemical Transportation Emergency Center, a 24-hour emergency information and/or emergency assistance to emergency responders.

CEILING LEVEL: The concentration that shall not be exceeded during any part of the working exposure.

DFG MAKs: Federal Republic of Germany Maximum Concentration Values in the workplace. Exposure limits are given as TWA (Time-Weighted Average) or PEAK (short-term exposure) values.

DFG MAK Germ Cell Mutagen Categories: 1: Germ cell mutagens that have been shown to increase the mutant frequency in the progeny of exposed humans. 2: Germ cell mutagens that have been shown to increase the mutant frequency in the progeny of exposed mammals. 3A: Substances that have been shown to induce genetic damage in germ cells of human of animals, or which produce mutagenic effects in somatic cells of mammals in vivo and have been shown to reach the germ cells in an active form. 3B: Substances that are suspected of being germ cell mutagens because of their genotoxic effects in mammalian somatic cell in vivo; in exceptional cases, substances for which there are no in vivo data, but that are clearly mutagenic in vitro and structurally related to known in vivo mutagens. 4: Not applicable (Category 4 carcinogenic substances are those with non-genotoxic mechanisms of action. By definition, germ cell mutagens are genotoxic. Therefore, a Category 4 for germ cell mutagens cannot apply. At some time in the future, it is conceivable that a Category 4 could be established for genotoxic substances with primary targets other than DNA [e.g. purely aneugenic substances] if research results make this seem sensible.) 5: Germ cell mutagens, the potency of which is considered to be so low that, provided the MAK value is observed, their contribution to genetic risk for humans is expected not to be significant.

DFG MAK Pregnancy Risk Group Classification: Group A: A risk of damage to the developing embryo or fetus has been unequivocally demonstrated. Exposure of pregnant women can lead to damage of the developing organism, even when MAK and BAT (Biological Tolerance Value for Working Materials) values are observed. Group B: Currently available information indicates a risk of damage to the developing embryo or fetus must be considered to be probable. Damage to the developing organism cannot be excluded when pregnant women are exposed, even when MAK and BAT values are observed. Group C: There is no reason to fear a risk of damage to the developing embryo or fetus when MAK and BAT values are observed.

**DFG MAK Pregnancy Risk Group Classification (continued): Group D:** Classification in one of the groups A-C is not yet possible because, although the data available may indicate a trend, they are not sufficient for final evaluation.

IDLH: Immediately Dangerous to Life and Health. This level represents a concentration from which one can escape within 30-minutes without suffering escape-preventing or permanent injury.

LOO: Limit of Ouantitation.

**NE:** Not Established. When no exposure guidelines are established, an entry of NE is made for reference.

NIC: Notice of Intended Change.

# KEY ACRONYMS (continued):

NIOSH CEILING: The exposure that shall not be exceeded during any part of the workday. If instantaneous monitoring is not feasible, the ceiling shall be assumed as a 15-minute TWA exposure (unless otherwise specified) that shall not be exceeded at any time during a workday.

(unless otherwise specified) that shall not be exceeded at any time during a workday. **NIOSH RELs:** NIOSH's Recommended Exposure Limits.

**PEL:** OSHA's Permissible Exposure Limits. This exposure value means exactly the same as a TLV, except that it is enforceable by OSHA. The OSHA Permissible Exposure Limits are based in the 1989 PELs and the June, 1993 Air Contaminants Rule (Federal Register: 58: 35338-35351 and 58: 40191). Both the current PELs and the vacated PELs are indicated. The phrase, "Vacated 1989 PEL" is placed next to the PEL that was vacated by Court Order.

SKIN: Used when a there is a danger of cutaneous absorption.

STEL: Short Term Exposure Limit, usually a 15-minute time-weighted average (TWA) exposure that should not be exceeded at any time during a workday, even if the 8-hr TWA is within the TLV-TWA, PEL-TWA or REL-TWA.

**TLV:** Threshold Limit Value. An airborne concentration of a substance that represents conditions under which it is generally believed that nearly all workers may be repeatedly exposed without adverse effect. The duration must be considered, including the 8-hour.

**TWA:** Time Weighted Average exposure concentration for a conventional 8-hr (TLV, PEL) or up to a 10-hr (REL) workday and a 40-hr workweek.

WEEL: Workplace Environmental Exposure Limits from the AIHA.

#### HAZARD RATINGS:

#### HAZARDOUS MATERIALS IDENTIFICATION SYSTEM HAZARD

**RATINGS:** This rating system was developed by the National Paint and Coating Association and has been adopted by industry to identify the degree of chemical hazards.

HEALTH HAZARD: 0 Minimal Hazard: No significant health risk, irritation of skin or eyes not anticipated. Skin Irritation: Essentially non-irritating, Mechanical irritation may occur. PII or Draize = 0. Eye Irritation: Essentially non-irritating, minimal effects clearing in < 24 hours. Mechanical irritation may occur. Draize = 0. Oral Toxicity LD<sub>50</sub> Rat: > 5000 mg/kg. Dermal Toxicity LD<sub>50</sub> Rat or Rabbit: > 2000 mg/kg. Inhalation Toxicity 4-hrs LC<sub>50</sub> Rat: > 20 mg/L. 1 Slight Hazard: Minor reversible injury may occur, may irritate the stomach if swallowed; may defat the skin and exacerbate existing dermatitis. Skin Irritation: Slightly or mildly irritating. PII or Draize > 0 < 5. Eye Irritation: Slightly to mildly irritating, but reversible within 7 days. Draize > 0 ≤ 25. Oral Toxicity LD<sub>50</sub> Rat: > 500-5000 mg/kg. Dermal Toxicity LD<sub>50</sub> Rat or Rabbit: > 1000-2000 mg/kg. Inhalation Toxicity LC<sub>50</sub> + hrs Rat: > 2-20 mg/L. 2 Moderate Hazard: Temporary or transitory injury may occur; prolonged exposure may affect the CNS. Skin Irritation: Moderately irritating; primary irritant; sensitizer. PII or Draize ≥ 5, with no destruction of dermal tissue. Eye Irritation: Moderately to severely irritating: reversible corneal opacity; corneal involvement or irritation clearing in 8-21 days. Draize = 26-100, with reversible effects. Oral Toxicity LD<sub>50</sub> Rat: > 50-500 mg/kg. Dermal Toxicity LD<sub>50</sub> Rat or Rabbit: > 200-1000 mg/kg. Inhalation Toxicity LC<sub>50</sub> 4-hrs Rat: > 0.5-2 mg/L.

# **DEFINITIONS OF TERMS (Continued)**

#### HAZARDOUS MATERIALS IDENTIFICATION SYSTEM HAZARD HAZARDOUS MATERIALS IDENTIFICATION SYSTEM HAZARD RATINGS (continued):

HEALTH HAZARD (continued): 3 Serious Hazard: Major injury likely unless prompt action is taken and medical treatment is given; high level of toxicity; corrosive. Skin Irritation: Severely irritating and/or corrosive; may cause destruction of dermal tissue, skin burns, and dermal necrosis. PII or Draize > 5-8, with destruction of tissue. Eye Irritation: Corrosive, irreversible destruction of ocular tissue; corneal involvement or irritation persisting for more than 21 days. Draize > 80 with effects irreversible in 21 days. Oral Toxicity  $LD_{50}$  Rat: > 1-50 mg/kg. Dermal Toxicity  $LD_{50}$  Rat or Rabbit: > 20-200 mg/kg. Inhalation Toxicity  $LC_{50}$  4-hrs Rat: > 0.05-0.5 mg/L. 4 Severe Hazard: Lifethreatening; major or permanent damage may result from single or repeated exposure; extremely toxic; irreversible injury may result from brief contact. Skin Irritation: Not appropriate. Do not rate as a 4, based on skin irritation alone. Eye Irritation: Not appropriate. Do not rate as a 4, based on eye irritation alone. Oral Toxicity  $LD_{50}$  Rat:  $\leq 1$  mg/kg. Dermal Toxicity  $LD_{50}$  Rat or Rabbit:  $\leq 20$  mg/kg. Inhalation Toxicity  $LC_{50}$  4-hrs Rat:  $\leq 0.05$  mg/L.

FLAMMABILITY HAZARD: 0 Minimal Hazard: Materials that will not burn in air when exposure to a temperature of 815.5°C (1500°F) for a period of 5 minutes. 1 Slight Hazard: Materials that must be pre-heated before ignition can occur. Material requires considerable pre-heating, under all ambient temperature conditions before ignition and combustion can occur. This usually includes the following: Materials that will burn in air when exposed to a temperature of  $815.5^{\circ}$ C ( $1500^{\circ}$ F) for a period of 5 minutes or less; Liquids, solids and semisolids having a flash point at or above 93.3°C (200°F) (i.e. OSHA Class IIIB); and Most ordinary combustible materials (e.g. wood, paper, etc.). 2 Moderate Hazard: Materials that must be moderately heated or exposed to relatively high ambient temperatures before ignition can occur. Materials in this degree would not, under normal conditions, form hazardous atmospheres in air, but under high ambient temperatures or moderate heating may release vapor in sufficient quantities to produce hazardous atmospheres with air. This usually includes the following: Liquids having a flash-point at or above 37.8°C (100°F); Solid materials in the form of course dusts that may burn rapidly but that generally do not form explosive atmospheres; Solid materials in a fibrous or shredded form that may burn rapidly and create flash fire hazards (e.g. cotton, sisal, hemp); and Solids and semisolids (e.g. viscous and slow flowing as asphalt) that readily give off flammable vapors. 3 Serious Hazard: Liquids and solids that can be ignited under almost all ambient temperature conditions. Materials in this degree produce hazardous atmospheres with air under almost all ambient temperatures, or, unaffected by ambient temperature, are readily ignited under almost all conditions. This usually includes the following: Liquids having a flash point below 22.8°C (73°F) and having a boiling point at or above 38°C (100°□F) and those liquids having a flash point at or above 22.8°C (73°F) and below 37.8°C (100°F) (i.e. OSHA Class IB and IC); Materials that on account of their physical form or environmental conditions can form explosive mixtures with air and are readily dispersed in air (e.g., dusts of combustible solids, mists or droplets of flammable liquids); and Materials that burn extremely rapidly, usually by reason of self-contained oxygen (e.g. dry nitrocellulose and many organic peroxides). 4 Severe Hazard: Materials that will rapidly or completely vaporize at atmospheric pressure and normal ambient temperature or that are readily dispersed in air, and that will burn readily. This usually includes the following: Flammable gases; Flammable cryogenic materials; Any liquid or gaseous material that is liquid while under pressure and has a flash point below 22.8°C (73°F) and a boiling point below 37.8°C (100°F) (i.e. OSHA Class IA); and Materials that ignite spontaneously when exposed to air at a temperature of 54.4°C (130°F) or below (pyrophoric).

PHYSICAL HAZARD: 0 Water Reactivity: Materials that do not react with water. Organic Peroxides: Materials that are normally stable, even under fire conditions and will not react with water. Explosives: Substances that are Non-Explosive. Compressed Gases: No Rating. Pyrophorics: No Rating. Oxidizers: No 0 rating. Unstable Reactives: Substances that will not polymerize, decompose, condense, or self-react.). 1 Water Reactivity: Materials that change or decompose upon exposure to moisture. Organic Peroxides: Materials that are normally stable, but can become unstable at high temperatures and pressures. These materials may react with water, but will not release energy violently. Explosives: Division 1.5 & 1.6 explosives. Substances that are very insensitive explosives or that do not have a mass explosion hazard. Compressed Gases: Pressure below OSHA definition. Pyrophorics: No Rating. Oxidizers: Packaging Group III oxidizers; Solids: any material that in either concentration tested, exhibits a mean burning time less than or equal to the mean burning time of a 3:7 potassium bromate/cellulose mixture and the criteria for Packing Group I and II are not met. Liquids: any material that exhibits a mean pressure rise time less than or equal to the pressure rise time of a 1:1 nitric acid (65%)/cellulose mixture and the criteria for Packing Group I and II are not met. Unstable Reactives: Substances that may decompose condense, or self-react, but only under conditions of high temperature and/or pressure and have little or no potential to cause significant heat generation or explosion hazard. Substances that readily undergo hazardous polymerization in the absence of inhibitors. 2 Water Reactivity: Materials that may react violently with water. Organic Peroxides: Materials that, in themselves, are normally unstable and will readily undergo violent chemical change, but will not detonate. These materials may also react violently with water. Explosives: Division 1.4 explosives. Explosive substances where the explosive effects are largely confined to the package and no projection of fragments of appreciable size or range are expected. An external fire must not cause virtually instantaneous explosion of almost the entire contents of the package. Compressed Gases: Pressurized and meet OSHA definition but < 514.7 psi absolute at 21.1°C (70°F) [500 psig]. Pyrophorics: No Rating. Oxidizers: Packing Group II oxidizers. Solids: any material that, either in concentration tested, exhibits a mean burning time of less than or equal to the mean burning time of a 2:3 potassium bromate/cellulose mixture and the criteria for Packing Group I are not met. Liquids: any material that exhibits a mean pressure rise time less than or equal to the pressure rise of a 1:1 aqueous sodium chlorate solution (40%)/cellulose mixture and the criteria for Packing Group I are not met. Reactives: Substances that may polymerize, decompose, condense, or self-react at ambient temperature and/or pressure, but have a low potential (or low risk) for significant heat generation or explosion. Substances that readily form peroxides upon exposure to air or oxygen at room temperature. 3 Water Reactivity: Materials that may form explosive reactions with water. Organic Peroxides: Materials that are capable of detonation or explosive reaction, but require a strong initiating source or must be heated under confinement before initiation; or materials that react explosively with water. Explosives: Division 1.3 explosives. Explosive substances that have a fire hazard and either a minor blast hazard or a minor projection hazard or both, but do not have a mass explosion hazard. Compressed Gases: Pressure ≥ 514.7 psi absolute at 21.1°C (70°F) [500 psig]. Pyrophorics: No Rating. Oxidizers: Packing Group I oxidizers. Solids: any material that, in either concentration tested, exhibits a mean burning time less than the mean burning time of a 3:2 potassium bromate/cellulose mixture. Liquids: any material that spontaneously ignites when mixed with cellulose in a 1:1 ratio, or which exhibits a mean pressure rise time less than the pressure rise time of a 1:1 perchloric acid (50%)/cellulose mixture. Unstable Reactives: Substances that may polymerize, decompose, condense, or self-react at ambient temperature and/or pressure and have a moderate potential (or moderate risk) to cause significant heat generation or explosion. 4 Water Reactivity: Materials that react explosively with water without requiring heat or confinement, Organic Peroxides: Materials that are readily capable of detonation or explosive decomposition at normal temperature and pressures. Explosives: Division 1.1 & 1.2 explosives. Explosive substances that have a mass explosion hazard or have a projection hazard. A mass explosion is one that affects almost the entire load instantaneously.

# RATINGS (continued):

PHYSICAL HAZARD (continued): 4 (continued): Compressed Gases: No Rating. Pyrophorics: Add to the definition of Flammability 4. Oxidizers: No 4 rating. Unstable Reactives: Substances that may polymerize, decompose, condense, or self-react at ambient temperature and/or pressure and have a high potential (or high risk) to cause significant heat generation or explosion. Pyrophorics: Add to the definition of Flammability 4. Oxidizers: No 4 rating. Unstable Reactives: Substances that may polymerize, decompose, condense, or self-react at ambient temperature and/or pressure and have a high potential (or high risk) to cause significant heat generation or explosion.

#### NATIONAL FIRE PROTECTION ASSOCIATION HAZARD RATINGS:

<u>HEALTH HAZARD</u>: 0 Materials that, under emergency conditions, would offer no hazard beyond that of ordinary combustible materials. Gases and vapors with an LC<sub>50</sub> for acute inhalation toxicity greater than 10,000 ppm. Dusts and mists with an LC50 for acute inhalation toxicity greater than 200 mg/L. Materials with an LD50 for acute dermal toxicity greater than 2000 mg/kg. Materials with an LD50 for acute oral toxicity greater than 2000 mg/kg. Materials essentially non-irritating to the respiratory tract, eyes, and skin. 1 Materials that, under emergency conditions, can cause significant irritation. Gases and vapors with an LC<sub>50</sub> for acute inhalation toxicity greater than 5,000 ppm but less than or equal to 10,000 ppm. Dusts and mists with an LC<sub>50</sub> for acute inhalation toxicity greater than 10 mg/L but less than or equal to 200 mg/L. Materials with an LD<sub>50</sub> for acute dermal toxicity greater than 1000 mg/kg but less than or equal to 2000 mg/kg. Materials that slightly to moderately irritate the respiratory tract, eyes and skin. Materials with an LD50 for acute oral toxicity greater than 500 mg/kg but less than or equal to 2000 mg/kg. 2 Materials that, under emergency conditions, can cause temporary incapacitation or residual injury. Gases with an  $LC_{50}$  for acute inhalation toxicity greater than 3,000 ppm but less than or equal to 5,000 ppm. Any liquid whose saturated vapor concentration at 20°C (68°F) is equal to or greater than one-fifth its LC50 for acute inhalation toxicity, if its LC50 is less than or equal to 5000 ppm and that does not meet the criteria for either degree of hazard 3 or degree of hazard 4. Dusts and mists with an LC50 for acute inhalation toxicity greater than 2 mg/L but less than or equal to 10 mg/L. Materials with an LD50 for acute dermal toxicity greater than 200 mg/kg but less than or equal to 1000 mg/kg. Compressed liquefied gases with boiling points between -30°C (-22°F) and -55°C (-66.5°F) that cause severe tissue damage, depending on duration of exposure. Materials that are respiratory irritants. Materials that cause severe, but reversible irritation to the eyes or are lachrymators. Materials that are primary skin irritants or sensitizers. Materials whose LD<sub>50</sub> for acute oral toxicity is greater than 50 mg/kg but less than or equal to 500 mg/kg. 3 Materials that, under emergency conditions, can cause serious or permanent injury. Gases with an LC<sub>50</sub> for acute inhalation toxicity greater than 1,000 ppm but less than or equal to 3,000 ppm. Any liquid whose saturated vapor concentration at 20°C (68°F) is equal to or greater its LC<sub>50</sub> for acute inhalation toxicity, if its LC50 is less than or equal to 3000 ppm and that does not meet the criteria for degree of hazard 4. Dusts and mists with an LC50 for acute inhalation toxicity greater than 0.5 mg/L but less than or equal to 2 mg/L. Materials with an LD50 for acute dermal toxicity greater than 40 mg/kg but less than or equal to 200 mg/kg. Materials that are corrosive to the respiratory tract. Materials that are corrosive to the eyes or cause irreversible corneal opacity. Materials corrosive to the skin. Cryogenic gases that cause frostbite and irreversible tissue damage. Compressed liquefied gases with boiling points below -55°C (-66.5°F) that cause frostbite and irreversible tissue damage. Materials with an LD50 for acute oral toxicity greater than 5 mg/kg but less than or equal to 50 mg/kg. 4 Materials that, under emergency conditions, can be lethal. Gases with an LC50 for acute inhalation toxicity less than or equal to 1,000 ppm. Any liquid whose saturated vapor concentration at 20°C (68°F) is equal to or greater than ten times its  $LC_{50}$  for acute inhalation toxicity, if its  $LC_{50}$  is less than or equal to 1000 ppm. Dusts and mists whose  $LC_{50}$  for acute inhalation toxicity is less than or equal to 0.5 mg/L. Materials whose LD50 for acute dermal toxicity is less than or equal to 40 mg/kg. Materials whose LD50 for acute oral toxicity is less than or equal to 5 mg/kg.

FLAMMABILITY HAZARD: 0 Materials that will not burn under typical fire conditions, including intrinsically noncombustible materials such as concrete, stone, and sand. Materials that will not burn in air when exposed to a temperature of 816°C (1500°F) for a period of 5 minutes in according with Annex D of NFPA 704. 1 Materials that must be preheated before ignition can occur. Materials in this degree require considerable preheating, under all ambient temperature conditions, before ignition and combustion can occur: Materials that will burn in air when exposed to a temperature of 816°C (1500°F) for a period of 5 minutes in according with Annex D of NFPA 704. Liquids, solids, and semisolids having a flash point at or above 93.4°C (200°F) (i.e. Class IIIB liquids). Liquids with a flash point greater than 35°C (95°F) that do not sustain combustion when tested using the Method of Testing for Sustained Combustibility, per 49 CFR 173, Appendix H or the UN Recommendations on the Transport of Dangerous Goods, Model Regulations (current edition) and the related Manual of Tests and Criteria (current edition). Liquids with a flash point greater than 35°C (95°F) in a watermiscible solution or dispersion with a water non-combustible liquid/solid content of more than 85% by weight. Liquids that have no fire point when tested by ASTM D 92, Standard Test Method for Flash and Fire Points by Cleveland Open Cup, up to the boiling point of the liquid or up to a temperature at which the sample being tested shows an obvious physical change. Combustible pellets with a representative diameter of greater than 2 mm (10 mesh). Most ordinary combustible materials. Solids containing greater than 0.5% by weight of a flammable or combustible solvent are rated by the closed cup flash point of the solvent. 2 Materials that must be moderately heated or exposed to relatively high ambient temperatures before ignition can occur. Materials in this degree would not under normal conditions form hazardous atmospheres with air, but under high ambient temperatures or under moderate heating could release vapor in sufficient quantities to produce hazardous atmospheres with air. Liquids having a flash point at or above 37.8°C (100°F) and below 93.4°C (200°F) (i.e. Class II and Class IIIA liquids.) Solid materials in the form of powders or coarse dusts of representative diameter between 420 microns (40 mesh) and 2 mm (10 mesh) that burn rapidly but that generally do not form explosive mixtures with air. Solid materials in fibrous or shredded form that burn rapidly and create flash fire hazards, such as cotton, sisal, and hemp. Solids and semisolids that readily give off flammable vapors. Solids containing greater than 0.5% by weight of a flammable or combustible solvent are rated by the closed cup flash point of the solvent. 3 Liquids and solids that can be ignited under almost all ambient temperature conditions. Materials in this degree produce hazardous atmospheres with air under almost all ambient temperatures or, though unaffected by ambient temperatures, are readily ignited under almost all conditions. Liquids having a flash point below 22.8°C (73°F) and having a boiling point at or above 37.8°C (100°F) and those liquids having a flash point at or above 22.8°C (73°F) and below 37.8°C (100°F) (i.e. Class IB and IC liquids). Materials that on account of their physical form or environmental conditions can form explosive mixtures with air and are readily dispersed in air. Flammable or combustible dusts with representative diameter less than 420 microns (40 mesh). Materials that burn with extreme rapidity, usually by reason of self-contained oxygen (e.g. dry nitrocellulose and many organic peroxides). Solids containing greater than 0.5% by weight of a flammable or combustible solvent are rated by the closed cup flash point of the solvent. 4 Materials that will rapidly or completely vaporize at atmospheric pressure and normal ambient temperature or that are readily dispersed in air and will burn readily. Flammable gases. Flammable cryogenic materials. Any liquid or gaseous materials that is liquid while under pressure and has a flash point below 22.8°C (73°F) and a boiling point below 37.8°C (100°F) (i.e. Class IA liquids). Materials that ignite when exposed to air, Solids containing greater than 0.5% by weight of a flammable or combustible solvent are rated by the closed cup flash point of the

# **DEFINITIONS OF TERMS (Continued)**

# NATIONAL FIRE PROTECTION ASSOCIATION HAZARD RATINGS (continued):

INSTABILITY HAZARD: 0 Materials that in themselves are normally stable, even under fire conditions. Materials that have an instantaneous power density (product of heat of reaction and reaction rate) at  $250^{\circ}$ C ( $482^{\circ}$ F) below 0.01 W/mL. Materials that do not exhibit an exotherm at temperatures less than or equal to 500°C (932°F) when tested by differential scanning calorimetry. 1 Materials that in themselves are normally stable, but that can become unstable at elevated temperatures and pressures. Materials that have an instantaneous power density (product of heat of reaction and reaction rate) at 250°C (482°F) at or above 0.01 W/mL and below 10 W/mL. 2 Materials that readily undergo violent chemical change at elevated temperatures and pressures. Materials that have an instantaneous power density (product of heat of reaction and reaction rate) at 250°C (482°F) at or above 10 W/mL and below 100W/mL. 3 Materials that in themselves are capable of detonation or explosive decomposition or explosive reaction, but that require a strong initiating source or that must be heated under confinement before initiation. Materials that have an estimated instantaneous power density (product of heat of reaction and reaction rate) at 250°C (482°F) at or above 100 W/mL and below 1000 W/mL. Materials that are sensitive to thermal or mechanical shock at elevated temperatures and pressures. 4 Materials that in themselves are readily capable of detonation or explosive decomposition or explosive reaction at normal temperatures and pressures. Materials that are sensitive to localized thermal or mechanical shock at normal temperatures and pressures. Materials that have an estimated instantaneous power density (product of heat of reaction and reaction rate) at 250°C (482°F) of 1000 W/mL or greater.

#### FLAMMABILITY LIMITS IN AIR:

Much of the information related to fire and explosion is derived from the National Fire Protection Association (NFPA). Flash Point: Minimum temperature at which a liquid gives off sufficient vapor to form an ignitable mixture with air near the surface of the liquid or within the test vessel used. Autoignition Temperature: Minimum temperature of a solid, liquid, or gas required to initiate or cause self-sustained combustion in air with no other source of ignition. LEL: Lowest concentration of a flammable vapor or gas/air mixture that will ignite and burn with a flame. UEL: Highest concentration of a flammable vapor or gas/air mixture that will ignite and burn with a flame.

#### TOXICOLOGICAL INFORMATION:

Human and Animal Toxicology: Possible health hazards as derived from human data, animal studies, or from the results of studies with similar compounds are presented. <u>LDs</u>: Lethal Dose (solids & liquids) that kills 50% of the exposed animals. LCs₀: Lethal Concentration (gases) that kills 50% of the exposed animals. <u>ppm</u>: Concentration expressed in parts of material per million parts of air or water. <u>mg/m²</u>: Concentration expressed in weight of substance per volume of air. <u>mg/kg</u>: Quantity of material, by weight, administered to a test subject, based on their body weight in <u>kg</u>. <u>TDLo</u>: Lowest dose to cause a symptom. <u>TCLo</u>: Lowest concentration to cause a symptom. <u>TDo</u>, <u>LDLo</u>, and <u>LDo</u>, or <u>TC</u>, <u>TCo</u>, <u>LCLo</u>, and <u>LCo</u>: Lowest dose (or concentration) to cause lethal or toxic effects. Cancer Information: <u>LARC</u>: International Agency for Research on Cancer. <u>NTP</u>: National Toxicology Program. <u>RTECS</u>: Registry of Toxic Effects of Chemical Substances. <u>IARC</u> and NTP rate chemicals on a scale of decreasing potential to cause human cancer with rankings from 1 to 4. Subrankings (2A, 2B, etc.) are also used. **Other Information**: <u>BEI</u>: ACGIH Biological Exposure Indices, represent the levels of determinants which are most likely to be observed in specimens collected from a healthy worker who has been exposed to chemicals to the same extent as a worker with inhalation exposure to the TLV.

**REPRODUCTIVE INFORMATION:** A <u>mutagen</u> is a chemical that causes permanent changes to genetic material (DNA) such that the changes will propagate through generational lines. An <u>embryotoxin</u> is a chemical that causes damage to a developing embryo (i.e. within the first eight weeks of pregnancy in humans), but the damage does not propagate across generational lines. A <u>teratogen</u> is a chemical that causes damage to a developing fetus, but the damage does not propagate across generational lines. A <u>reproductive toxin</u> is any substance that interferes in any way with the reproductive process.

#### ECOLOGICAL INFORMATION:

EC: Effect concentration in water. BCF: Bioconcentration Factor, which is used to determine if a substance will concentrate in life forms that consume contaminated plant or animal matter. TLm: Median threshold limit. log Kow or log Koc: Coefficient of Oil/Water Distribution is used to assess a substance's behavior in the environment.

**REGULATORY INFORMATION:** This section explains the impact of various laws and regulations on the material.

U.S.:

EPA: U.S. Environmental Protection Agency. ACGIH: American Conference of Governmental Industrial Hygienists, a professional association that establishes exposure limits. OSHA: U.S. Occupational Safety and Health Administration. NIOSH: National Institute of Occupational Safety and Health, which is the research arm of OSHA. DOT: U.S. Department of Transportation. TC: Transport Canada. SARA: Superfund Amendments and Reauthorization Act. TSCA: U.S. Toxic Substance Control Act. CERCLA: Comprehensive Environmental Response, Compensation, and Liability Act. Marine Pollutant status according to the DOT; CERCLA or Superfund; and various state regulations. This section also includes information on the precautionary warnings that appear on the material's package label.

#### CANADA:

WHMIS: Canadian Workplace Hazardous Materials Information System. <u>TC</u>: Transport Canada. <u>DSL/NDSL</u>: Canadian Domestic/Non-Domestic Substances List.