

# SILVERBACK INK®

## SAFETY DATA SHEET

Prepared to U.S. OSHA, CMA, ANSI, Canadian WHMIS 2015 (HPR-GHS), European Union CLP EC 1272/2008 and REACH, Australian NOHSC, New Zealand HNSO, the Korean ISHA (Notice 2009-68), Singapore SS586 - 2: 2008 and SS 586 - 3: 2008 Standards, Chinese GB 20576 ~ GB 20602-2006, Japanese JIS Z7253, Taiwanese Standards and the Global Harmonization Standard, and requirements under Chemical Control Regulations of Argentina, Chile, Colombia, Costa Rica, Panama, Honduras, Venezuela, Uruguay, Peru and Paraguay.

### 1. IDENTIFICATION of the SUBSTANCE or PREPARATION

#### IDENTIFICATION OF SUBSTANCE/PREPARATION:

<u>TRADE NAME (AS LABELED):</u>	<b>Silverback Ink White</b>
<u>CHEMICAL NAME/CLASS:</u>	Water-Based Ink
<u>SYNONYMS:</u>	Not Applicable
<u>PRODUCT USE:</u>	Various Uses
<u>U.N. NUMBER:</u>	None Allocated
<u>U.N. DANGEROUS GOODS CLASS/SUBSIDIARY RISK:</u>	None Allocated
<u>HAZCHEM CODE (AUSTRALIA):</u>	None Allocated
<u>POISONS SCHEDULE NUMBER (AUSTRALIA):</u>	None Allocated

#### COMPANY/UNDERTAKING IDENTIFICATION:

<u>U.S. SUPPLIER/MANUFACTURER'S NAME:</u>	<b>SILVERBACK INK CORP</b>
<u>ADDRESS:</u>	4869 Fountain Ave Los Angeles, CA 90029, USA
<u>INFORMATION PHONE:</u>	1-323-669-8282

NOTE: ALL United States Occupational Safety and Health Administration Standard (29 CFR 1910.1200), U.S. State equivalent Standards, Canadian WHMIS [Controlled Products Regulations], the European Union CLP EC 1272/2008 and the Global Harmonization Standard, Australian [NOHSC:2012 (2003)], and Japanese Industrial Standard (JIS Z 7250: 2000) required information is included in appropriate sections based on the Global Harmonization Standard. This product has been classified in accordance with the hazard criteria of the countries listed above.

### 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 2-15, EU CLP Regulation (EC) 1272/2008, Japanese JIS Z7253: 2012 and Singapore Standards. For information on Korean ISHA and New Zealand HSNO classification, see below.

Classification: Eye Irritation Cat. 2A      Signal Word: Warning      Hazard Statement Codes: H319  
Precautionary Statement Codes: P264, P280, P305 + P351 + P338, P337 + P313  
Hazard Symbol/Pictogram: GHS07



KOREAN ISHA (Notice 2009-68) LABELING AND CLASSIFICATION: Classified in accordance with ISHA Notice 2009-68. Under ISHA, the following differences in classification are applicable.

Classification: Eye Irritation Cat. 2      Signal Word: Warning      Hazard Statement Codes: H319  
Precautionary Statement Codes: P264, P280, P305 + P351 + P338, P337 + P313  
Hazard Symbol/Pictogram: GHS07



NEW ZEALAND HAZARDOUS SUBSTANCES and NEW ORGANISMS ACT (HNSO) CHEMICAL CLASSIFICATION:

Product Group Standard: Not Otherwise Classified, Subsidiary Hazard  
Classification: 6.4A: Irritating to the eye.

See Section 16 for full text of classification of product and components

**EMERGENCY OVERVIEW: Product Description:** This product is a white liquid that has a mild odor. **Health Hazards:** The primary health hazard associated with this product is the potential for mild irritation of contaminated tissue. Eye contact may cause irritation. The inks may stain skin, eyes, other contaminated tissue, and objects. The Titanium Dioxide component is a suspect carcinogen by inhalation of respirable particles. Due to the liquid form of this product, this hazard is not expected to be significant. **Flammability Hazards:** This product is not flammable. If involved in a fire, the product may decompose to produce calcium, carbon, nitrogen and nitrogen oxides. **Reactivity Hazards:** This product is not reactive. **Environmental Hazards:** Not tested. This product may have adverse effects when released into the environment. **Emergency Recommendations:** Emergency responders must wear the personal protective equipment suitable for the situation to which they are responding.

### 3. COMPOSITION and INFORMATION ON INGREDIENTS

Chemical Name	CAS #	European EINECS #	Japanese MITI/ENC #	Korean ECL #	New Zealand NZIoC #	Chinese IECSC Inventory	Taiwan NESCI ECS Inventory	Australian AICS	% w/w	LABEL ELEMENTS GHS under U.S. OSHA, Canadian WHMIS HPR-GHS & EU Classification (1272/2008), Japanese, Taiwan, Chinese and Korean Regulations Korean ISHA Classification Hazard Statement Codes
Titanium Dioxide	13463-67-7	236-675-5	1-558	KE-33900	Group Listing	Listed	Listed	Listed	45.0	GHS under U.S. OSHA, Canadian WHMIS & EU CLP 1272/2008, KOREAN ISHA Classification: Carcinogenic Category 2 Hazard Statement Codes: H351i
Proprietary Suspension Agent			Excepted as a Mineral	Proprietary	Group Listing	Listed	Listed	Listed	20.0	GHS under U.S. OSHA, Canadian WHMIS HPR-2015, Korean ISHA & EU CLP 1272/2008 Classification: Not Classified
Proprietary Dispersing Agent					Group Listing	Listed	Listed	Listed	12.0	GHS under Korean ISHA & EU CLP 1272/2008 Classification: Not Applicable Under U.S. OSHA, Canadian WHMIS HPR-2015 only Classification: Combustible Dust Hazard
Proprietary Alcohol						Listed	Listed	Listed	10.0	HARMONISED CLASSIFICATION - ANNEX VI of REGULATION (EC) No 1272/2008 (CLP00) Classification: Flammable Liquid Category 2, Eye Irritant Category 2A, STOT SE 3 Hazard Statement Codes: H225, H319, H336 GHS under Korean ISHA Classification: Flammable Liquid Category 2, Eye Irritant Category 2, STOT (Inhalation-Narcotic Effect) SE 3 Hazard Statement Codes: H225, H319, H336
Water	7732-18-5	231-791-2	Not Identified in the Listing	KE-35400	Excepted	Listed	Listed	Listed	Balance	GHS under U.S. OSHA, Canadian WHMIS & EU CLP 1272/2008, KOREAN ISHA Classification: Not Classified

See Section 16 for full text of classification. See Section 15 for information on other country inventory listing of components, as applicable

### 4. FIRST-AID MEASURES

**PROTECTION OF FIRST AID RESPONDERS:** Rescuers should be taken for medical attention, if necessary. Only trained personnel should administer supplemental oxygen and/or cardio-pulmonary resuscitation, if necessary.

**DESCRIPTION OF FIRST AID MEASURES:** Victim(s) must be taken for medical attention. Take copy of label and MSDS to physician or other health professional with victim(s). Remove victim(s) to fresh air, as quickly as possible.

**Skin Exposure:** If this product contaminates the skin, begin decontamination with running water. Minimum flushing is for 20 minutes. The contaminated individual must seek medical attention if any adverse effects occur after flushing.

**Eye Exposure:** If this product enters the eyes, open contaminated individual's eyes while under gently running water. Use sufficient force to open eyelids. Have contaminated individual "roll" eyes. Minimum flushing is for 20 minutes. Contaminated individual must seek medical attention if adverse effect continues after flushing.

**Inhalation:** If aerosols are inhaled, remove victim to fresh air. The contaminated individual must seek medical attention if any adverse effects occur.

**Ingestion:** If swallowed, CALL PHYSICIAN OR POISON CONTROL CENTER FOR MOST CURRENT INFORMATION.

**DESCRIPTION OF FIRST AID MEASURES (continued):**

**Ingestion (continued):** If professional advice is not available, do not induce vomiting. Never induce vomiting or give diluents (milk or water) to someone who is unconscious, having convulsions, or unable to swallow. If victim is convulsing, maintain an open airway and obtain immediate medical attention.

**IMPORTANT SYMPTOMS AND EFFECTS:** See Sections 2 (Hazard Information) and 11 (Toxicological Information) for information.

**MEDICAL CONDITIONS AGGRAVATED BY EXPOSURE:** Skin or respiratory conditions may be aggravated by exposure to this product.

**INDICATION OF IMMEDIATE MEDICAL ATTENTION AND SPECIAL TREATMENT IF NEEDED:** Treat symptoms and eliminate exposure.

### 5. FIRE-FIGHTING MEASURES

**FLASH POINT:** Not flammable.

**AUTOIGNITION TEMPERATURE:** Not applicable.

**FLAMMABLE LIMITS (in air by volume, %):** Not applicable.

**FIRE EXTINGUISHING MEDIA:** Unless incompatibilities exist for surrounding materials, carbon dioxide, water spray, 'ABC' type chemical extinguishers, foam, dry chemical and halon extinguishers can be used to fight fires involving this product.

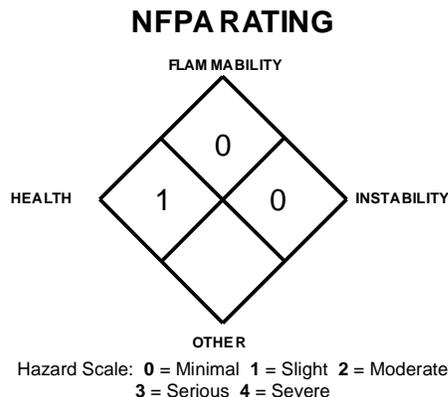
## 5. FIRE-FIGHTING MEASURES (Continued)

UNSUITABLE FIRE EXTINGUISHING MEDIA: None known.

SPECIAL HAZARDS ARISING FROM THE SUBSTANCE: When involved in a fire, this material may decompose and produce irritating vapors and toxic gases (e.g., calcium, carbon, nitrogen and nitrogen oxides).

Explosion Sensitivity to Mechanical Impact or Static Discharge: Not sensitive.

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. Due to the presence of pigment, the runoff water from this product can discolor contaminated objects. If possible, prevent runoff water from entering storm drains, bodies of water, or other environmentally sensitive areas. If necessary, rinse fire-response equipment with soapy water before returning it to service.



## 6. ACCIDENTAL RELEASE MEASURES

PERSONAL PRECAUTIONS AND EMERGENCY PROCEDURES: Uncontrolled releases should be responded to by trained personnel using pre-planned procedures. Proper protective equipment should be used. Call CHEMTREC (1-800-424-9300) for emergency assistance. Or if in Canada, call CANUTEC (613-996-6666). 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.

PERSONAL PROTECTIVE EQUIPMENT: Proper protective equipment should be used.

Small Spills: Wear rubber gloves, splash goggles, and appropriate body protection.

Large Spills: 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 and boots, hard hat, and Self-Contained Breathing Apparatus.**

METHODS FOR CLEAN-UP AND CONTAINMENT:

Small Spills: Carefully absorb spill using polypads or other non-reactive absorbent. Place spilled material in appropriate container for disposal, sealing tightly. Remove all residue before decontamination of spill area.

Large Spills: Access to the spill area should be restricted. For large spills, dike or otherwise contain spill and absorb spill with polypads or other non-reactive absorbent material. Monitor area for combustible vapor levels.

All Spills: Place all spill residue in a double plastic bag or other containment and seal. Decontaminate the area thoroughly. Do not mix with wastes from other materials. Dispose of in accordance with applicable Federal, State, and local procedures (see Section 13, Disposal Considerations). For spills on water, contain, minimize dispersion and collect. Dispose of recovered material and report spill per regulatory requirements.

ENVIRONMENTAL PRECAUTIONS: Avoid release to the environment. Run-off water may be contaminated by other materials and should be contained to prevent possible environmental damage.

REFERENCE TO OTHER SECTIONS: 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, drink, smoke, or apply cosmetics while handling this product. Avoid breathing aerosols from the product. Keep away from incompatible materials (see Section 10, Stability and Reactivity). Containers of this product must be properly labeled. Use in a well-ventilated location. Remove contaminated clothing.

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. Have appropriate extinguishing equipment in the storage area (such as sprinkler systems or portable fire extinguishers). Empty containers may contain residual product; therefore, empty containers should be handled with care.

SPECIFIC USE(S): This product is for use as an ink. Follow all industry standards for use of this product.

PROTECTIVE PRACTICES DURING MAINTENANCE OF CONTAMINATED EQUIPMENT: Follow practices indicated in Section 6 (Accidental Release Measures). Make certain that application equipment is locked and tagged-out safely. Always use this product in areas where adequate ventilation is provided. Decontaminate equipment thoroughly, before maintenance begins. Collect all rinsates and dispose of according to applicable Federal, State, or local procedures, or applicable standards.

## 8. EXPOSURE CONTROLS - PERSONAL PROTECTION

EXPOSURE LIMITS/CONTROL PARAMETERS:

Ventilation and Engineering Controls: Use with adequate ventilation to ensure exposure levels are maintained below the limits provided in this section. Use local exhaust ventilation. Normal office ventilation conforming to the American Society of Heating, Refrigerating, and Air Conditioning Engineers (ASHRAE) Standards is adequate under normal circumstances of use.

## 8. EXPOSURE CONTROLS - PERSONAL PROTECTION (Continued)

### EXPOSURE LIMITS/CONTROL PARAMETERS (continued):

**Ventilation and Engineering Controls (continued):** Persons using this material should consult a qualified Ventilation Engineer and/or Industrial Hygienist if concerns about exposure arise. If necessary, refer to Australian National Code of Practice for the Control of Workplace Hazardous Substances [NOHSC: 2007 (1994)] for further information.

### Workplace Exposure Limits/Control Parameters:

CHEMICAL NAME	CAS #	EXPOSURE LIMITS IN AIR							
		ACGIH-TLVs		OSHA-PELs		NIOSH-RELs		NIOSH	OTHER
		TWA mg/m <sup>3</sup>	STEL mg/m <sup>3</sup>	TWA mg/m <sup>3</sup>	STEL mg/m <sup>3</sup>	TWA mg/m <sup>3</sup>	STEL mg/m <sup>3</sup>	IDLH mg/m <sup>3</sup>	mg/m <sup>3</sup>
Proprietary Suspension Agent		NE	NE	NE	NE	NE	NE	NE	NE
Proprietary Dispersing Agent		NE	NE	NE	NE	NE	NE	NE	NE
Proprietary Alcohol		492	984	980	500 ppm (vacated 1989 PEL)	980	1225	2000 ppm (based on LEL)	DFG MAK: TWA = 500 ppm PEAK = 2•MAK, 15 min., average value, 1-hr interval, 4 per shift DFG MAK Pregnancy Risk Classification: C
Titanium Dioxide	13463-67-7	10	NE	15 (total dust)	NE	See Pocket Guide Appendix A	NE	5000	Carcinogen: IARC-2B, MAK-3A, NIOSH-Ca, TLV-A4

NE = Not Established.

See Section 16 for Definitions of Other Terms Used

**International Occupational Exposure Limits:** The following international limits are in place for some components of this product. Limits may have changed since time of preparation of this SDS and should be checked with competent authorities of individual countries.

#### ISOPROPYL ALCOHOL:

Australia: TWA = 400 ppm (983 mg/m<sup>3</sup>), STEL = 500 ppm (1230 mg/m<sup>3</sup>), JUL 2008  
 Austria: MAK-TMW 200 ppm (500 mg/m<sup>3</sup>); KZW = 800 ppm (2000 mg/m<sup>3</sup>), 2007  
 Belgium: TWA = 400 ppm (997 mg/m<sup>3</sup>), MAR 2002  
 Belgium: STEL = 500 ppm (1248 mg/m<sup>3</sup>), MAR 2002  
 Denmark: TWA = 200 ppm (490 mg/m<sup>3</sup>), MAY 2011  
 France: VLE = 400 ppm (980 mg/m<sup>3</sup>), FEB 2006  
 Germany: MAK = 500 mg/m<sup>3</sup> (200 mL/m<sup>3</sup>), 2005  
 Hungary: TWA = 500 mg/m<sup>3</sup>, STEL = 2000 mg/m<sup>3</sup>, Skin, SEP 2000  
 Iceland: TWA = 200 ppm (490 mg/m<sup>3</sup>), skin, NOV 2011  
 Japan: OEL-C = 400 ppm (980 mg/m<sup>3</sup>), MAY 2009  
 Korea: TWA = 400 ppm (980 mg/m<sup>3</sup>), STEL = 500 ppm (1225 mg/m<sup>3</sup>), 2006  
 Mexico: TWA = 400 ppm (980 mg/m<sup>3</sup>); STEL = 500 ppm (1225 mg/m<sup>3</sup>), 2004  
 The Netherlands: MAC-TGG = 650 mg/m<sup>3</sup>, 2003  
 New Zealand: TWA = 400 ppm (983 mg/m<sup>3</sup>); STEL = 500 ppm (1230 mg/m<sup>3</sup>), JAN 2002  
 Peru: TWA = 200 ppm (491 mg/m<sup>3</sup>); STEL = 400 ppm (983 mg/m<sup>3</sup>), JUL 2005  
 The Philippines: TWA = 400 ppm (980 mg/m<sup>3</sup>), JAN 1993  
 Poland: MAC(TWA) = 900 mg/m<sup>3</sup>, MAC(STEL) = 1200 mg/m<sup>3</sup>, JAN 1999  
 Russia: TWA = 10 mg/m<sup>3</sup>, STEL = 50 mg/m<sup>3</sup>, JUN 2003  
 Sweden: TWA = 150 ppm (350 mg/m<sup>3</sup>); STEL = 250 ppm (600 mg/m<sup>3</sup>), JUN 2005  
 Switzerland: MAK-W = 200 ppm (500 mg/m<sup>3</sup>), KZG-W = 400 ppm (1000 mg/m<sup>3</sup>), DEC 2006  
 Turkey: TWA = 200 ppm (500 mg/m<sup>3</sup>), JAN 1993  
 United Kingdom: TWA = 400 ppm (999 mg/m<sup>3</sup>); STEL = 500 ppm (1250 mg/m<sup>3</sup>), OCT 2007

#### ISOPROPYL ALCOHOL (continued):

In Argentina, Bulgaria, Colombia, Jordan, Singapore, Vietnam check ACGIH TLV  
**TITANIUM DIOXIDE:**  
 ARAB Republic of Egypt: TWA = 15 mg/m<sup>3</sup>, JAN 1993  
 Austria: MAK-TMW = 5 mg/m<sup>3</sup>, KZW = 10 mg/m<sup>3</sup>, resp, 2007  
 Belgium: TWA = 10 mg/m<sup>3</sup>, MAR 2002  
 Denmark: TWA = 6 mg(Ti)/m<sup>3</sup>, MAY 2011  
 France: VME = 10 mg/m<sup>3</sup>, FEB 2006  
 Iceland: TWA = 6 mg(Ti)/m<sup>3</sup>, NOV 2011  
 Japan: OEL = 1 mg/m<sup>3</sup> (resp. dust), 4 mg/m<sup>3</sup> (total dust), MAY 2012  
 Korea: TWA = 10 mg/m<sup>3</sup>, 2006  
 Mexico: TWA = 10 mg(Ti)/m<sup>3</sup>; STEL = 20 mg(Ti)/m<sup>3</sup>, 2004  
 The Netherlands: MAC-TGG = 10 mg/m<sup>3</sup>, 2003  
 New Zealand: TWA = 10 mg/m<sup>3</sup> (inspirable dust), JAN 2002  
 Norway: TWA = 5 mg/m<sup>3</sup>, JAN 1999  
 Peru: TWA = 10 mg/m<sup>3</sup>, JUL 2005  
 Poland: MAC(TWA) = 10 mg(Ti)/m<sup>3</sup>, MAC(STEL) = 30 mg(Ti)/m<sup>3</sup>, JAN 1999  
 Russia: TWA = 10 mg/m<sup>3</sup>, JUN 2003  
 Sweden: TWA = 5 mg/m<sup>3</sup> (total dust), JUN 2005  
 Switzerland: MAK-W = 3 mg/m<sup>3</sup>, resp, JAN2011  
 Turkey: TWA = 15 mg/m<sup>3</sup>, JAN 1993  
 United Kingdom: TWA = 10 mg/m<sup>3</sup> (inhal. dust), OCT 2007  
 United Kingdom: TWA = 4 mg/m<sup>3</sup> (resp. dust), OCT 2007  
 In Argentina, Bulgaria, Colombia, Jordan, Singapore, Vietnam check ACGIH TLV

**Biological Exposure Indices (ACGIH):** Currently, there are ACGIH Biological Exposure Indices (BEIs) determined for the components of this product, as follows:

CHEMICAL: DETERMINANT	SAMPLING TIME	BEI
Proprietary Alcohol • Acetone in urine	• End of Shift End of Workweek	• 40 mg/L

**Workplace Exposure Standards (New Zealand):** None established. Refer to the Hazardous Substances (Classes 6, 8, and 9 Controls) Regulations 2001 (Regulations 29-30).

**Exposure Standards Outside the Workplace (New Zealand):** Currently, there are no other exposure limits, such as TELS and EELS (See Section 12 [Ecological Information] for EEL information) established for components of this product.

**PERSONAL PROTECTIVE EQUIPMENT:** 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 Respiratory Protection Standard [29 CFR 1910.134]), equivalent standards of Canada (including CSA including CSA Respiratory Standard Z94.4-02 and CSA Standard Z94.3-07), standards of EU member states (including EN 529:2005 for respiratory PPE, CEN/TR 15419:2006 for hand protection, and CR 13464:1999 for face/eye protection), or standards of Australia (including AS/NZS 1715:1994 for respiratory PPE, AS/NZS 4501.2:2006 for protective clothing, AS/NZS 2161.1:2000 for glove selection, and AS/NZS 1336:1997 for eye protection), or standards of Japan (including JIS T 8116:2005 for glove selection, JIS T 8150:2006 for respiratory PPE, JIS T 8147:2003 for eye protectors, and JIS T 8030:2005 for protective clothing). Please reference applicable regulations and standards in each jurisdiction for relevant details.

**Respiratory Protection:** Respiratory protection is not generally needed when using this product. Maintain airborne contaminant concentrations below limits listed in this section, if applicable. In instances where inhalable mists or sprays of product may be generated, and respiratory protection is necessary, use only respiratory protection authorized in appropriate regulations.

## 8. EXPOSURE CONTROLS - PERSONAL PROTECTION (Continued)

### PERSONAL PROTECTIVE EQUIPMENT (continued):

**Respiratory Protection (continued):** Oxygen levels below 19.5% are considered IDLH by U.S. OSHA. In such atmospheres, use of a full-facepiece pressure/demand SCBA or a full facepiece, SAR with auxiliary self-contained air supply is required under OSHA's Respiratory Protection Standard (1910.134-1998).

**Eye Protection:** Depending on the use of this product, splash goggles or safety glasses may be worn. Use goggles or safety glasses for spill response, as stated in Section 6 (Accidental Release Measures) of this SDS. If necessary, refer to appropriate regulations when selecting eye protection.

**Hand Protection:** Wear butyl rubber, neoprene, or nitrile rubber or latex gloves for routine use. If necessary, refer to appropriate regulations for further information.

**Body Protection:** Use body protection appropriate for task, such as a lab coat. If necessary, use body protection appropriate for task (e.g., Tyvek suit, rubber apron). 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.

## 9. PHYSICAL and CHEMICAL PROPERTIES

**FORM:** Liquid.

**MOLECULAR FORMULA:** Mixture.

**ODOR:** Mild.

**VAPOR DENSITY (air = 1):** Not established.

**SPECIFIC GRAVITY (water = 1):** Not established.

**SOLUBILITY IN WATER:** Soluble

**VAPOR PRESSURE:** Not established.

**OXIDIZING PROPERTIES:** Not applicable.

**COEFFICIENT OF OIL/WATER DISTRIBUTION (PARTITION COEFFICIENT):** Not established.

**HOW TO DETECT THIS SUBSTANCE (identification properties):** The odor and color of this product may be distinguishing characteristics to identify it event of a spill.

**COLOR:** White.

**MOLECULAR WEIGHT:** Mixture.

**ODOR THRESHOLD:** Not established.

**EVAPORATION RATE (n-BuAc = 1):** < 1

**MELTING/FREEZING POINT:** Not established.

**BOILING POINT:** Not established.

**pH:** Not established.

**EXPLOSIVE PROPERTIES:** Not applicable.

## 10. STABILITY and REACTIVITY

**CHEMICAL STABILITY:** Stable under conditions of normal temperature and pressure.

**DECOMPOSITION PRODUCTS:** **Combustion:** If exposed to extremely high temperatures, this product can decompose to generate calcium, carbon, nitrogen and nitrogen oxides. **Hydrolysis:** None known.

**MATERIALS WITH WHICH SUBSTANCE IS INCOMPATIBLE:** Strong oxidizers, water-reactive materials.

**POSSIBILITY OF HAZARDOUS REACTION OR POLYMERIZATION:** Will not occur.

**CONDITIONS TO AVOID:** Exposure to or contact with extreme temperatures and incompatible chemicals.

## 11. TOXICOLOGICAL INFORMATION

**SYMPTOMS OF EXPOSURE BY ROUTE OF EXPOSURE:** The most significant routes of occupational exposure are inhalation and contact with skin and eyes. The symptoms of exposure to this material, via route of entry, are as described below.

**Inhalation:** This product does not normally present a significant inhalation hazard under anticipated circumstances of use. Inhalation of vapors, mists, or sprays of this material, may mildly irritate the nose, throat, and other tissues of the respiratory system.

**Contact with Skin or Eyes:** Skin contact may cause mild irritation in sensitive individuals. Repeated or prolonged skin exposure may cause dermatitis (dry, red skin). Eye contact with this material can moderately irritate the eyes, causing discomfort, tearing, and redness. Because the eye tissue may be stained, vision may be temporarily blurred.

**Skin Absorption:** No component is known to be absorbed via intact skin.

**Ingestion:** Though not anticipated to be a significant route of occupational exposure, ingestion of large quantities of this material may cause nausea, vomiting, diarrhea, and discoloration of the mouth, teeth, and tissues of the throat.

**Injection:** Accidental injection of this liquid (as may occur by a puncture with a contaminated object) will cause local pain, irritation, and redness.

**HEALTH EFFECTS OR RISKS FROM EXPOSURE:** In the event of exposure, the following symptoms may be observed:

**Acute:** The ink may stain hair, skin, and other contaminated tissue. Eye contact will cause moderate irritation. Ingestion of large amounts may cause nausea, vomiting, diarrhea.

**Chronic:** Repeated or prolonged skin exposure may cause dermatitis (dry, red skin).

**TARGET ORGANS:** **Acute:** Skin, central nervous system, eyes. **Chronic:** Skin.

**TOXICITY DATA:** Specific toxicology data currently available for components of this product in greater than 1% concentration are as follows.

### HYDROXYETHYL CELLULOSE:

LDLo (Intravenous-Woman) 5100 mg/kg/6 days-intermittent: Brain and Coverings: increased intracranial pressure; Behavioral: somnolence (general depressed activity); Vascular: change in plasma or blood volume

TDLo (Intraperitoneal-Mouse) 500 mg/kg: female 3-7 day(s) after conception: Reproductive: Fertility: post-implantation mortality (e.g. dead and/or resorbed implants per total number of implants)

### HAZARDOUS MATERIAL IDENTIFICATION SYSTEM

<b>HEALTH HAZARD</b>	(BLUE)	1
----------------------	--------	---

<b>FLAMMABILITY HAZARD</b>	(RED)	0
----------------------------	-------	---

<b>PHYSICAL HAZARD</b>	(YELLOW)	0
------------------------	----------	---

### PROTECTIVE EQUIPMENT

EYES	RESPIRATORY	HANDS	BODY
	SEE SECTION 8		SEE SECTION 8

For Routine Industrial Use and Handling Applications

Hazard Scale: 0 = Minimal 1 = Slight 2 = Moderate  
3 = Serious 4 = Severe \* = Chronic hazard

# 11. TOXICOLOGICAL INFORMATION (Continued)

## TOXICITY DATA (continued):

### ISOPROPYL ALCOHOL:

LDLo (oral, man) = 5272 mg/kg; Behavioral: coma; Vascular: BP lowering not characterized in autonomic section; Lungs, Thorax, or Respiration: chronic pulmonary edema  
LDLo (oral, human) = 3570 mg/kg; Behavioral: coma; Lungs, Thorax, or Respiration: respiratory depression; Gastrointestinal: nausea or vomiting  
TDLo (oral, man) = 14,432 mg/kg; Behavioral: coma; Vascular: BP lowering not characterized in autonomic section; Lungs, Thorax, or Respiration: dyspnea  
TDLo (oral, human) = 223 mg/kg; Behavioral: hallucinations, distorted perceptions; Cardiac: pulse rate; Vascular: BP lowering not characterized in autonomic section  
TDLo (oral, infant) = 13 gm/kg; Behavioral: somnolence (general depressed activity), irritability; Gastrointestinal: nausea or vomiting  
LDLo (unreported, man) = 2770 mg/kg  
Skin Irritancy (rabbit) = 500 mg; mild  
Eye Irritancy (rabbit) = 100 mg; severe  
Eye Irritancy (rabbit) = 16 mg  
Eye Irritancy (rabbit) = 10 mg; moderate  
LD<sub>50</sub> (oral, rat) = 5045 mg/kg  
LD<sub>50</sub> (oral, mouse) = 3600 mg/kg  
LD<sub>50</sub> (oral, rabbit) = 6410 mg/kg  
LD<sub>50</sub> (skin, rabbit) = 12,800 mg/kg  
LD<sub>50</sub> (intravenous, rat) = 1099 mg/kg  
LD<sub>50</sub> (intravenous, mouse) = 1509 mg/kg  
LD<sub>50</sub> (intravenous, rabbit) = 1184 mg/kg  
LD<sub>50</sub> (intraperitoneal, rat) = 2735 mg/kg  
LD<sub>50</sub> (intraperitoneal, mouse) = 4477 mg/kg  
LD<sub>50</sub> (intraperitoneal, rabbit) = 667 mg/kg  
LD<sub>50</sub> (intraperitoneal, guinea pig) = 2560 mg/kg  
LD<sub>50</sub> (intraperitoneal, hamster) = 3444 mg/kg  
LDLo (oral, dog) = 1537 mg/kg; Gastrointestinal: nausea or vomiting  
LDLo (oral, cat) = 6 mL/kg  
LCLo (inhalation, rat) = 16,000 ppm/4 hours  
LCLo (inhalation, mouse) = 12,800 ppm/3 hours  
LDLo (intravenous, cat) = 1963 mg/kg  
LDLo (parenteral, frog) = 20 g/kg; Peripheral Nerve and Sensation: spastic paralysis with or without sensory change; Behavioral: somnolence (general depressed activity)  
LDLo (subcutaneous, mouse) = 6000 mg/kg  
LDLo (intravenous, dog) = 5120 mg/kg  
TDLo (oral, rat) = 6480 mg/kg/male 26 weeks pre; Reproductive effects  
TCLo (inhalation, rat) = 10,000 ppm/7 hours/female 1–19 days post; Teratogenic effects  
Cytogenetic Analysis (*Saccharomyces cerevisiae*) = 200 mmol/tube  
Cytogenetic Analysis (inhalation, rat) = 1030 µg/m<sup>3</sup>/16 weeks/intermittent

### TITANIUM DIOXIDE:

Standard Draize Test (Skin-Human) 300 µg/3 days-intermittent: Mild  
TC (Inhalation-Rat) 10 mg/m<sup>3</sup>/18 hours/2 years-intermittent: Tumorigenic: carcinogenic by RTECS criteria; Lungs, Thorax, or Respiration: tumors  
LD (Intratracheal-Rat) > 100 µg/kg; Lungs, Thorax, or Respiration: structural or functional change in trachea or bronchi; Blood: changes in serum composition (e.g. TP, bilirubin, cholesterol); Biochemical: Enzyme inhibition, induction, or change in blood or tissue levels: other Enzymes  
TD (Intramuscular-Rat) 260 mg/kg/84 weeks-intermittent: Tumorigenic: equivocal tumorigenic agent by RTECS criteria; Blood: lymphoma, including Hodgkin's disease; Tumorigenic: tumors at site of application  
TDLo (Oral-Rat) 60 mg/kg; Gastrointestinal: hypermotility, diarrhea, other changes  
TDLo (Intramuscular-Rat) 360 mg/kg/2 years-intermittent: Tumorigenic: neoplastic by RTECS criteria; Blood: lymphoma, including Hodgkin's disease; Tumorigenic: tumors at site of application  
TDLo (Intratracheal-Rat) 1.25 mg/kg; Vascular: regional or general arteriolar constriction; Lungs, Thorax, or Respiration: other changes

### TITANIUM DIOXIDE (continued):

TDLo (Intratracheal-Rat) 1.6 mg/kg; Lungs, Thorax, or Respiration: other changes  
TDLo (Intratracheal-Rat) 5 mg/kg; Lungs, Thorax, or Respiration: other changes; Biochemical: Metabolism (Intermediary): effect on inflammation or mediation of inflammation  
TDLo (Intratracheal-Mouse) 100 mg/kg; Tumorigenic: increased incidence of tumors in susceptible strains  
TCLo (Inhalation-Rat) 1 mg/kg; Lungs, Thorax, or Respiration: other changes; Biochemical: Metabolism (Intermediary): effect on inflammation or mediation of inflammation  
TCLo (Inhalation-Rat) 250 mg/m<sup>3</sup>/6 hours/4 weeks-intermittent: Lungs, Thorax, or Respiration: chronic pulmonary edema, other changes  
TCLo (Inhalation-Rat) 50 mg/m<sup>3</sup>/6 hours/13 weeks-intermittent: Lungs, Thorax, or Respiration: structural or functional change in trachea or bronchi  
TCLo (Inhalation-Rat) 10 mg/m<sup>3</sup>/6 hours/13 weeks-intermittent: Lungs, Thorax, or Respiration: fibrosis (interstitial), other changes; Biochemical: Metabolism (Intermediary): effect on inflammation or mediation of inflammation  
TCLo (Inhalation-Rat) 10 mg/m<sup>3</sup>/13 weeks-intermittent: Lungs, Thorax, or Respiration: other changes; Biochemical: Metabolism (Intermediary): effect on inflammation or mediation of inflammation  
TCLo (Inhalation-Rat) 50 mg/m<sup>3</sup>/13 weeks-intermittent: Lungs, Thorax, or Respiration: sputum; Blood: changes in cell count (unspecified); Biochemical: Enzyme inhibition, induction, or change in blood or tissue levels: dehydrogenases  
TCLo (Inhalation-Rat) 250 mg/m<sup>3</sup>/13 weeks-intermittent: Lungs, Thorax, or Respiration: other changes; Blood: changes in cell count (unspecified); Biochemical: Enzyme inhibition, induction, or change in blood or tissue levels: dehydrogenases  
TCLo (Inhalation-Rat) 274 mg/m<sup>3</sup>/5 days-intermittent: Lungs, Thorax, or Respiration: changes in lung weight; Biochemical: Enzyme inhibition, induction, or change in blood or tissue levels: multiple enzyme effects, Metabolism (Intermediary): effect on inflammation or mediation of inflammation  
TCLo (Inhalation-Rat) 250 mg/m<sup>3</sup>/6 hours/2 years-intermittent: Tumorigenic: carcinogenic by RTECS criteria; Lungs, Thorax, or Respiration: tumors  
TCLo (Inhalation-Mouse) 10 mg/m<sup>3</sup>/6 hours/13 weeks-intermittent: Lungs, Thorax, or Respiration: other changes; Biochemical: Metabolism (Intermediary): effect on inflammation or mediation of inflammation  
TCLo (Inhalation-Mouse) 10 mg/m<sup>3</sup>/6 hours/13 weeks-intermittent: Lungs, Thorax, or Respiration: structural or functional change in trachea or bronchi  
TCLo (Inhalation-Mouse) 10 mg/m<sup>3</sup>/13 weeks-intermittent: Lungs, Thorax, or Respiration: other changes; Biochemical: Metabolism (Intermediary): effect on inflammation or mediation of inflammation  
TCLo (Inhalation-Mouse) 50 mg/m<sup>3</sup>/13 weeks-intermittent: Lungs, Thorax, or Respiration: sputum; Biochemical: Enzyme inhibition, induction, or change in blood or tissue levels: dehydrogenases  
TCLo (Inhalation-Mouse) 250 mg/m<sup>3</sup>/13 weeks-intermittent: Lungs, Thorax, or Respiration: sputum; Blood: changes in cell count (unspecified); Biochemical: Enzyme inhibition, induction, or change in blood or tissue levels: dehydrogenases  
TCLo (Inhalation-Hamster) 250 mg/m<sup>3</sup>/13 weeks-intermittent: Lungs, Thorax, or Respiration: sputum; Blood: changes in cell count (unspecified); Biochemical: Enzyme inhibition, induction, or change in blood or tissue levels: dehydrogenases  
TCLo (Inhalation-Hamster) 250 mg/m<sup>3</sup>/6 hours/13 weeks-intermittent: Lungs, Thorax, or Respiration: structural or functional change in trachea or bronchi  
DNA Damage (Human Lung) 100 µg/plate  
DNA Damage (Human Lung) 20 µg/disk/4 hours  
Sister Chromatid Exchange (Human Lymphocyte) 2 µmol/L/72 hours  
Micronucleus Test (Human Lymphocyte) 5 µmol/L/72 hours  
Micronucleus Test (Intraperitoneal-Mouse) 3 gm/kg/3 days-continuous  
Micronucleus Test (Hamster Ovary) 5 µmol/L  
DNA Inhibition (Hamster Lung) 500 mg/L  
Sister Chromatid Exchange (Hamster Ovary) 1 µmol/L

**CARCINOGENIC POTENTIAL OF COMPONENTS:** Components of this product are listed by agencies tracking the carcinogenic potential of chemical compounds, as follows;

**ISOPROPYL ALCOHOL:** ACGIH TLV-A4 (Not Classifiable as a Human Carcinogen); IARC-3 (Not Classifiable as to Carcinogenicity to Humans)

**TITANIUM DIOXIDE:** ACGIH TLV-A4 (Not Classifiable as a Human Carcinogen); IARC-2B (Possibly Carcinogenic to Humans); MAK-3A (Substances for Which the Criteria for Classification in Category 4 or 5 are fulfilled but for which the database is insufficient for the establishment of a MAK value); NIOSH-Ca (Potential Occupational Carcinogen with No Further Categorization)

The remaining components of this product are not found on the following lists: U.S. EPA, U.S. NTP, U.S. OSHA, U.S. NIOSH, GERMAN MAK, IARC, and ACGIH, and therefore are neither considered to be nor suspected to be cancer-causing agents by these agencies.

**IRRITANCY OF PRODUCT:** Acute exposure to this material via skin contact, eye contact, and inhalation may mildly irritate contaminated tissue.

**SENSITIZATION TO THE PRODUCT:** The components of this product are not known to be human skin or respiratory sensitizers.

**REPRODUCTIVE TOXICITY INFORMATION:** Listed below is information concerning the effects of components of this product on the human reproductive system.

**Mutagenicity:** No data are available on possible mutagenic effects from this product and components.

**Embryotoxicity:** The components of this product are not reported to produce embryotoxic effects in humans. The Proprietary Alcohol component has produced fetotoxicity (reduced fetal weight) in rats exposed by inhalation, in the absence of maternal toxicity. Reduced survival in the early postnatal period has been observed in the offspring of rats exposed to high oral doses, in the presence of minimal maternal toxicity. Rats were exposed by inhalation to 0, 3500, 7000 or 10000 ppm during days 1-19 of pregnancy. Maternal toxicity was observed at the 2 high doses, but not at 3500 ppm. Fetal weights were significantly reduced in a concentration related manner at all treatment levels. At 7000 and 10000 ppm, teratogenicity and/or embryotoxicity were observed.

**Teratogenicity:** No data are available on possible teratogenic effects from this product and components.

## 11. TOXICOLOGICAL INFORMATION (Continued)

### REPRODUCTIVE TOXICITY INFORMATION (continued):

**Reproductive Toxicity:** The components of this product are not reported to cause reproductive effects in humans. In a two-generation study of Proprietary Alcohol, rats were orally dosed with 0, 100, 500 or 1000 mg/kg/day for 10 weeks prior to mating. Females were dosed during mating, gestation and lactation and males were dosed during mating through delivery of the last litter sired. In the first generation, a significant reduction was observed in the live birth index and the survival index on days 1 and 4 for the offspring of animals exposed to 1000 mg/kg/day, as well as the survival rate of off-spring. Only minimal maternal toxicity (increased liver weight) was observed at 500 mg/kg/day. At 1000 mg/kg/day, 2/30 females in the first generation (P1) and 2/26 females died in the second generation.

## 12. ECOLOGICAL INFORMATION

ALL WORK PRACTICES MUST BE AIMED AT ELIMINATING ENVIRONMENTAL CONTAMINATION.

**MOBILITY:** This product has not been tested for mobility in soil. It is expected to be somewhat mobile in soil. The following information is available for the Proprietary Alcohol component.

**PROPRIETARY ALCOHOL:** The Koc of Proprietary Alcohol is estimated as 25, using a measured log Kow of 0.05 and a regression-derived equation. According to a classification scheme, this estimated Koc value suggests that Proprietary Alcohol is expected to have very high mobility in soil.

**PERSISTENCE AND BIODEGRADABILITY:** This product has not been tested for persistence or biodegradability. It is expected that some biodegradation will occur to this product; however, no specific information is known. The following information is available for the Proprietary Alcohol component.

**PROPRIETARY ALCOHOL:** Based on a classification scheme, an estimated Koc value of 25, determined from a log Kow of 0.05 and a regression-derived equation, indicates that Proprietary Alcohol is expected to have very high mobility in soil. Volatilization of Proprietary Alcohol from moist soil surfaces is expected to be an important fate process given a Henry's Law constant of  $8.10 \times 10^{-6}$  atm-cu m/mole. The potential for volatilization of Proprietary Alcohol from dry soil surfaces may exist based upon a vapor pressure of 45.4 mmHg. Proprietary Alcohol is readily degraded in aerobic systems; the range of half-lives for aerobic degradation using a sewage sludge inoculum are < 1 day to 48 days. Proprietary Alcohol has also been shown to be readily degraded under anaerobic conditions. Volatilization from water surfaces is expected based upon a Henry's Law constant of  $8.10 \times 10^{-6}$  atm-cu m/mole. Using this Henry's Law constant and an estimation method, volatilization half-lives for a model river and model lake are 57 hours and 29 days, respectively. Proprietary Alcohol is readily degraded in aerobic systems; the range of half-lives for aerobic degradation using a sewage sludge inoculum are < 1 day to 48 days. Proprietary Alcohol has also been shown to be readily degraded under anaerobic conditions. According to a model of gas/particle partitioning of semi-volatile organic compounds in the atmosphere, Proprietary Alcohol, which has a vapor pressure of 45.4 mm Hg at 25°C, is expected to exist solely as a vapor in the ambient atmosphere. Vapor-phase Proprietary Alcohol is degraded in the atmosphere by reaction with photochemically-produced hydroxyl radicals; the half-life for this reaction in air is estimated to be 3.2 days, calculated from its rate constant of  $5.07 \times 10^{-12}$  cu cm/molecule-sec at 25°C.

**BIO-ACCUMULATION POTENTIAL:** This product has not been tested for bio-accumulation potential. The following information is available for the Proprietary Alcohol component.

**PROPRIETARY ALCOHOL:** An estimated BCF of 3 was calculated for Proprietary Alcohol, using a log Kow of 0.05 and a regression-derived equation. According to a classification scheme, this BCF suggests the potential for bioconcentration in aquatic organisms is low.

**ECOTOXICITY:** This product has not been tested for the effects of this product on terrestrial or aquatic organisms if released to the environment. Plants may be discolored and damaged (depending on the severity of the contamination). The following aquatic toxicity data are available for the Proprietary Alcohol component (not all available data are given in the SDS. Contact Silverback Ink for information on other available data).

**PROPRIETARY ALCOHOL:**

EC<sub>50</sub> (*Daphnia magna*) 3,010 mg/L  
LC<sub>50</sub> (*Artemia salina*) 24 hours = 16,700 mg/L  
LC<sub>50</sub> (*Streptocephalus proboscideus*) 24 hours = 11,600 mg/L  
LC<sub>50</sub> (*Daphnia magna*) 24 hours = 9,500 mg/L

**PROPRIETARY ALCOHOL (continued):**

LC<sub>50</sub> (*Brachionus calyciflorus*) 24 hours = 28,600 mg/L  
LC<sub>50</sub> (*Crangon crangon* brown shrimp) 98 hours = 1,150 mg/L  
LC<sub>50</sub> (goldfish) 24 hours = > 500 mg/L  
LC<sub>50</sub> (fathead minnow) 96 hours = 11,130 mg/L

**RESULTS OF PBT AND vPvB ASSESSMENT:** No data available. PBT and vPvB assessments are part of the chemical safety report required for some substances in European Union Regulation (EC) 1907/2006, Article 14.

**OTHER ADVERSE EFFECTS:** This product does not contain any constituents with known ozone depletion potential.

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

## 13. DISPOSAL CONSIDERATIONS

**DISPOSAL METHODS:** It is the responsibility of the generator to determine at the time of disposal whether the product meets the criteria of a hazardous waste per regulations of the area in which the waste is generated and/or disposed of. Waste disposal must be in accordance with appropriate Federal, State, and local regulations. This product, if unaltered by use, may be disposed of by treatment at a permitted facility or as advised by your local hazardous waste regulatory authority. Shipment of wastes must be done with appropriately permitted and registered transporters.

**DISPOSAL CONTAINERS:** Waste materials must be placed in and shipped in appropriate 5-gallon or 55-gallon poly or metal waste pails or drums. Permeable cardboard containers are not appropriate and should not be used. Ensure that any required marking or labeling of the containers be done to all applicable regulations.

**PRECAUTIONS TO BE FOLLOWED DURING WASTE HANDLING:** Wear proper protective equipment when handling waste materials.

**U.S. EPA WASTE NUMBER:** Not applicable to wastes consisting only of this product.

**EUROPEAN WASTE CODES:** Wastes from MFSU and Removal of Printing Inks: 08 03 99: Wastes Not Otherwise Specified

## 14. TRANSPORTATION INFORMATION

**U.S. DEPARTMENT OF TRANSPORTATION:** This product is not classified as dangerous goods, per U.S. DOT regulations, under 49 CFR 172.101.

**TRANSPORT CANADA, TRANSPORTATION OF DANGEROUS GOODS REGULATIONS:** This product is NOT classified as dangerous goods, per regulations of Transport Canada.

**INTERNATIONAL AIR TRANSPORT ASSOCIATION SHIPPING INFORMATION (IATA):** This product is NOT classified as dangerous goods.

## 14. TRANSPORTATION INFORMATION (Continued)

INTERNATIONAL MARITIME ORGANIZATION SHIPPING INFORMATION (IMO): This product is NOT classified as dangerous goods.

EUROPEAN AGREEMENT CONCERNING THE INTERNATIONAL CARRIAGE OF DANGEROUS GOODS BY ROAD (ADR): This product is NOT classified by the United Nations Economic Commission for Europe to be dangerous goods.

AUSTRALIAN FEDERAL OFFICE OF ROAD SAFETY CODE FOR THE TRANSPORTATION OF DANGEROUS GOODS BY ROAD OR RAIL: This product is NOT classified as dangerous goods, per regulations of the Australian Federal Office of Road Safety.

TRANSPORT IN BULK ACCORDING TO ANNEX II OF MARPOL 73/78 AND THE IBC CODE: Not applicable.

ENVIRONMENTAL HAZARDS: This product is neither environmentally hazardous according to the criteria of the UN Model Regulations (as reflected in the IMDG Code, ADR, RID, and ADN); no component meets the criteria of environmentally hazardous.

## 15. REGULATORY INFORMATION

### UNITED STATES REGULATIONS:

U.S. SARA Reporting Requirements: The 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 as follows:

CHEMICAL NAME	SARA 302 (40 CFR 355, Appendix A)	SARA 304 (40 CFR Table 302.4)	SARA 313 (40 CFR 372.65)
Proprietary Alcohol (mfg-strong acid process)	No	No	Yes

U.S. SARA Threshold Planning Quantity (TPQ): There are no specific Threshold Planning Quantities for this material. The default Federal SDS submission and inventory requirement filing threshold of 10,000 lb (4,540 kg) may apply, per 40 CFR 370.20.

U.S. CERCLA Reportable Quantity (RQ): Not applicable.

U.S. TSCA Inventory Status: The components of this product are listed on the TSCA Inventory.

U.S. Hazardous Air Pollutant (HAPs): The components of this product are not listed by the EPA under section 112(b) of the Clean Air Act as a 'HAP'.

Other U.S. Federal Regulations: Not applicable.

California Safe Drinking Water And Toxic Enforcement Act (Proposition 65): The Titanium Dioxide component, (airborne, unbound particles of respirable size) is on the California Proposition 65 Lists. If airborne particles of this product are generated, the following warning must be on packaging and labeling of this product: WARNING! This product contains a component known to the State of California to cause cancer. As a liquid, this warning is not expected to be required.

### CANADIAN REGULATIONS:

Canadian DSL/NDSL Inventory Status: The components of this product are listed on the DSL Inventory.

Canadian Environmental Protection Act (CEPA) Priority Substances Lists: The Proprietary Alcohol component is listed as a Substance With Greatest Potential For Human Exposure Substance on Environment Canada/Health Canada Pilot Project List (CEPA 1999, Section 73). Meets categorization criteria: \*may present, to individuals in Canada, the greatest potential for exposure; or \*are persistent or bioaccumulative in accordance with the regulations, and inherently toxic to human beings or to non-human organisms, as determined by laboratory or other studies.

Canadian WHMIS HPR 2015 Classification and Symbols: See the following section for classification and symbols under WHMIS.

### EUROPEAN UNION REGULATIONS:

Safety, Health, and Environmental Regulations/Legislation Specific for the Product: Currently, there is no specific legislation pertaining to this product.

Chemical Safety Assessment: No data available. The chemical safety assessment is required for some substances according to European Union Regulation (EC) 1907/2006, Article 14.

### AUSTRALIAN REGULATIONS:

Australian Inventory Of Chemical Substances (AICS) Status: The components of this product are listed on the AICS.

Hazardous Substances Information System (HSIS): The components of this product are not listed in the HSIS.

Standard for the Uniform Scheduling of Drugs and Poisons: Not applicable.

### CHINESE REGULATIONS:

Chinese Inventory of Existing Chemical Substances Status: Components are listed on the Chinese Inventory of Existing Chemical Substances (IECSC).

### JAPANESE REGULATIONS:

Japanese ENCS Inventory: Components are not on the ENCS Inventory.

Japanese Ministry of Economy, Trade, and Industry (METI) Status: Components are not listed as Class I Specified Chemical Substances, Class II Specified Chemical Substances, or Designated Chemical Substances by the Japanese METI.

Poisonous and Deleterious Substances Control Law: Components are not listed as a Specified Poisonous Substance under the Poisonous and Deleterious Substances Control Law.

### KOREAN REGULATIONS:

Korean Existing Chemical Substances Inventory Status: Components are listed on the Korean Existing Chemicals List, as indicated in composition tables in Section 3 (Composition and Information on Ingredients).

### NEW ZEALAND REGULATIONS:

New Zealand Inventory of Chemicals (NZIoC): The components of this product are on the NZIoC.

### MEXICAN REGULATIONS:

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

### TAIWANESE REGULATIONS:

Taiwan Existing Chemical Substances Inventory Status: Components are listed on the Taiwan Existing Chemicals List.

## 16. OTHER INFORMATION

**ANSI LABELING (Z129.1): CAUTION! MAY CAUSE SERIOUS EYE IRRITATION. MAY CAUSE SKIN AND RESPIRATORY TRACT IRRITATION. MAY DISCOLOR CONTAMINATED SKIN, EYES, HAIR, AND CLOTHES.** Use with adequate ventilation. Avoid contact of liquid with skin, eyes, and clothing. Avoid exposure to vapors, mists, or sprays. Wash thoroughly after handling. Wear appropriate hand and eye protection. **FIRST-AID:** In case of contact, immediately flush skin with plenty of water. Remove contaminated clothing and shoes. If inhaled, remove to fresh air. If swallowed, do not induce vomiting. Get medical attention if irritation develops or persists or if any other adverse effect occurs. **IN CASE OF FIRE:** Use water fog, dry chemical, or CO<sub>2</sub>, or alcohol foam. **IN CASE OF SPILL:** Absorb spill with inert materials (e.g., polypads, dry sand). Rinse area with soapy water. Consult Safety Data Sheet for additional information.

**GLOBAL HARMONIZATION LABELING AND CLASSIFICATION:** Classified in accordance with Global Harmonization Standard under U.S. OSHA Hazard Communication Standard, Canadian WHMIS HPR-GHS 2-15, EU CLP Regulation (EC) 1272/2008, Japanese JIS Z7253: 2012 and Singapore Standards. For information on Korean ISHA and New Zealand HSN0 classification, see below.

**Classification:** Eye Irritation, Category 2A

**Hazard Statements:** H319: Causes serious eye irritation.

**Precautionary Statements:**

**Prevention:** P264: Wash thoroughly after handling. P280: Wear protective gloves/protective clothing/eye protection/face protection.

**Response:** P305 + P351 + P338: IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing. P337 + P313: If eye irritation persists: get medical advice/attention.

**Storage:** None

**Disposal:** None

**Signal Word:** Warning

**Hazard Symbols/Pictograms:** GHS07

**KOREAN ISHA (Notice 2009-68) LABELING AND CLASSIFICATION:** Classified in accordance with ISHA Notice 2009-68. Under ISHA, the following differences in classification are applicable. Refer to information given under the Global Harmonization Standard Classification.

**Classification:** Eye Irritation, Category 2

**Hazard Statement Codes:** H319: Causes serious eye irritation.

**Precautionary Statements:**

**Prevention:** P264: Wash thoroughly after handling. P280: Wear protective gloves/protective clothing/eye protection/face protection.

**Response:** P305 + P351 + P338: IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing. P337 + P313: If eye irritation persists: get medical advice/attention.

**Storage:** None

**Disposal:** None

**Signal Word:** Warning

**Hazard Symbols/Pictograms:** GHS07

**NEW ZEALAND HAZARDOUS SUBSTANCES and NEW ORGANISMS ACT (HNSO) CHEMICAL CLASSIFICATION:**

The product is classified as follows under the regulation:

**Classification:**

6.4A: Irritation to the eye.

**CLASSIFICATION INFORMATION FOR COMPONENTS:**

**CLP Regulation (EC) 1272/2008**

**Proprietary Alcohol:**

**Classification:** Flammable Liquid Category 2, Eye Irritant Category 2A, Specific Target Organ Toxicity (Inhalation-Central Nervous System) Single Exposure Category 3

**Hazard Statements:** H225: Highly flammable liquid and vapour. H319: Causes serious eye irritation. H336: May cause drowsiness or dizziness.

**Titanium Dioxide:**

**Classification:** Carcinogenic Category 2

**Hazard Statements:** H350i: May cause cancer by inhalation.

**All Remaining Components:**

**Classification:** An official classification for this substance has not been published under CLP 1272/2008 and a self-classification is not applicable.

**NEW ZEALAND HAZARDOUS SUBSTANCES and NEW ORGANISMS ACT (HNSO) CHEMICAL CLASSIFICATION:**

**Carbon Black:**

**Classification:** 6.3B: Mildly irritating to the skin. 6.4A: Irritating to the eye. 6.7B: Suspected human carcinogen.

**Dehydroacetic Acid:**

**Classification:** 6.1D (oral): Substances that are acutely toxic. 9.3C: Harmful to terrestrial vertebrates.

**Proprietary Alcohol:**

**Classification:** 3.1B: Flammable liquids: high hazard. 6.1E (oral): Substances that are acutely toxic. 6.3C: Mildly irritating to the skin. 6.4A: Irritating to the eye.

**Phenoxyethanol:**

**Classification:** 6.1D (oral, dermal, inhalation): Substances that are acutely toxic. 6.4A: Irritating to the eye. 6.8B: Suspected human reproductive or developmental toxicants. 9.3C: Harmful to terrestrial vertebrates.

**REVISION DETAILS:** September 2016: Up-date of entire SDS to include additional country classification. Review of SDS and up-date as applicable all exposure limits, toxicity data and other information.

**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.

**PREPARED BY:** CHEMICAL SAFETY ASSOCIATES, Inc., PO Box 1961, Hilo, HI 96721, 800/441-3365

**DATE OF PRINTING:** October 11, 2016

The data in this Safety Data Sheet is true and accurate to the best of Silverback Ink knowledge. However, since data, safety standards, and government regulations are subject to change conditions of handling, use, or misuse are beyond Silverback Ink control, Silverback Ink MAKES NO WARRANTY, EITHER EXPRESSED OR IMPLIED, WITH RESPECT TO THE COMPLETENESS OR CONTINUING ACCURACY OF THE INFORMATION CONTAINED HEREIN AND DISCLAIMS ALL LIABILITY FOR RELIANCE THEREON. The user is required to comply with all laws and regulations relating to the purchase, use, storage, and disposal of the product. User must be familiar with and follow generally accepted safe handling procedures of chemicals, and is solely responsible for any effects caused by its misuse or mixing of this chemical with any other substance.

# DEFINITIONS OF TERMS

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

**CAS #:** This is the Chemical Abstract Service Number that uniquely identifies each constituent.

## EXPOSURE LIMITS IN AIR:

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

**DFG MAK Germ Cell Mutagen Categories:** 1: Germ cell mutagens which have been shown to increase the mutant frequency in the progeny of exposed humans. 2: Germ cell mutagens which have been shown to increase the mutant frequency in the progeny of exposed mammals. 3A: Substances which 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 which 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 which 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 cause 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. **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.

**LOQ:** Limit of Quantitation.

**MAK:** Federal Republic of Germany Maximum Concentration Values in the workplace.

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

**NIC:** Notice of Intended Change.

**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.

**NIOSH RELS:** NIOSH's Recommended Exposure Limits.

**PEL-Permissible Exposure Limit:** 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 there is a danger of cutaneous absorption.

**STEL-Short Term Exposure Limit:** 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:** 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.

## 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 HAZARDS: 0 (Minimal Hazard):** No significant health risk, irritation of skin or eyes not anticipated. *Skin Irritation:* Essentially non-irritating. PII or Draize = "0". *Eye Irritation:* Essentially non-irritating, or minimal effects which clear in < 24 hours [e.g. mechanical irritation]. 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; slightly or mildly irritating. *Skin Irritation:* Slightly or mildly irritating. *Eye Irritation:* Slightly or mildly irritating. *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> 4-hrs Rat.* > 2-20 mg/L; 2 (Moderate Hazard): Temporary or transitory injury may occur. *Skin Irritation:* Moderately irritating; primary irritant; sensitizer. PII or Draize > 0, < 5. *Eye Irritation:* Moderately to severely irritating and/or corrosive; reversible corneal opacity; corneal involvement or irritation clearing in 8-21 days. Draize > 0, < 25. *Oral Toxicity LD<sub>50</sub> Rat.* > 500-500 mg/kg. *Dermal Toxicity LD<sub>50</sub>Rat or Rabbit.* > 200-1000 mg/kg. 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 destroy dermal tissue, cause skin burns, 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<sub>50</sub> Rat.* > 1-50 mg/kg. *Dermal Toxicity LD<sub>50</sub>Rat or Rabbit.* > 20-200 mg/kg. *Inhalation Toxicity LC<sub>50</sub> 4-hrs Rat.* > 0.05-0.5 mg/L; 4 (Severe Hazard): Life-threatening; major or permanent damage may result from single or repeated exposure. *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<sub>50</sub> Rat.* ≤ 1 mg/kg. *Dermal Toxicity LD<sub>50</sub>Rat or Rabbit.* ≤ 20 mg/kg. *Inhalation Toxicity LC<sub>50</sub> 4-hrs Rat.* ≤ 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 require considerable pre-heating, under all ambient temperature conditions before ignition and combustion can occur. Including: Materials that will burn in air when exposed to a temperature of 815.5°C (1500°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] (e.g. OSHA Class IIIB, or; 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 in air, including: Liquids having a flash-point at or above 37.8°C [100°F]; Solid materials in the form of coarse 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; Solids and semisolids 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, including: Liquids having a flash point below 22.8°C [73°F] and having a boiling point at or above 38°C [100°F] and below 37.8°C [100°F] [e.g. OSHA Class IB and IC];**

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

**FLAMMABILITY HAZARD (continued): 3 (Serious Hazard) [CONTINUED]:** 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]; 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 which will burn readily, including: 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] [e.g. OSHA Class IA; Material that ignite spontaneously when exposed to air at a temperature of 54.4°C [130°F] or below [e.g. 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. *Unstable Compressed Gases:* No Rating. *Pyrophorics:* No Rating. *Oxidizers:* No "0" rating allowed. *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. *Explosives:* Division 1.5 & 1.6 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; *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 explosive 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 – Explosive substances where the explosive effect 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:* Packaging Group II; *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. *Unstable Reactives:* Substances that may polymerize, decompose, condense, or self-react at ambient temperature and/or pressure, but have a low potential 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 1.2 – 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:* Packaging Group I; *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 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-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. *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 to cause significant heat generation or explosion.).

## NATIONAL FIRE PROTECTION ASSOCIATION HAZARD RATINGS:

**HEALTH HAZARD: 0 (materials that, under emergency conditions, would offer no hazard beyond that of ordinary combustible materials):** Gases and vapors whose LC<sub>50</sub> for acute inhalation toxicity is greater than 10,000 ppm. Dusts and mists whose LC<sub>50</sub> for acute inhalation toxicity is greater than 200 mg/L. Materials whose LD<sub>50</sub> for acute dermal toxicity is greater than 2000 mg/kg. Materials whose LD<sub>50</sub> for acute oral toxicity is greater than 2000 mg/kg. Materials that are essentially non-irritating to the respiratory tract, eyes and skin. 1 (materials that, under emergency conditions, can cause significant irritation): Gases and vapors whose LC<sub>50</sub> for acute inhalation toxicity is greater than 5,000 ppm but less than or equal to 10,000 ppm. Dusts and mists whose LC<sub>50</sub> for acute inhalation toxicity is greater than 10 mg/L but less than or equal to 200 mg/L. Materials whose LD<sub>50</sub> for acute dermal toxicity is greater than 1000 mg/kg but less than or equal to 2000 mg/kg. Materials whose LD<sub>50</sub> for acute oral toxicity is greater than 500 mg/kg but less than or equal to 2000 mg/kg. Materials that cause slight to moderate irritation to the respiratory tract, eyes and skin. 2 (materials that, under emergency conditions, can cause temporary incapacitation or residual injury): Gases and vapors whose LC<sub>50</sub> for acute inhalation toxicity is greater than 3,000 ppm but less than or equal to 5,000 ppm. Dusts and mists whose LC<sub>50</sub> for acute inhalation toxicity is greater than 2 mg/L but less than or equal to 10 mg/L. Materials whose LD<sub>50</sub> for acute dermal toxicity is greater than 200 mg/kg but less than or equal to 1000 mg/kg. 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. Any liquid whose saturated vapor concentration at 20°C (68°F) is equal to or greater than one-fifth its LC<sub>50</sub> for acute inhalation toxicity, if its LC<sub>50</sub> 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. 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. 3 (materials that, under emergency conditions, can cause serious or permanent injury): Gases and vapors whose LC<sub>50</sub> for acute inhalation toxicity is greater than 1,000 ppm but less than or equal to 3,000 ppm. Dusts and mists whose LC<sub>50</sub> for acute inhalation toxicity is greater than 0.5 mg/L but less than or equal to 2 mg/L. Materials whose LD<sub>50</sub> for acute dermal toxicity is greater than 40 mg/kg but less than or equal to 200 mg/kg. Materials whose LD<sub>50</sub> for acute oral toxicity is greater than 5 mg/kg but less than or equal to 50 mg/kg. Any liquid whose saturated vapor concentration at 20°C (68°F) is equal to or greater than one-fifth its LC<sub>50</sub> for acute inhalation toxicity, if its LC<sub>50</sub> is less than or equal to 3000 ppm and that does not meet the criteria for degree of hazard 4.

## DEFINITIONS OF TERMS (Continued)

### NATIONAL FIRE PROTECTION ASSOCIATION HAZARD RATINGS (continued):

**HEALTH HAZARD (continued): 3 (continued):** Compressed liquefied gases with boiling points between -30°C (-22°F) and -55°C (-66.5°F) that cause frostbite and irreversible tissue damage. Materials that are respiratory irritants. Cryogenic gases that cause frostbite and irreversible tissue damage. Materials that are corrosive to the respiratory tract. Materials that are corrosive to the eyes or cause irreversible corneal opacity. Materials that are corrosive to the skin. **4 (materials that, under emergency conditions, can be lethal):** Gases and vapors whose LC<sub>50</sub> for acute inhalation toxicity less than or equal to 1,000 ppm. Dusts and mists whose LC<sub>50</sub> for acute inhalation toxicity is less than or equal to 0.5 mg/L. Materials whose LD<sub>50</sub> for acute dermal toxicity is less than or equal to 40 mg/kg. Materials whose LD<sub>50</sub> for acute oral toxicity is less than or equal to 5 mg/kg. Any liquid whose saturated vapor concentration at 20°C (68°F) is equal to or greater than one-fifth its LC<sub>50</sub> for acute inhalation toxicity, if its LC<sub>50</sub> is less than or equal to 1000 ppm.

**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 accordance with Annex D. **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 accordance with Annex D. 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 Recommendation 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 water-miscible solution or dispersion with a water non-combustible liquid/solid content of more than 85 percent 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 a 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). Solids containing greater than 0.5 percent by weight of a flammable or combustible solvent are rated by the closed up flash point of the solvent. Most ordinary combustible materials. **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 in 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 percent 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) 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 a 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 percent 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 percent by weight of a flammable or combustible solvent are rated by the closed cup flash point of the solvent.

**INSTABILITY HAZARD: 0** Materials that in themselves are normally stable, even under fire conditions: Materials that have an estimated instantaneous power density (product of heat of reaction and reaction rate) at 250°C (482°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 estimated 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 estimated 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.

### NATIONAL FIRE PROTECTION ASSOCIATION HAZARD RATINGS (continued):

**INSTABILITY HAZARD (continued): 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 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. Materials that are sensitive to localized thermal or mechanical shock at normal temperatures and pressures.

### 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 vapors to form an ignitable mixture with air. **Autoignition Temperature**: The minimum temperature required to initiate combustion in air with no other source of ignition. **LEL** - the lowest percent of vapor in air, by volume, that will explode or ignite in the presence of an ignition source. **UEL** - the highest percent of vapor in air, by volume, that will explode or ignite in the presence of an ignition source.

### 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. Definitions of some terms used in this section are: **LD<sub>50</sub>** - Lethal Dose (solids & liquids) which kills 50% of the exposed animals; **LC<sub>50</sub>** - Lethal Concentration (gases) which kills 50% of the exposed animals; **ppm** concentration expressed in parts of material per million parts of air or water; **mg/m<sup>3</sup>** concentration expressed in weight of substance per volume of air; **mg/kg** quantity of material, by weight, administered to a test subject, based on their body weight in kg. Other measures of toxicity include **TDLo**, the lowest dose to cause a symptom and **TCLo** the lowest concentration to cause a symptom; **TDo**, **LDLo**, and **LDo**, or **TC**, **TCo**, **LCLo**, and **LCo**, the lowest dose (or concentration) to cause lethal or toxic effects.

**Cancer Information:** The sources are: **IARC** - the International Agency for Research on Cancer; **NTP** - the National Toxicology Program, **RTECS** - the Registry of Toxic Effects of Chemical Substances, **OSHA** and **CAL/OSHA**. IARC 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:** **BEI** - 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 TOXICITY INFORMATION:

A **mutagen** is a chemical that causes permanent changes to genetic material (DNA) such that the changes will propagate through generational lines. An **embryotoxin** 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 **teratogen** is a chemical that causes damage to a developing fetus, but the damage does not propagate across generational lines. A **reproductive toxin** is any substance that interferes in any way with the reproductive process.

### ECOLOGICAL INFORMATION:

EC is the effect concentration in water. **BCF** = Bioconcentration Factor, which is used to determine if a substance will concentrate in lifeforms which consume contaminated plant or animal matter. **TL<sub>m</sub>** = median threshold limit; Coefficient of Oil/Water Distribution is represented by **log K<sub>ow</sub>** or **log K<sub>oc</sub>** and is used to assess a substance's behavior in the environment.

### REGULATORY INFORMATION:

#### U.S. and CANADA:

**ACGIH:** American Conference of Governmental Industrial Hygienists, a professional association which establishes exposure limits.

This section explains the impact of various laws and regulations on the material. **EPA** is the U.S. Environmental Protection Agency. **NIOSH** is the National Institute of Occupational Safety and Health, which is the research arm of the U.S. Occupational Safety and Health Administration (**OSHA**).

**WHMIS** is the Canadian Workplace Hazardous Materials Information System. **DOT** and **TC** are the U.S. Department of Transportation and the Transport Canada, respectively. Superfund Amendments and Reauthorization Act (**SARA**); the Canadian Domestic/Non-Domestic Substances List (**DSL/NDSL**); the U.S. Toxic Substance Control Act (**TSCA**); Marine Pollutant status according to the **DOT**; the Comprehensive Environmental Response, Compensation, and Liability Act (**CERCLA** or **Superfund**); and various state regulations. This section also includes information on the precautionary warnings which appear on the material's package label. **OSHA** - U.S. Occupational Safety and Health Administration.

#### EUROPEAN and INTERNATIONAL:

The **DFG** is the Federal Republic of Germany's Occupation Health Agency, similar to the U.S. **OSHA**. **EU** is the European Union (formerly known as the **EEC**, European Economic Community). **EINECS**: This is the European Inventory of Now-Existing Chemical Substances. The **ARD** is the European Agreement Concerning the International Carriage of Dangerous Goods by Road and the **RID** are the International Regulations Concerning the Carriage of Dangerous Goods by Rail. **AICS** is the Australian Inventory of Chemical Substances. **METI** is the Japanese Ministry of Economy, Trade, and Industry.