SAFETY DATA SHEET



DATE ISSUED: 8/7/2018
SDS REF. No: 6H00 SERIES

6H00 SERIES 2K POLYURTHANE

1. PRODUCT AND COMPANY IDENTIFICATION

PRODUCT NAME: 6H00 SERIES HIGH SOLIDS 2K POLYURETHANE

PRODUCT CODE: 6H00 SERIES

PRODUCT USE: Industrial Solventborne Paint

MANUFACTURER
Cardinal Industrial Finishes

1329 Potrero Ave

S. El Monte, CA, 626 444-9274 **24 HR. EMERGENCY TELEPHONE NUMBER CHEMTREC (US Transportation)**: (800)424-9300 **CHEMTREC (International** : 1(202)483-7616

Transportation) WEB: WWW.CARDINALPAINT.COM

2. HAZARDS IDENTIFICATION

PICTOGRAMS



SIGNAL WORD: DANGER

HAZARD STATEMENTS:

H226 Flammable liquid and vapor.

H319 Causes serious eye irritation.

H336 May cause drowsiness or dizziness.

H360 May be fertility or the unborn child.

H372 Causes damage to organs through prolonged or repeated exposure.

H402 Harmful to aquatic life.

PRECAUTIONARY STATEMENTS:

P233 Keep container tightly closed.

P264 Wash thoroughly after handling.

P280 Wear protective gloves/protective clothing/eye protection/face protection.

P304 + P340 IF INHALED: Remove victim to fresh air and keep at rest in a position comfortable for breathing.

P312 Call a POISON CENTER or doctor/physician if you feel unwell.

P337 + P313 If eye irritation persists: Get medical advice/attention.

P403 Store in a well-ventilated place.

P501 Dispose of in accordance with Local, Regional, State, Federal and International Regulations.

R40 Limited evidence of a carcinogenic effect.

S36 Wear suitable protective clothing.

S37 Wear suitable gloves.

3. COMPOSITION/INFORMATION ON INGREDIENTS

Chemical Name	Weight %	CAS Number
Acetone	20% - 25%	67-64-1

n-Butyl Acetate	10% - 15%	123-86-4	

The follow substances may be present in varying quantities depending on color.

Titanium Dioxide	0% - 60%	13463-67-7	
Carbon Black	0% - 40%	1333-86-4	

4. FIRST AID MEASURES

Description of first and measures.

EYES CONTACT: Flush with large quantities of water for 15 to 30 minutes. Remove contact lenses. Keep eyes wide open while rising. If eye irritation persists: Get medical attention.

SKIN CONTACT: Wash exposed area with mild soap and water for 15 to 30 minutes. Remove contaminated clothing. Repeated exposure may cause dryness or cracking.

INGESTION: Rinse mouth. Do NOT induce vomiting. Keep victim warm and seek immediate attention.

INHALATION: Remove to fresh air and keep in a position comfortable to breath. Call a doctor/physician if you feel unwell. Get medical attention.

Most important symptoms and effects, both acute and delayed. Symptoms/injuries: Eye irritation

Symptoms/injuries after inhalation: May cause drowsiness or dizziness.

Symptoms/injuries after eve contact: Cause serious eve irritation.

Symptoms/injuries after ingestion: Ingestion may cause nausea, vomiting and diarrhea.

Indication of any immediate medical attention and special treatment needed.

If medical advice is needed, have product container or label on hand.

5. FIRE FIGHTING MEASURES

pressure modes.

SUITABLE EXTINGUISHING MEDIA: In the event of a fire, use specifically suitable extinguishing agents. Suitable extinguishing media: Foam, alcohol resistant foam, CO2, water fog. Unsuitable extinguishing media: Do not use heavy water stream. A heavy water stream my spread burning liquid.

FIRE FIGHTING PROCEDURE: Firefighting instructions: Use water spray or fog for cooling exposed containers. Exercise caution when fighting any chemical fire. Prevent fire-fighting water from entering the environment. Protection during firefighting: Firefighters should wear full protective gear. Do not enter fire area without proper protective equipment, including self-contained breathing apparatus with full face piece operated in pressure demand or other positive

UNUSUAL FIRE AND EXPLOSION HAZARD : Fire hazard: Highly flammable/liquid or vapor.

Explosive hazard: May form flammable/explosive vapor-air mixture.

6. ACCIDENTAL RELEASE MEASURES

PERSONAL PRECAUTIONS, PROTECTIVE EQUIPMENT AND EMERGENCY PROCEDURES:

General measures: Remove ignition sources. Use special care to avoid static electric charges. No smoking.

FOR NON-EMERGENCY PERSONNEL:

For non-Emergency procedures: Evacuate unnecessary personnel.

FOR EMERGENCY RESPONDERS:

Equip cleanup crew with proper protection. Avoid breathing fume, vapors.

ENVIROMENTAL PRECAUTIONS:

Prevent entry to sewers and public waters.

METHODS AND MATERIAL FOR CONTAINMENT AND CLEAN UP:

Collect damaged aerosols and use absorbent and/or inert material, then place in suitable container.

7. HANDLING AND STORAGE

PRECAUTIONS FOR SAFE HANDLING: Additional hazards when processed: Handle empty containers with care because residual vapors are flammable.

Precautions for safe handling: Wash hands and other exposed areas with mild soap and water before eating, drinking or smoking and when you are leaving work. Provide good ventilation in process area to prevent formation of vapor. No smoking. Use only non-sparking tools. Use outdoors or in a well ventilated area. Avoid breathing fume, vapors. Hygiene measures: Wash Skin thoroughly after handling.

CONDITIONS FOR SAFE STORAGE, INCLUDING INCOMPATIBILITIES: Storage conditions: Store in a dry, cool and well-ventilated place away from: Heat sources. Direct sunlight.

Incompatible products: Strong bases. Strong acids.

Incompatible materials: Source of ignition. Direct sunlight. Heat Sources.

8. EXPOSURE CONTROLS\PERSONAL PROTECTION

(5= 51.1)		
Acetone(67-64-1)		
USA ACGIH	ACGIH STEL TLV	750 ppm
USA ACGIH	ACGIH TWA TLV	500 ppm
USA NIOSH	NIOSH STEL (Table Z-1)	1,000 ppm, 2,400 mg/m3
USA NIOSH	NIOSH TWA	250 ppm, 590 mg/m3
USA OSHA	OSHA TWA (Table Z-1)	1,000 ppm, 2,400 mg,m3
Aluminum Hydroxide(21645-51-2)		
USA ACGIH	ACGIH (TLV) TWA	10 mg/m3 (Total dust), 3 mg/m3
		(Repairable fraction)
USA OSHA	OSHA (PEL) TWA	15 mg/m3 (Tptal dust), 5 mg/m3
		(Repairable fraction)
Carbon Black(1333-86-4)		
USA ACGIH	ACGIH TLV (mg/m3)	3.0 mg/m3
USA OSHA	OSHA PEL (mg/m3)	3.5 mg/m3
Crystalline Silica(14808-60-7)		
USA ACGIH	ACGIH (TLV) TWA	.025 mg/m3
Lithium Chloride(7447-41-8)		
USA OSHA	OSHA	Not Established.
n-Butyl Acetate(123-86-4)		·
USA ACGIH	ACGIH STEL	200 ppm
USA ACGIH	ACGIH TWA	150 ppm
USA OSHA	OSHA PEL (Table Z-1)	150 ppm, 710 mg/m3
n-Methyl-2-pyrrolidone(872-50-4)	· · · · · · · · · · · · · · · · · · ·	· · · · · · · · · · · · · · · · · · ·
USA ACGIH	ACGIH PEL	N/E
USA OSHA	OSHA TWA	N/E
Phenylethane(100-41-4)		
USA ACGIH	ACGIH STEL	125 ppm
USA ACGIH	ACGIH TWA	20 ppm
USA NIOSH	NIOSH REL	100 ppm, 435 mg/m3
USA NIOSH	NIOSH REL (ST)	125 ppm, 545 mg/m3
USA OSHA	OSHA STEL	125 ppm, 545 mg/m3
USA OSHA	OSHA TWA (Table Z-1)	100 ppm, 435 mg/m3
Styrene(100-42-5)		т = ос ррин, нос инд, инс
USA ACGIH	ACGIH STEL (ppm)	40 ppm
USA ACGIH	ACGIH TWA (ppm)	20 ppm
USA OSHA	OSHA TWA (ppm)	100 ppm
Titanium Dioxide(13463-67-7)	TEF /	. FF
PEL (Permissible Exposure Limit)	OSHA TWA	15 mg/m3
TLV	ACGIH TWA	10 mg/m3
Xylene(1330-20-7)		1 · · · · g ₁ · · · · ·
USA ACGIH	ACGIH STEL	150 ppm
USA ACGIH	ACGIH TWA	100 ppm
USA OSHA	OSHA TWA (Table Z-1)	100 PPM, 435 mg/m3
337. 33117.	JOS. IN THIN (TODIC Z I)	100 1111, 100 mg/mb

PERSONAL PROTECTIVE EQUIPMENT

RESPIRATORY PROTECTION: If TLV of the product or any component is exceeded, a NIOSH approved dust respirator is advised in absence of environmental control. OSHA Regulations also permit other NIOSH dust respirators under specified conditions. (See your Safety Equipment Supplier) Engineering or administrative controls should be implemented to reduce exposure.

HAND PROTECTION REMARKS : The suitability for a specific workplace should be discussed with the producers of the protective gloves.

EYES PROTECTION: Eye wash bottle with pure water.

Tightly fitting safety goggles.

Where face-shield and protective suit for abnormal processing problems.

SKIN AND BODY PROTECTION: Wear impervious clothing. Choose body protection according to the amount and concentration of the dangerous substance at the work place.

WORK HYGIENIC PRACTICES: When using do not eat or drink. When using do not smoke. Wash hands before breaks and at the end of workday.

9. PHYSICAL AND CHEMICAL PROPERTIES

Physical state	:	Liquid
Color	:	Various colors depending on the pigmentation.
Odor	:	Characteristic. Sweet. Mint like.
Odor threshold	:	No data available.
Ph	:	N/A - See Technical Data Sheet
Evaporation rate	:	Slower Than Ether
Melting point	:	-94.7 C (-138.46 F)
Freezing point	:	No data available.
Boiling point	:	133.0 deg F TO 281.0 deg F
Flash point	:	-4.00 deg F
Lower explosion limit	:	1.7
Upper explosion limit	:	12.8
Vapor pressure	:	185 mm Hg
Vapor density	:	Heavier than air
Relative density	:	No data available.
Density	:	10.5336
Solubility	:	No data available.
Partion coefficient: n-	:	No data available.
octanol/water		
Autoignition temperature	:	No data available.
Decomposition temperature	:	No data available.

10. STABILITY AND REACTIVITY

REACTIVITY: No dangerous reaction known under conditions of normal use.

CHEMICAL STABILITY: Stable under normal conditions.

CONDITIONS TO AVOID: Heat, flames and sparks. Extremely high temperatures and direct sunlight.

INCOMPATIBLE MATERIALS: Avoid contact with strong oxidizing agents.

HAZARDOUS DECOMPOSITION PRODUCTS: Carbon dioxide (CO2), carbon monoxide (CO), oxides of nitrogen (NOx), dense black smoke.

11. TOXICOLOGICAL INFORMATION

Acetone(67-64-1)	
Aspiration toxicity	Remarks: Symptoms of overexposure may be headache, dizziness, tiredness, nausea and vomiting., Concentrations substantially above TLV value may cause neurotic effects., Solvents may degrease the skin.
Carcinogenicity	Species: mouse, (female), Application Route: Dermal; Exposure time: .365 d (90%) or 424 d (100%), Dose: 0.1ml 90(71mg) or 100% (79mg), Frequency of Treatment: 3 times a wk, NOAEL: 79; Result: did not display carcinogenic properties., Carcinogenicity-Assessment: Not classified as a human carcinogen.
Germ cell mutagenicity	Test Type: mammalian cell gene mutation assay. Test species: Mouse Lymphoma, Metabolic activation: Without metabolic activation; Method: OECD Guideline 476; Result: negative; Test Type: Ames test, Metabolic activation: Without metabolic activation; Method: OECD Guideline 471; Result: negative, Test Type: Chromosome aberration test in vitro, Test species: Chinese hamster ovary (CHO), Metabolic activation: Without metabolic activation; Method: OECD Guideline 473; Result: negative; Genotoxicity in vivo: Test Type: I vivo micronucleus test.

	Test species: Mouse, Application Route: Oral, Exposure: 13 wk, Dose: 5,000, 10,000, 20,000
	ppm, Result: negative
Germ cell mutagenicity Assessment	Animal testing did not show any mutagenic effects.
LC50 (rat) Inhalation	76 mg/l (4 h exposure)
LD50 (rat) Oral	5,800 mg/kg; Symptoms: tremors
LD50 Dermal	>7,426 mg/kg
Repeated dose	Species: mouse, male, NOAEL: 20,000, Application Route: Oral, Exposure time: 13 wk,
exposure	Number of exposures: daily, Dose: 1250, 2500, 5000, 10000, 20000, Method OECD Test
	Guideline 408, GLP: No data available.; Species: mouse, female, NAOEL 20000, LAOEL:
	50000; Application Route: Oral, Exposure time: 13 wk, Number of exposures: daily, Dose: 1250, 2500, 5000, 10000, 20000, Method OECD Test Guideline 408, GLP: No data available;
	Repeated dose toxicity Assessment: causes mild skin irritation., Causes serious eye irritation.
Reproductive toxicity	Effects on fertility: Species: rat, male; Application Route: oral; Dose: 0, 5,000, 10,000 mg/l;
	Frequency of Treatment: 7 days/week; General Toxicity - Parent: LOAEL: 10,000; Fertility:
	10,000; Effects on fetal development: Species: rat; Application Route: Inhalation; Dose: 0,
	440, 2200, 11,000 ppm; Frequency of Treatment: 7 days/week; General Toxicity Material: NOAEC: 2,200 ppm; Teragenicity: NOAEC: 2,200 ppm; Embryo-fetal toxicity:: NOAEC: 2,200
	ppm; Result: No teratogenic potential. GLP: No data available.; Reproductive toxicity
	Assessment: Did not show teratogenic effects in animal experiments.
Respiratory or skin	Test type: Maximization test, Species: guinea pig, Assessment: Does not cause skin
sensitization	sensitization. Result: Did not cause sensitization on laboratory animals.
Serious eye damage/eye irritation	Species: rabbit, Result: Slightly irritating to eyes, Exposure time: 24 h, Classification: Irritating to eyes, Remarks: Eye irritation.
Skin	Species: rabbit, Exposure time: 24 h, Classification: Not irritating to skin, Method: In vivo,
corrosion/irritation	Result: Mild irritation, Remarks: Repeated or prolonged contact with the mixture may cause
	removal natural fat from the skin resulting in desiccation of the skin.
STOT - single exposure	Exposure routes: Inhalation (vapor); Assessment: May cause drowsiness or dizziness.
STOT- repeated	No data available.
exposure Aluminum Hydroxide(21	645-51-2)
Additional Information	RTECS: BD0940000 Nausea, Vomiting, and Constipation.
Aspiration hazard	No data available.
Carcinogenicity	IARC: No components of this product present at levels greater than or equal to 0.1% is
	identified as probable, possible or confirmed human carcinogen by IARC. ACGIH: No
	component of this product present at levels greater than or equal to 0.1% is identified as a carcinogen or potential carcinogen by ACGIH. NTP: No component of this product present at
	levels greater than or equal to 0.1% is identified as a known or anticipated carcinogen by
	NTP. OSHA: No component of this product present at levels greater than or equal to 0.1% is
	identified as a carcinogen or potential carcinogen by OSHA.
Dermal	No data available.
Germ cell mutagenicity	Mouse lymphocyte Result- negative Mutagenicity (micronucleus test) Rat - male Result:
Inhalation	negative No data available.
LD50 Oral - Rat -	>5,000 mg/kg, Oral - Rat - female
female - Acute toxicity	, 3, 3,
Reproductive toxicity	No data available.
Respiratory or skin	Maximization Test (GPMT) - Guinea pig Result- Does not cause skin sensitization.(OECD Test
sensitization Serious eye	Guideline 406) Eyes - Rabbit Result: No eye irritation (OECD Test Guideline 405)
damage/eye irritation	Lyes Rabbit Result. No eye irritation (OLOD Test Guideline 403)
Skin	Skin - Rabbit Result: No skin irritation - 4 h (OECD Test Guideline 404)
corrosion/irritation	
Specific target organ	No data available.
toxicity - repeated exposure	
Specific target organ	No data available.
toxicity - single	
exposure	
Amorphous Silica(7631-	
Additional toxicological information	The product is not subject to classification according ot internally approved calculation methods for preparations: When used and handled according to specifications, the product
miormation	does not have any harmful effects according to our experience and information provided to
	us.
Irritant of skin	Not irritating (rabbit) (OCED 404)
Irritating of eyes	Not irritating (rabbit) (OCED 405)
LC0 - Inhalative	>140->2000 mg/m3 / 4 h (Rat) (OCED 403)
LD50 - Dermal - Rabbit LD50 - Oral - Rat	>5000 mg/kg (Rabbit) >5000 mg/kg (Rat) (OECD 401)
LDJU - OIGI - Kal	>3000 mg/kg (nat) (OECD 401)

Other information - Oral	=> 1340 mg/kg/day
Sensitization	Not sensitization (guinea pig) (OCED 406)
Carbon Black(1333-86-4	
ACGIH	ACGIH The American Conference of Governmental Industrial Hygienists classifies carbon black as A4, Not Classifiable as a Human Carcinogen.
Carcinogenicity Classification	GHS- Not a hazardous substance or preparation according to the Global Harmonized System (GHS).
Human Epidemiology	Results of epidemiological studies of carbon black production workers suggest that cumulative exposure to carbon black may result in small decrements in lung function, as measured by FEV1. A recent U.S. respiratory morbidity study suggested a 27 mL decline in FEV1 from a 1 mg/m3 (inhalable fraction) exposure over a 40-year period. An older European investigation suggested an exposure to 1 mg/m3 (inhalable fraction) of carbon black over a 40-year working-lifetime will result in a 48 mL decline in FEV1. In contrast, normal age related decline over a similar period of time would be approximately 1200 ml. The relationship between symptoms and exposure to carbon black is less clear. In the U.S. study, 9% of the highest exposure group (in contrast to 5% of the unexposed group) reported symptoms consistent with chronic bronchitis. In the European study, methodological limitations in the administration of the questionnaire limit the drawing of definitive conclusions about symptoms.
Human Epidemiology - cont	Since this IARC evaluation of carbon black, Sorahan and Harrington 16) re-analyzed the UK study data using an alternative exposure hypothesis and found a positive association with carbon black exposure in two of the five plants. The same exposure hypothesis was applied by Morfeld and McCunney 17-18) to the German cohort; in contrast, they found no association between carbon black exposure and lung cancer risk and, thus, no support for the alternative exposure hypothesis used by Sorahan and Harrington 16).
Human Epidemiology - cont.	Morfeld and McCunney 19) applied a Bayesian approach to unravel the role of uncontrolled confounders and identified smoking and prior exposure to occupational carcinogens received before being hired in the carbon black industry as main causes of the observed lung cancer excess risk. Overall, as a result of these detailed investigations, no causative link between carbon black exposure and cancer risk in humans has been demonstrated. This view is consistent with the IARC evaluation in 2006. Several epidemiological and clinical studies of workers in the carbon black production industries show no evidence of clinically significant adverse health effects due to occupational exposure to carbon black. No dose response relationship was observed in workers exposed to carbon black.
Human Epidemiology - cont.	This study, however, indicated a link between carbon black and small opacities on chest films, with negligible effects on lung function. A study on carbon black production workers in the UK 10) found an increased risk of lung cancer in two of the five plants studied; however, the increase was not related to the dose of carbon black. Thus, the authors did not consider the increased risk in lung cancer to be due to carbon black exposure. A German study of carbon black workers at one plant 11-14) found a similar increase in lung cancer risk but, like the 2001 UK study 10), found no association with carbon black exposure. In contrast, a large US study 15) of 18 plants showed a reduction in lung cancer risk in carbon black production workers. Based upon these studies, the February 2006 Working Group at IARC concluded that the human evidence for carcinogenicity was inadequate 1).
IARC	IARC In 1995 IARC concluded, "There is inadequate evidence in humans for the carcinogenicity of carbon black." Based on rat inhalation studies IARC concluded that there is, "sufficient evidence in experimental animals for the carcinogenicity of carbon black," IARC's overall evaluation was that, "Carbon black is possibly carcinogenic to humans (Group 2B)". This conclusion was based on IARC's guidelines, which require such a classification if one species exhibits carcinogenicity in two or more studies. IARC performed another review in 2006, and again classified carbon black as possibly carcinogenic to humans (Group 2B). In its 1987 review IARC concluded, "There is sufficient evidence in experimental animals for the carcinogenicity of carbon black extracts." Carbon black extracts are classified as, possibly carcinogenic to humans (Group 2B). >8000 mg/kg
Mutagenic Effects and Germ Cell Mutagenicity	In an experimental investigation, mutational changes in the hprt gene were reported in alveolar epithelial cells in the rat following inhalation exposure to carbon black. This observation is believed to be rat specific and a consequence of "lung overload" which led to chronic inflammation and release of genotoxic oxygen species. This mechanism is considered to be a secondary genotoxic effect and thus, carbon black itself would not be considered to be mutagenic. Carbon black is not suitable to be tested in bacterial (Ames test) and other in vitro systems because of its insolubility in aqueous solutions. When tested, however, results for carbon black showed no mutagenic effects. Organic solvent extracts of carbon black can, however, contain traces of polycyclic aromatic hydrocarbons (PAHs). A study to examine the bioavailability of these PAHs showed that PAHs are very tightly bound to carbon black and not bioavailable.
NIOSH	NIOSH The U.S. National Institute of Occupational Safety and Health (NIOSH) 1978 criteria document on carbon black recommends that only carbon blacks with PAH contaminant levels greater than 0.1% require the measurement of PAHs in air. As some PAHs are possible human

	carcinogens, NIOSH recommends an exposure limit of 0.1 mg/m3 for PAHs in air, measured as the cyclohexane-extractable fraction.
NTP	NTP Carbon black is not designated a carcinogen by the U.S. National Toxicology Program
	(NTP), the U.S. Occupational Safety and Health Administration (OSHA) or the European Union (EU).
Reproductive and	No experimental studies on effects of carbon black on fertility and reproduction have been
Teratogenic Effects	located. However, based on toxic kinetic data, carbon black is deposited in the lungs and
	based on its specific physicochemical properties (insolubility, low absorption potential), it is
	not likely to distribute in the body to reach reproductive organs, embryo and/or fetusunder in
	vivo conditions. Therefore, no adverse effects of carbon black to fertility/reproduction or to
	fetal development are expected. No effects have been reported in long-term animal studies.
Sensitization	No animal data is available. No cases in humans have been reported.
STOT- repeated	Therefore, no STOT, Repeated exposure classification is made.
exposure	
STOT- single exposure	Inhalation studies with the rat showed lung effects (see Section 11.2 and 11.3), these effects
	are believed to be the effects of "lung overload" 1 and these effects are believed to be specific
	to the species. In addition, the European CLP Regulation states that no classification is
	necessary if the mechanism is not relevant to humans. 4) Also, the CLP Guidance on
	classification and labeling states that the "lung overload" mechanism is not relevant to
	humans. 4) Therefore, no STOT, Repeated Exposure classification is made
Crystalline Silica(14808	-60-/)
Acute toxicity - Dermal	No data available.
Acute toxicity -	No data available.
Inhalation	DTECC: VV/7220000 Purkeyed into 112
Additional Information	RTECS: VV7330000 Prolonged inhalation of crystalline silica may result in silicosis, a disabling
	pulmonary fibrosis characterized by fibrotic changes and military nodules in the lungs, a dry
	cough, shortness of breath, emphysema, decreased chest expansion, and increased
	susceptibility to tuberculosis. In advanced stages, loss of appetite, pleuritic pain, and total
	incapacity to work. Advanced silicosis may result in death due to cardiac failure or destruction of lung tissue. Crystalline silica is classified as group 1 "known to be carcinogenic to humans"
	by IARC and "sufficient evidence" of carcinogenicity by the NTP., The chronic health risks are
	associated with reparable particles of 3-4 um over protracted periods of time. Currently, there
	is a limited understanding of the mechanisms of quartz toxicity, including its mechanisms for
	lung carcinogenicity.
Additional Information	Additional studies are needed to determine whether the cell transforming activity of quartz is
(cont.)	related to its carcinogenic potential. Liver - Irregularities - Based on Human Evidence Liver -
(conci)	Irregularities - Based on Human Evidence.
Aspiration hazard	No data available.
Carcinogenicity	Limited evidence of carcinogenicity in human studies IARC: 1 - Group 1: Carcinogenic to
- ,	humans (Quartz) ACGIH: No component of this product present at levels greater than or
	equal to 0.1% is identified as a carcinogen or potential carcinogen by ACGIH. NTP: Known to
	be human carcinogen (Quartz) OSHA: No component of this product present at levels greater
	than or equal to 0.1% is identified as a carcinogen or potential carcinogen by OSHA.
Germ Cell	No data available.
mutagenicity	
Reproductive toxicity	No dada available.
Respiratory or skin sensitization	No data available.
	No data available.
Serious eye damage/eye irritation	וייט טמנמ מימוומטוכ.
Skin	No data available.
corrosion/irritation	The data diffuldation
Specific target organ	Inhalation - May cause damage to organs through prolonged or repeated exposure.
toxicity - repeated	The state of the s
exposure	
Specific target organ	No data available.
toxicity - single	
exposure	
Lithium Chloride(7447-4	41-8)
Additional Information	RTECS: OJ5950000 To the best of our knowledge, the chemical, physical, and toxicological
	properties have not been thoroughly investigated. Stomach - Irregularities - Based on Human
	Evidence Stomach - Irregularities - Based on Human Evidence.
Aspiration hazard	No data available.
Carcinogenicity	IARC: No component of this product present at levels greater than or equal to 0.1% is
	identified as probable, possible or confirmed human carcinogen by IARC. ACGIH: No
	I compare the first and the second of the se
	component of this product present at levels greater than or equal to 0.1% is identified as a
	carcinogen or potential carcinogen by ACGIH. NTP: No component of this product present at
	carcinogen or potential carcinogen by ACGIH. NTP: No component of this product present at levels greater than or equal to 0.1% is identified as a known or anticipated carcinogen by
	carcinogen or potential carcinogen by ACGIH. NTP: No component of this product present at

r .	
Dermal	No data available.
Germ cell mutagenicity	No data available.
Inhalation	No data available.
LD50 Oral - Rat -	526 mg/kg, Oral - Rat
Acute Toxicity	
Reproductive toxicity	No data available.
Respiratory or skin	No data available.
	NO data available.
sensitization	
Serious eye	No data available.
damage/eye irritation	
Skin	No data available.
O	NO data available.
corrosion/irritation	
Specific target organ	No data available.
toxicity - repeated	
exposure	
Specific target organ	No data available.
toxicity - single	
exposure	
n-Butyl Acetate(123-86-	4)
Aspiration hazard	No data available.
Carcinogenicity	No data available.
Inhalation	No data available.
LD-50 Dermal -	
	> 16ml/kg
(Rabbit)	
LD-50 Oral - (Rat)	14,130 mg/kg
Mutagenicity	In vitro: No data available. In vivo: No data available.
Other adverse effects:	No data available.
Repeated dose toxicity	No data available.
Reproductive toxicity	No data available.
Respiratory or skin	Skin Sensitization:, (Guinea Pig) - non-sensitizing.
	Skiii Selistization., (Guinea Fig) - non-selistizing.
sensitization	
Serious eye	(Rabbit, 24 h): none
damage/eye irritation	
Skin	(Rabbit, 24 h): none
	(Rabbit, 24 ii). Holie
corrosion/irritation	
Specific target organ	No data available.
toxicity - repeated	
exposure	
	N. C.
Specific target organ	Narcotic effect.
toxicity - single	
exposure	
n-Methyl-2-pyrrolidone(872-50-4)
Aspiration Hazard	Not Applicable.
Assessment other	Assessment of STOT single: Causes temporary irritation of the respiratory tract. Irritation /
acute effects	corrosion Assessment of irritating effects: Eye contact causes irritation. Skin contact causes
	irritation. Causes temporary irritation of the respiratory tract. EU-classification Skin Species:
	rabbit Result: Slightly irritating. Method: Draize test Literature data. The European Union (EU)
	has classified this substance with 'Irritating to skin' (R38). Eye Species: rabbit Result: Irritant.
	Method: Draize test Literature data. Sensitization Assessment of sensitization: Skin sensitizing
	effects were not observed in animal studies. Mouse Local Lymph Node Assay (LLNA) Species:
	mouse Result: Non-sensitizing. Method: OECD Guideline 429 The product has not been
	· · · · · · · · · · · · · · · · · · ·
	tested. The statement has been derived from substances/products of a similar structure or
	composition.
Carcinogenicity	Assessment of carcinogenicity: In long-term animal studies in which the substance was given
]	by inhalation, a carcinogenic effect was not observed. In long-term studies in rats in which
	the substance was given by feed, a carcinogenic effect was not observed. In long-term
	studies in rodents exposed to high doses, a tumorigenic effect was found; however, these
	results are thought to be due to a rodent-specific liver effect that is not relevant to humans.
	The whole of the information assessable provides no indication of a carcinogenic effect.
Genetic toxicity	Assessment of mutagenicity: The substance was not mutagenic in bacteria. No mutagenic
Correcte toxicity	
	effect was found in various tests with mammalian cell culture and mammals.
LC50 Inhalation - Rat	> 5.1 mg/l (OECD Guideline 403) Exposure time: 4 h An aerosol was tested. Limit
	concentration test only (LIMIT test). No mortality was observed.
LD50 Dermal - Rat	5,000 mg/m3; Species: rat (male/female) Value: > 5,000 mg/kg (OECD Guideline 402)
LD30 DCITIGI Rat	
	Literature data.
LD50 Oral - Rat	4,150 mg/kg (OECD Guideline 401) Literature data.
Repeated dose toxicity	Assessment of repeated dose toxicity: After repeated exposure the prominent effect is local
, , , , , , , , , , , , , , , , , , , ,	irritation. The substance may cause damage to the testes after repeated inhalation of high
	doses. Experiment

Reproductive toxicity	Assessment of reproduction toxicity: As shown in animal studies, the product may cause damage to the testes after repeated high exposures that cause other toxic effects.
Symptoms of Exposure	Medical conditions aggravated by overexposure Data available do not indicate that there are medical conditions that are generally recognized as being aggravated by exposure to this substance/product.
Teragenicity	Assessment of Teratogenicity: The substance caused malformations/developmental toxicity in laboratory animals.
Phenylethane(100-41-4)	
Aspiration toxicity	May be fatal if swallowed and enters airways.
Carcinogenicity	Species: mouse, (male and female) Application Route: Inhalation Exposure time: 103 wk
caremogeritatey	Activity duration: 6 h Dose: 0, 75, 250, 750 ppm Frequency of Treatment: 5 days/week NOAEL: 250 ppm Method: OECD Test Guideline 453 Result: evidence of carcinogenic activity Symptoms: increased incidences of alveolar/bronchiolar neoplasm's, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment: Carcinogenicity classification not possible from current data.
Germ cell mutagenicity	Genotoxicity in vitro, Test Type: Chromosome aberration test in vitro Test species: Chinese hamster ovary (CHO) Metabolic activation: with and without metabolic activation Method: OECD Test Guideline 473 Result: negative GLP: no : Test Type: Mammalian cell gene mutation assay Test species: mouse lymphoma cells Metabolic activation: with and without metabolic activation Method: OECD Test Guideline 476 Result: negative GLP: yes Genotoxicity in vivo: Test Type: In vivo micronucleus test species: mouse (male) Application Route: Oral Method: OECD Test Guideline 474 Result: negative GLP: yes Test Type: DNA damage and/or repair Test species: mouse (male and female)Application Route: Inhalation Method: OECD Test Guideline 486 Result: negative GLP: yes Germ cell mutagenicity Assessment: In vivo tests did not show mutagenic effects
LC50 (Mouse, Male)	10 mg/l Assessment: The component/mixture is moderately toxic after short term inhalation.
LD50 (rabbit)	
	15,433 mg/kg
Repeated dose toxicity Reproductive toxicity	Species: rat, male and female NOAEL: 75 mg/kg Application Route: Oral Exposure time: 28 d Dose: 75, 250 and 750 mg/kg bw/day Method: OECD Test Guideline 407 GLP: yes Symptoms: Increased kidney and liver weights Effects on fertility: Test Type: One generation study Species: rat, male and female
	Application Route: Inhalation Dose: 0, 100, 500 and 1000 ppm Duration of Single Treatment: 6 h General Toxicity - Parent: NOAEC: 1,000 ppm General Toxicity F1: NOAEC: 100 ppm Symptoms: Reduced fetal weight. Reduced offspring weight gain. Method: OECD Test Guideline 415 Result: No reproductive effects. GLP: yes Effects on fetal development: Species: rat Application Route: Inhalation Dose: 0, 100, 500, 1000, 2000 ppm Duration of Single Treatment: 15 d General Toxicity Maternal: NOAEC: 500 ppm Teratogenicity: NOAEC: 2,000 ppm Developmental Toxicity: NOAEC: 500 ppm Symptoms: Reduced body weight Method: OECD Test Guideline 414 Result: Developmental toxicity occurred at maternal toxicity dose levels GLP: No data available Reproductive toxicity - Assessment: No toxicity to reproduction Did not show teratogenic effects in animal experiments.
Respiratory or skin sensitization	Remarks: No data available
Serious eye damage/eye irritation	Species: rabbit Result: Mild eye irritation Remarks: No data available
Skin corrosion/irritation	Species: rabbit Result: Mild skin irritation
STOT - repeated exposure	Target Organs: Auditory system Assessment: May cause damage to organs through prolonged or repeated exposure., The substance or mixture is classified as specific target organ toxicant, repeated exposure, category 2.
STOT - single exposure	No data available.
Styrene(100-42-5)	
Irritation / corrosion - Eye	Species: Rabbit; Result: non-irritant; Method: BASF - Test
Irritation / corrosion - Sensitization	Species: Guinea pig; Result: non-sensitization; Method: OECD Guideline 406.
Irritation / corrosion - Skin	Species: Rabbit; Result: non-irritant; Method: BASF - Test
LC50 Dermal - Rat	Not determined
LC50 Inhalation - Rat	Exposure time 4 h ; not determined
LD50 Oral - Rat	>5,000 mg/kg
Titanium Dioxide(13463	
Carcinogenicity	In lifetime inhalation studies rats were exposed for 2 years to respectively 10, 50, 250 mg/m3 of respirable TiO2.
Dermal ALD (rabbit)	>10000 mg/m3
Eye irritation	slight irritation
Inhalation 4 h ALC	>6.82 mg/l
ORAL ALD (rat)	>2400 mg/kg
Sensitation	Did not cause Sensitation on laboratory animals.
Skin irritation	slight irritation

Xylene(1330-20-7)	
Acute dermal toxicity	Acute toxicity estimate: 1,100 mg/kg Method: Expert judgment.
Acute inhalation	Acute toxicity estimate, 4631 ppm Exposure time, 4 h Test atmosphere: gas Method;
toxicity	Calculation method.
Acute toxicity Product	Acute oral toxicity: Acute toxicity estimate: 3,523 mg/kg Method: Calculation method.
Aspiration Toxicity	May be fatal if swallowed and enters airways.
Carcinogenicity	Species: mouse, (male and female) Application Route: Oral Exposure time: 103 wk Dose: 0, 500 or 1000 mg/kg Frequency of Treatment: 5 days/week Method: Directive 67/548/EEC, Annex V, B.32. Result: did not display carcinogenic properties GLP: No data available, Carcinogenicity - Assessment: Animal testing did not show any carcinogenic effects.
Germ cell mutagenicity	12:00:00 AM
Germ cell mutagenicity Assessment	Animal testing did not show any mutagenic effects.
LC50 (rat, male) Inhalation	6700 ppm Exposure time: 4 h Method: Directive 67/548/EEC, Annex V, B.2. GLP: No data available Assessment: The substance or mixture is classified as specific target organ toxicant, single exposure, category 3 with respiratory tract irritation. Remarks: Acutely Toxic Category 4
LC50 (rat, male) Oral	3,523 mg/kg Method: EU Method B.1 (Acute Toxicity, Oral) Target Organs: Kidney, Bladder GLP: no
Repeated dose toxicity	Species: rat, male and female NOAEL: 250 mg/kg Application Route: Oral Exposure time: 103 wk Number of exposures: 5 d/wk Dose: 0, 250 or 500 mg/kg Assessment: The substance or mixture is classified as specific target organ toxicant, repeated exposure, category 2.
Reproductive toxicity	Effects on fertility: Test Type: Two-generation study Species: rat, male and female Application Route: Inhalation Dose: 0, 25, 100 and 500 ppm Duration of Single Treatment: 6 h Frequency of Treatment: 7 days/week General Toxicity - Parent: NOAEC: > 500 ppm General Toxicity F1: NOAEC: > 500 ppm Early Embryonic Development: NOAEC: > 500 ppm Result: No reproductive effects. Effects on fetal development: Species: rat Application Route: Inhalation Dose: 0, 100, 500, 1000 or 2000 ppm Duration of Single Treatment: 14 d Frequency of Treatment: 6 hr/day General Toxicity Maternal: NOAEC: 500 ppm Teratogenicity: NOAEC: > 2,000 Developmental Toxicity: NOAEC: 100 ppm Result: No teratogenic effects., Developmental toxicity occurred at maternal toxicity dose levels Reproductive toxicity - Assessment: Animal testing did not show any effects on fertility. Damage to fetus not classifiable
Respiratory or skin sensitization	Remarks: No data available
Serious eye damage/eye irritation	Species: rabbit Result: Mild eye irritation
Skin corrosion/irritation	Species: rabbit Exposure time: 24 h Result: Irritating to skin Remarks: Skin irritation, Category 2
STOT - repeated exposure	Target Organs: Liver, Kidney, Central nervous system Assessment: May cause damage to organs through prolonged or repeated exposure.
STOT - single exposure	No data available.

12. ECOLOGICAL INFORMATION

Acetone(67-64-1)	Acetone(67-64-1)		
Bioacculative potential	Partition coefficient: n-octanol/water: log Pow: -0.24		
EC50 (Daphnia magna	7,630 mg/l (Exposure time 48 h); Test substance: Acetone		
(Water flea))			
LC50 (Oncorhynchus	6,100 mg/l (Exposure time: 48 h)		
mykiss (rainbow			
trout))			
Mobility in soil	No data available.		
Other adverse effects	No data Available. Regulation: 40 CFR Protection of Environment; Part 82 Protection of		
	Stratospheric Ozone - CAA Section 602 Class I Substances., Additional ecological information:		
	No data available.		
Persistence and	Biodegradability: Remarks: No data available		
degradability			
Toxicity to algee	Remarks: No data available		
Aluminum Hydroxide(21			
Bioaccumulative	Inert material.		
potential			
EC50 - Daphnia -	>10,000 mg/l, Daphnia magna (Water flea) (OECD Test Guideline 202)		
Toxicity to daphnia and			
other aquatic			
invertebrates			
EC50 - Fish - Toxicity	>10,000 mg/l, Fish		
to fish			
Mobility in soil	Inert material.		

NOEC - Toxicity to algae	>0.004 mg/l, 72 h, Pseudokirchneriella subcapitata (algae) - (OECD Test Guideline 201)
Other adverse effects	None known.
Persistence and	Non-degradable
degradability	
Amorphous Silica(7631-	
Additional ecological	General notes: Do not allow product to reach ground water, water course or sewage system.
information	
Bioaccumulative potential	No further relevant information available.
EC50 - Algae	>10000 mg/l (Scenedesmus subspicatus) (72 h) (OCED 201) comparable substance
EC50 - Daphnia magna	>1000 mg/l (Daphnia magna) (24 h) (OCED 202)
LCO - Zebra fish	10000 mg/l (zebra fish) (96 h) (static) (OCED203)
Mobility in soil	No further relevant information available.
Persistence and	The product is chemically and biologically inert. By the insolubility in water there is a
durability	separation at every filtration and sedimentation process.
Carbon Black(1333-86-4	
Behavior in water treatment plants	Activated sludge, EC0 (3 h) > 800 mg/L. DEV L3 (TTC test)
Bioaccumulation Potential	Potential bioaccumulation is not expected because of the physicochemical properties of the substance
EC50 (Scenedesmus subspicatus)	> 10,000 mg/L, OECD (Guideline 201)
EC50 Daphnia magna (waterflea)	>5600 mg/l (24 h) OECD (Guideline 202)
Environmental fate	Carbon black is an inert solid, stable and insoluble in water or organic solvents. Its vapor
Ziivii oiiii eitai iate	pressure is negligible. Based on these properties it is expected that carbon black will not
	occur in air or water in relevant amounts. Also potential for distribution via water or air can be
	dismissed. The deposition in soil or sediments is therefore the most relevant compartment of
	fate in the environment.
LC50 Brachydanio reio	>1000 mg/l (96 h) OECD (Guideline 203)
(zebrafish)	
NOEC 50	> 10,000 mg/L, OECD (Guideline 201)
(Scenedesmus	
subspicatus)	
Crystalline Silica(14808-	
Bioaccumulative	No data available.
potential	No deba suscitable
Mobility in soil	No data available.
Other adverse effects Persistence and	No data available. No data available.
degradability	NO data available.
Results of PBT and	PBT/vPvB assessment not available as chemical safety assessment not required/not
vPvB assessment	conducted
Toxicity	No data available.
Lithium Chloride(7447-4	
Bioaccumulative	No data available.
potential	
EC50 - Daphnia magna	1.2 mg/l - 64 h, Daphnia magna (Water flea)
(Water flea) - to	
daphnia and other	
aquatic invertebrates	
LC50 - Ptychocheilus	17 mg/l - 96 h, -Ptychocheilus lucius
lucius - Toxicity to fish	
Mobility in soil	No data available.
Other adverse effects	An environmental hazard cannot be excluded in the event of unprofessional handling or
Davidat	disposal. Harmful to aquatic life.
Persistence and degradability	No data available.
Results of PBT and	PBT/vPvB assessment not available as chemical safety assessment not required/not
vPvB assessment	conducted.
n-Butyl Acetate(123-86-	
Bioaccumulative potential	No data available.
Chronic Toxicity	Fish: No data available. Aquatic invertebrates: No data available. Toxicity to Aquatic Plants: No data available.
LC-50 (Fathead	18 mg/l, (96 h)
Minnow) Acute Toxicity	

LC-50 (Water Flea)	44 mg/l , (48 h)
Aquatic invertebrates	
Mobility in soil	Known or predicted distribution to environmental compartments: No data available.
Other adverse effects	No data available.
Persistence and	83 % (28 d), Biological Oxygen Demand: BOD-5: 730 mg/g, Chemical Oxygen Demand:1,010
degradability	mg/g, BOD/COD ratio:72 %.
Results of PBT and	No data available.
vPvB assessment	
n-Methyl-2-pyrrolidone(8	
Additional information	Sum parameter Chemical oxygen demand (COD): (DIN 38409 Part 41) approx. 1,600 mg/g
	Biochemical oxygen demand (BOD) Incubation period 5 d: < 2 mg/g Absorbable organically-
	bound halogen (AOX): This product contains no organically-bound halogen.
Bioaccumulative	Assessment bioaccumulation potential Because of the n-octanol/water distribution coefficient
potential	(log Pow) accumulation in organisms is not to be expected.
EC50 (Algae)	> 500 mg/l, (72 h), Scenedesmus subspicatus (DIN 38412 Part 9) The details of the toxic
	effect relate to the nominal concentration.
EC50 (Daphnia)	> 1,000 mg/l, (24 h), Daphnia magna (DIN 38412 Part 11, static) The details of the toxic
1550 (6.1)	effect relate to the nominal concentration.
LD50 (fish)	> 500 mg/l, Salmo gairdneri, syn. O. mykiss (static) The details of the toxic effect relate to
17.55	the nominal concentration.
Microorganisms/Effect	Toxicity to microorganisms DIN EN ISO 8192 aquatic activated sludge, industrial/EC50 (0.5
on activated sludge	h): > 600 mg/l The details of the toxic effect relate to the nominal concentration.
Mobility in soil	Assessment transport between environmental compartments The substance will rapidly
	evaporate into the atmosphere from the water surface. Adsorption to solid soil phase is not
Dami's London	expected.
Persistence and	Assessment biodegradation and elimination (H2O) Readily biodegradable (according to OECD
degradability	criteria). Elimination information 73 % BOD of the ThOD (28 d) (OECD 301C; ISO 9408;
	92/69/EEC, C.4-F) (aerobic, Inoculum conforming to MITI requirements (OECD 301C))
	Readily biodegradable (according to OECD criteria). Assessment of stability in water In
Db (100 41 4)	contact with water the substance will hydrolyze slowly.
Phenylethane(100-41-4)	Daytition on efficient, restand Western Land Rose 2 02
Bioaccumulative	Partition coefficient: noctanol/water : log Pow: 2.92
potential	1. O may 1. Expressive binary 40 h. Tark Turney shakin had
EC50 (Daphnia magna	1.8 mg/l Exposure time: 48 h Test Type: static test
(Water flea)) EC50	5.4 mg/l Exposure time: 72 h Test Type: static test Analytical monitoring: yes Method: Static
(Pseudokirchneriella subcapitata)	GLP: yes
LC50 (Oncorhynchus	4.2 mg/l Exposure time: 96 h Test Type: semi-static test
mykiss (rainbow	4.2 mg/r Exposure time. 30 m rest Type. Serm-static test
trout))	
Mobility in soil	No data available.
Other adverse effects	Results of PBT and vPvB assessment : This substance is not considered to be persistent,
Other daverse effects	bioaccumulation nor toxic (PBT). This substance is not considered to be persistent,
	very bioaccumulation (vPvB).
Persistence and	Biodegradability: Inoculum: activated sludge Concentration: 22 mg/l Result: Readily
degradability	biodegradability: Inoculum: activated studge concentration: 22 mg/r Result: Readily biodegradable. Biodegradation: 70 % Exposure time: 28 d GLP: yes
Toxicity to daphnia and	(Daphnia): 3.6 mg/l Toxicity to bacteria : GLP: Remarks: No data available Ecotoxicology
other aquatic	Assessment Chronic aquatic toxicity: Harmful to aquatic life with long lasting effects.
invertebrates (Chronic	A second contents adjusted toxicity. The final to adjusted inc. With long labeling circuits.
toxicity)	
Styrene(100-42-5)	
Bioaccumulation	At present state of knowledge, no negative ecological effects are expected.
Chronic	No data available regarding toxicity to daphnids.
Chronic	No data available regarding toxicity to dish.
EC50 (Algae)	(72 h); No data available concerning toxicity for algae.
EC50 (Daphnia) Acute	(48 h) No data available regarding toxicity to daphnia.
LC50 Fish (Leuciscus	>100 mg/l (96 h)
idus) Acute	···· · · · · · · · · · · · · · · ·
Microorganisms	Toxicity to microorganisms: The inhibition of the degradation activity sludge is not antiquated
co. gamomo	when introduced to biological treatment plants in appropriate low concentrations.
Titanium Dioxide(13463-	
LC50 fish	Fathead minnow 96 h >1000 mg/l
Xylene(1330-20-7)	- ======
Bioaccumulative	Partition coefficient: noctanol/water : log Pow: 2.77 - 3.15
potential	Taradan coemelenc noctanoly nater i log row. 2.77 3.13
EC50	4.36 mg/l End point: Growth rate Exposure time: 73 h Test Type: static test Analytical
(Pseudokirchneriella	monitoring: yes
subcapitata)	

IC50 (Daphnia magna	1 mg/l Exposure time: 24 h Test Type: static test substance: Information given is based on	
(Water flea))	data obtained from similar substances. Method: OECD Test Guideline 202 GLP	
LC50 (Oncorhynchus	2.6 mg/l Exposure time: 96 h Test substance: Information given is based on data obtained	
mykiss (rainbow	from similar substances. Method: OECD Test Guideline 203 GLP: No data available	
trout))		
Mobility in soil	No data available.	
Persistence and	Biodegradability: Inoculum: activated sludge Result: Readily biodegradable. Biodegradation:	
degradability	72 % Exposure time: 20 d	

13. DISPOSAL CONSIDERATIONS

WASTE TREATMENT METHODS

GENERAL INFORMATION: No data available.

DISPOSAL METHOD: Dispose of waste and residues in accordance with Local, State, and Federal Regulations. Mix with compatible chemical which is less flammable and incinerate. Since emptied containers retain product residue, follow label warnings even after container is emptied. Residual vapors may explode on ignition; do not cut, drill, grind or weld or near this container.

14. TRANSPORT INFORMATION

*CHECK WITH YOUR CARRIER FOR ADDITIONAL RESTRCITIONS THAT MAY APPLY.

USDOT GROUND

DOT (DEPARTMENT OF TRANSPORTATION)

PROPER SHIPPING NAME (DOT): Paint, flammable liquid

HAZARDS CLASS: 3 UN/NA NUMBER: UN1263 PACKING GROUP: PG II

EMERGENCY RESPONSE GUIDE (ERG): 128

IATA (AIR)

DOT (INTERNATIONAL AIR TRANSPORTATION ASSOCIATION)

PROPER SHIPPING NAME: Paint, flammable liquid

HAZARDS CLASS: 3 UN/NA NUMBER: UN1263 PACKING GROUP: PG II

EMERGENCY RESPONSE GUIDE (ERG): 128

IMDG (OCEAN)

PROPER SHIPPING NAME: Paint, flammable liquid

HAZARDS CLASS: 3 UN/NA NUMBER: UN1263 PACKING GROUP: PG II

EMERGENCY RESPONSE GUIDE (ERG): 128

MARINE POLLUTANT: No

SPECIAL PRECAUTIONS: P210 Keep away from heat/sparks/open flames/hot surfaces. No smoking. P235 Keep cool.

15. REGULATORY INFORMATION

US FEDERAL REGULATIONS

All ingredients in Section #3 are TSCA (Toxic Substance Control Act) listed.

OSHA HAZARDS: Flammable liquid, Moderate skin irritant, Moderate eye irritant, Carcinogen.

EPCRA - Emergency

CERCLA REPORTABLE QUANTITY

This product contains:	Chemical CAS#
Carbon Black	1333-86-4
n-Butyl Acetate	123-86-4
Xylene	1330-20-7
Phenylethane	100-41-4

SARA 304 Extremely Hazardous Substances Reportable Quantity: This material does not contain any components

with a section 304 EHS RQ.

SARA TITLE III (SUPERFUND AMENDMENRS AND REAUTHORIZATION ACT)

SARA 311/312 Hazards: Fire Hazard, Acute Health Hazard, Chronic Health Hazard

SARA 313:

This product contains:	Chemical CAS#
Acetone	67-64-1
n-Butyl Acetate	123-86-4
Titanium Dioxide	13463-67-7
Carbon Black	1333-86-4

CLEAN AIR ACT:

This product contains:	Chemical CAS#
Styrene	100-42-5
Ethylene glycol mono phenyl ether	122-99-6
Phenylethane	100-41-4

INTERNATIONAL REGULATIONS

CLASSIFICATION ACCORDING TO REGULATION (EC) No. 1272/2008 (CLP):

Flam. Liq. Cat. 2; H226 Eye Irrit. Cat. 2; H319 STOT SE Cat. 3; H336 Reprod. Tox. Cat 1B; H360 STOT RE Cat. 1; H372 Aquatic Tox. Cat. 3; H402

NATIONAL REGULATIONS

This product contains:	Chemical CAS#
~Titanium Dioxide	13463-67-7
~Carbon Black	1333-86-4

IARC KEY

~ Indicates a chemical listed by IARC as a possible carcinogen.

STATE REGULATIONS **CALIFORNIA PROPOSITION 65**

This product contains:	Chemical CAS#
+n-Methylpyrrolidone	872-50-4
*Phenylethane	100-41-4
*Styrene	100-42-5

PROPOSTION 65 KEY

* MARNING Cancer – www P65Warnings.ca.gov



MARNING Reproductive Harm – www P65Warnings.ca.gov

+ MARNING Cancer and Reproductive Harm – www P65Warnings.ca.gov

Massachusetts Right to Know

This product contains Chemical CAS#

[^] Indicates a chemical listed by IARC as a carcinogen.

Acetone	67-64-1
Carbon Black	1333-86-4
n-Butyl Acetate	123-86-4
Xylene	1330-20-7
Phenylethane	100-41-4

Pennsylvania Right to Know

This product contains	Chemical CAS#
Acetone	67-64-1
Titanium Dioxide	13463-67-7
Carbon Black	1333-86-4
Amorphous Silica	7631-86-9
n-Butyl Acetate	123-86-4
Aluminum Hydroxide	21645-51-2
Lithium Chloride	7447-41-8
Xylene	1330-20-7
Phenylethane	100-41-4

New Jersey Right to Know

This product contains	Chemical CAS#
Acetone	67-64-1
Titanium Dioxide	13463-67-7
Carbon Black	1333-86-4
Amorphous Silica	7631-86-9
n-Butyl Acetate	123-86-4
Aluminum Hydroxide	21645-51-2
Lithium Chloride	7447-41-8
Xylene	1330-20-7
Phenylethane	100-41-4

16. OTHER INFORMATION

Other Product Information

% Volatile by Volume: 60.19 % Volatile by Weight: 39.37 % Solids by volume: 39.81 % Solids by Weight: 60.63 % Exempt by Volume: 38.86 % Exempt by Weight: 24.34

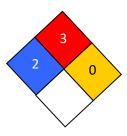
VOC CONTENT: Excluding Exempt VOC: 310 Including Exempt VOC: 190

HMIS RATING

Health :	2*
Flammability :	3
Reactivity :	0
Personal Protection:	Н

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NFPA CODES



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