SAFETY DATA SHEET



 DATE ISSUED :
 3/30/2016

 SDS REF. No :
 6200 SERIES

6200 SERIES POLYURETHANE

1. PRODUCT AND COMPANY IDENTIFICATION

PRODUCT NAME: 6200 SERIESPOLYURETHANE

PRODUCT CODE: 6200 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.

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	15% - 20%	67-64-1
Parachlorobenzotrifluoride	5% - 10%	98-56-6
P.M. Acetate	1% - 5%	108-65-6

n-Butyl Acetate	1% - 5%	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 aid 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 eye contact: Cause serious eye irritation.

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

Indication of any immediate medical attention and special treatment needed.

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

5. FIRE FIGHTING MEASURES

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 pressure modes.

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.

ENVIRONMENTAL 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

Acetone(67-64-1)		
USA ACGIH	ACGIH STEL TLV	750 ppm
	ACGIH TWA TLV	
USA ACGIH		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 (Respirable fraction)
USA OSHA	OSHA (PEL) TWA	15 mg/m3 (Total dust), 5 mg/m3 (Respirable 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
Cyclohexanone(108-94-1)	COLIN LE (Mg/ MO)	515 mg/m5
USA ACGIH	ACGIH (TLV) STEL	50 ppm
USA ACGIH	ACGIH (TLV) TWA	20 ppm
USA NIOSH	NIOSH (TLV) TWA	25 ppm, 100 mg/m3
USA OSHA	OSHA (OEL) Table Z-1 TWA	50 ppm, 200 mg/m3
	OSHA (OEL) Table Z-1 TWA	30 ppili, 200 flig/filis
Dibutyltin Dilaurate(77-58-7)	ACCIU CTEL	0.2/2
USA ACGIH	ACGIH STEL	0.2 mg/m3
USA ACGIH	ACGIH TWA	0.1 mg/m3
USA NIOSH	NIOSH REL	0.1 mg/m3
USA OSHA	OSHA PEL (Table Z-1)	0.1 mg/m3
USA OSHA	OSHA TWA (Table Z-1A)	0.1 mg/m3
Formaldehyde(50-00-0)		
USA ACGIH	ACGIH (TLV)	0.3 ppm
USA OSHA	OSHA (PEL) STEL	2 ppm
USA OSHA	OSHA (PEL) STEL	2 ppm STEL 15 min
USA OSHA	OSHA (PEL) TWA	0.75 ppm
Isobutyl Alcohol(78-83-1)		
USA ACGIH	ACGIH TWA	50 ppm
USA OSHA	OSHA PEL	100 ppm, 300 mg/m3
Methyl Amyl Ketone(110-43-0)		
USA ACGIH	ACGIH TLV TWA	50 ppm
USA OSHA	OSHA PEL (Table Z-1)	100 ppm, 465 mg/m3
Methyl Ethyl Ketone(78-93-3)	TOSTIN TEL (TUBIC E 1)	100 ppm, 103 mg/ms
USA ACGIH	ACGIH STEL (ppm)	300 ppm
USA ACGIH	ACGIH TWA (ppm)	200 ppm
USA OSHA	OSHA PEL (STEL) (ppm)	100 ppm
USA OSHA	OSHA PEL (STEL) (ppili) OSHA PEL TWA (mg/m3)	410 mg/m3
	OSHA FLL TWA (HIg/HIS)	1 410 Hig/III3
n-Butyl Acetate(123-86-4)	ACCIH CTEI	200 nnm
USA ACGIH	ACGIH STEL	200 ppm
USA ACGIH	ACGIH TWA	150 ppm
USA OSHA	OSHA PEL (Table Z-1)	150 ppm, 710 mg/m3
P.M. Acetate(108-65-6)	TATALL (MEEL) TIME	150
USA AIHA	AIAH (WEEL) TWA	50 ppm
Parachlorobenzotrifluoride(98-56-6)		
USA ACGIH	USA ACGIH	Conatins no substances with exposure limit values.
Styrene(100-42-5)		•
USA ACGIH	ACGIH STEL (ppm)	40 ppm
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USA ACGIH	ACGIH TWA (ppm)	20 ppm
USA OSHA	OSHA TWA (ppm)	100 ppm
TALC(14807-96-6)		
USA ACGIH	ACGIH (TLV) TWA	2 mg/m3
USA NIOSH	NIOSH (REL) TWA	2 mg/m3
USA OSHA	OSHA (Table Z-3) Mineral Dusts TWA	20 Millon particles per cubic foof
Titanium Dioxide(13463-67-7)		
PEL (Permissible Exposure Limit)	OSHA TWA	15 mg/m3
TLV	ACGIH TWA	10 mg/m3

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

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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 294.0 deg F
Flash point	:	-4.00 deg F
Lower explosion limit	:	.9
Upper explosion limit	:	12.8
Vapor pressure	:	185 mm Hg
Vapor density	:	Heavier than air
Relative density	:	No data available.
Density	:	12.8715
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 narcotic 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	Animal testing did not show any mutagenic effects.
Assessment	
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 exposure	Species: mouse, male, NOAEL: 20,000, 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.; 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; Tetragenicity: 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
sensitsation	sensitsation. Result: Did not cause sensitsation on laboratory animals.
Serious eye damage/eye irritation	Species: rabbit, Result: Slightly irritating to eyes, Exposure time: 24 h, Classification:
Skin corrosion/irritation	Irritating to eyes, Remarks: Eye irritation. Species: rabbit, Exposure time: 24 h, Classification: Not irritating to skin, Method: In
Skiii coi i osioriyii i itatiori	vivo, 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 exposure	No data available.
Aluminum Hydroxide(21645-	
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:
Tabalation	negative
Inhalation	No data available.
LD50 Oral - Rat - female - Acute toxicity	>5,000 mg/kg, Oral - Rat - female
Reproductive toxicity	No data available.
Respiratory or skin	Maximization Test (GPMT) - Guinea pig Result- Does not cause skin sensitization.(OECD
sensitization	Test Guideline 406)
Serious eye damage/eye	Eyes - Rabbit Result: No eye irritation (OECD Test Guideline 405)
irritation	·
Skin corrosion/irritation	Skin - Rabbit Result: No skin irritation - 4 h (OECD Test Guideline 404)

Specific target organ toxicity - repeated exposure	No data available.
Specific target organ toxicity - single exposure	No data available.
Amorphous Silica(7631-86-9)	
Additional toxicological	The product is not subject to classification according ot internally approved calculation
information	methods for preparations: When used and handled according to specifications, the
mormacion	product does not have any harmful effects according to our experience and information
	provided to us.
Irritant of skin	Not irritating (rabbit) (OCED 404)
Irritation of eyes	Not irritating (rabbit) (OCED 405)
LC0 - Inhalative	>140->2000 mg/m3 / 4 h (Rat) (OCED 403)
LD50 - Dermal - Rabbit	>5000 mg/kg (Rabbit)
LD50 - Oral - Rat	>5000 mg/kg (Rabbit) >5000 mg/kg (Rat) (OECD 401)
Other information - Oral	=> 1340 mg/kg/day
Sensitization	Not sensitizating (guinea pig) (OCED 406)
Carbon Black(1333-86-4)	Not sensitizating (guinea pig) (OCLD 400)
ACGIH	ACCIU The American Conference of Covernmental Industrial Hydionists classifies carbon
	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
Home - Faid ide	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) .l
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).
LD50 (Rat)	>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 Teratogenic Effects	No experimental studies on effects of carbon black on fertility and reproduction have been located. However, based on toxicokinetic 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 foetus under in vivo conditions. Therefore, no adverse effects of carbon black to fertility/reproduction or to foetal 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 exposure	Therefore, no STOT, Repeated exposure classification is made.
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
Cyclohexanone(108-94-1)	
Aspiration hazard	Solvent may degrease the skin.
Carcinogenicity	This product is or contains a component that is not classifiable as to its carcinogenicity based on its IARC, ACGIH, NTP, or EPA classification. IARC: 3 - Group 3: Not classifiable as to its carcinogenicity to humans (Cyclohexanone) 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.
Germ cell mutagenicity	Not mutagenic in Ames Test Ames test S. typhimurium Result: negative Human fibroblast Result: Laboratory experiments have shown mutagenic effects.
LC50 Inhalation - Rat	> 6.2 mg/l Rat - (4 h)
LD50 Dermal - Rabbit LD50 Oral - Rat - Acute toxicity	794 - 3,160 mg/kg 1534 mg/kg (Rat), Method: Standard Acute.
Reproductive toxicity	Overexposure may cause reproductive disorder(s) based on tests with laboratory animals.
Respiratory or skin sensitization	Test type: Maximization Test (GPMT), Species: guinea pig, Assessment: Does not cause skin sensitsation. Method: In vivo, Result: Does not cause skin sensitsation.
Serious eye damage/eye irritation	Eyes - Rabbit Result: Risk of serious damage to eyes, 24 h
Skin corrosion/irritation	Skin - Rabbit Result: Irritating to skin. (OECD Test Guideline 404)
Specific target organ toxicity - repeated exposure	Harmful if swallowed., Harmful in contact with skin., Harmful if inhaled., Causes skin irritation., Cuases serious eye damage.
Specific target organ toxicity - single exposure	No data available Acute inhalation toxicity - Breathing difficulties
Dibutyltin Dilaurate(77-58-7)	
Chronic Health Hazard	Dibutyltin compounds have shown reproductive and immunotoxic effects in laboratory animals. Abnormalities noted at necropsy of animals treated with 2000 mg/kg of dibutyltin dilaurate were hemorrhagic lungs, dark liver, dark kidneys, hemorrhage of gastric mucosa, hemorrhage of the large and small intestines, enlarged bile duct and behavioral and central nervous system effects. Decreased fertility was seen in hens
	following dietary administration equal to 78 mg/kg.
Eve irritation/correction	
Eye irritation/corrosion Inahaltion	Severe eye irritation. No data is available on the product itself.

LD50 - Rat (Ingestion)	> 2,000 mg/kg
Skin irritation/corrosion	Severe skin irritation. Corrosive to the skin of a rabbit.
	Severe skill illitation. Corrosive to the skill of a rappit.
Formaldehyde(50-00-0)	
Genotoxicity	Formaldehyde was found to be weakly mutagenic in a number of in vitro genotoxicity tests and positive in certain in vivo screening tests for mutagenicity. Formaldehyde did not cause birth defects in rats inhaling concentrations up to 10 ppm. However, a study using higher levels did show a slight but statistically significant reduction in male fetal body weight.
LD50 Dermal - Rabbit	270 mg/kg
LD50 Inhalation - Rat	0.31-0.59 mg/l (4 h) (Dust/ Mist)
LD50 Oral - Rat - Acute	100 mg/kg, Rat
toxicity	
Other Information	Lifetime inhalation of formaldehyde vapor at concentrations above 5 ppm for 6 hours per day, caused nasal tumors in laboratory animals. The International Agency for Research on Cancer (IARC) has classified formaldehyde as a Group 1 (known) human carcinogen based on epidemiological evidence linking formaldehyde exposure to the occurrence of nasopharyngeal cancer, a rare type of cancer. IARC also found limited evidence of cancer of the nasal cavity and paranasal sinuses and insufficient evidence for an association between formaldehyde and leukemia. Inhalation caused liver and kidney damage in laboratory animal tests.
Sensitization	Formaldehyde has been reported to cause pulmonary hypersensitivity in some individuals who were exposed to conceratrations know to cause irritation, however, no pulmonary
Chin/Free in 1991	sensitization has been demonstrated in laboratory animal studies.
Skin/Eye irritation	Can cause severe eye and moderate skin irritation.
Specific Target Organ Toxicity - Repeated exposure	Repeated skin exposure to solutions of 2% or more formaldehyde has caused skin allergic reactions.
Specific Target Organ Toxicity - Single	No data.
Isobutyl Alcohol(78-83-1)	
Carcinogenicity Data:	The ingredient(s) of this product is (are) not classified as carcinogenic by ACGIH, IARC, OSHA or NTP.
LC50 Inhalation - Rat	8000 ppm; (4 h)
LD50 Dermal - Rabbit	3400 mg/kg
LD50 Oral - Rat (Acute	2460 mg/kg
Toxicity)	3, 3
Mutagenicity Data:	No adverse mutagenicity effects are anticipated.
Reproductive Data:	No adverse reproductive effects are anticipated.
Respiratory / Skin Sensitization Data:	None known.
Synergistic Materials:	Alcohols may interact synergistically with chlorinated solvents (example - carbon tetrachloride, chloroform, bromotrichloromethane), dithiocarbamates (example - disulfiram), dimethylnitrosamine and thioacetamide.
Tetragenicity Data:	No adverse Tetragenicity effects are anticipated.
Methyl Amyl Ketone(110-43-0)	
Aspiration hazard	May be harmful if swallowed and enters airways.
Carcinogenicity	No data available.
LD50 Dermal - (Rat)	>2,000 mg/kg
LD50 Inhalation - (Rat)	>16.7 mg/l (4 h)
LD-50 Oral - (Rat)	1,600 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 sensitization	Skin Sensitization:, (Mouse) - non-sensitizing.
Serious eye damage/eye irritation	(Rabbit, 24 h): slight.
Skin corrosion/irritation	(Rabbit, 24 h): moderate.
Specific target organ toxicity -	No data available.
repeated exposure Specific target organ toxicity -	No data available.
single exposure	ito data avaliable.
Methyl Ethyl Ketone(78-93-3)	In 1 1 M. 1 1 . C17 . II
Aspiration toxicity	Product: May be harmful if swallowed and enters airways.
Carcinogenicity	Remarks: This information is not available, Carcinogenicity-Assement: Not classified as a human carcinogen.
Further information	
Turtilei illioittiation	Product Remarks: Symptoms of overexposure may be headache, diaainess, titedness, nausea and vomiting.,
Germ cell mutagenicity	Product Remarks: Symptoms of overexposure may be headache, diaainess, titedness,

	metabolic activation, Method OECD Test Guideline 471
LC50 (mouse) inhalation	320 mg/l (4 h exposure)
LC50 (rat) Oral	3737 mg/kg
LD50 (rabbit) dermal	6,480 mg/kg
Reproductive toxicity	Effects on fetal development, Species: rat female, Application Route: Inhalation, Dose: 400, 1000, 3000 ppm,
Respiratory or skin	Test Type: Buehler Test, Species guinea pig, Method OECD Test Guideline 406, Result:
sensitsation	Did not cause sensitization on laboratory animals.
Serious eye damage/ eye	Remarks: Severe skin irritation, Species rabbit, Exposure time 24 h, Result: Irritation to
irritation	eyes
Skin corrosion/irritation	Remarks: Moderate skin irritation, Species rabbit, Exposure time 24 h, Result: Mild skin irritation
STOT - repeated exposure	Product: No data available, Components: No data available.
STOT - single exposure	Product: Target Organs: Central Nervous system, Components: Exposure routes: Inhalation, Product: Target Organs: Central Nervous system
n-Butyl Acetate(123-86-4)	
Aspiration hazard	No data available.
Carcinogenicity	No data available.
Inhalation	No data available.
LD-50 Dermal - (Rabbit)	> 16ml/kg
LD-50 Oral - (Rat) Mutagenicity	14,130 mg/kg In vitro: No data available. In vivo: No data available.
Other adverse effects:	No data available. In vivo. No data available.
Repeated dose toxicity	No data available.
Reproductive toxicity	No data available.
Respiratory or skin sensitization	Skin Sensitization:, (Guinea Pig) - non-sensitizing.
Serious eye damage/eye irritation	(Rabbit, 24 h): none
Skin corrosion/irritation	(Rabbit, 24 h): none
Specific target organ toxicity -	No data available.
repeated exposure	
Specific target organ toxicity - single exposure	Narcotic effect.
P.M. Acetate(108-65-6)	
Aspiration hazard	No data available.
Carcinogenicity	No data available.
Carcinogenicity LC50 - Inhalation Rat	No data available. >4345 ppm (Rat, 6 h)
Carcinogenicity LC50 - Inhalation Rat LD50 - Dermal - Rabbit	No data available. >4345 ppm (Rat, 6 h) >5000 mg/kg
Carcinogenicity LC50 - Inhalation Rat LD50 - Dermal - Rabbit LD50 - Oral - Rat	No data available. >4345 ppm (Rat, 6 h) >5000 mg/kg 6,190 mg/kg
Carcinogenicity LC50 - Inhalation Rat LD50 - Dermal - Rabbit LD50 - Oral - Rat Mutagenicity	No data available. >4345 ppm (Rat, 6 h) >5000 mg/kg 6,190 mg/kg In vitro: No data available. In vivo: No data available.
Carcinogenicity LC50 - Inhalation Rat LD50 - Dermal - Rabbit LD50 - Oral - Rat Mutagenicity Other adverse effects	No data available. >4345 ppm (Rat, 6 h) >5000 mg/kg 6,190 mg/kg In vitro: No data available. In vivo: No data available. No data available.
Carcinogenicity LC50 - Inhalation Rat LD50 - Dermal - Rabbit LD50 - Oral - Rat Mutagenicity Other adverse effects Repeated dose toxicity	No data available. >4345 ppm (Rat, 6 h) >5000 mg/kg 6,190 mg/kg In vitro: No data available. In vivo: No data available. No data available. No data available.
Carcinogenicity LC50 - Inhalation Rat LD50 - Dermal - Rabbit LD50 - Oral - Rat Mutagenicity Other adverse effects	No data available. >4345 ppm (Rat, 6 h) >5000 mg/kg 6,190 mg/kg In vitro: No data available. In vivo: No data available. No data available. No data available. No data available.
Carcinogenicity LC50 - Inhalation Rat LD50 - Dermal - Rabbit LD50 - Oral - Rat Mutagenicity Other adverse effects Repeated dose toxicity Reproductive toxicity. Respiratory or skin sensitization	No data available. >4345 ppm (Rat, 6 h) >5000 mg/kg 6,190 mg/kg In vitro: No data available. In vivo: No data available. No data available. No data available.
Carcinogenicity LC50 - Inhalation Rat LD50 - Dermal - Rabbit LD50 - Oral - Rat Mutagenicity Other adverse effects Repeated dose toxicity Reproductive toxicity. Respiratory or skin	No data available. >4345 ppm (Rat, 6 h) >5000 mg/kg 6,190 mg/kg In vitro: No data available. In vivo: No data available. No data available. No data available. No data available.
Carcinogenicity LC50 - Inhalation Rat LD50 - Dermal - Rabbit LD50 - Oral - Rat Mutagenicity Other adverse effects Repeated dose toxicity Reproductive toxicity. Respiratory or skin sensitization Serious eye damage/eye	No data available. >4345 ppm (Rat, 6 h) >5000 mg/kg 6,190 mg/kg In vitro: No data available. In vivo: No data available. No data available. No data available. No data available. Skin Sensitization:, (Guinea Pig) - non-sensitizing (Rabbit): very slight Specified substance(s) 2-methoxy-1-methylethyl acetate (Rabbit, 4 h): none (Rabbit, 24 h): none.
Carcinogenicity LC50 - Inhalation Rat LD50 - Dermal - Rabbit LD50 - Oral - Rat Mutagenicity Other adverse effects Repeated dose toxicity Reproductive toxicity. Respiratory or skin sensitization Serious eye damage/eye irritation Skin corrosion/irritation Specific target organ toxicity -	No data available. >4345 ppm (Rat, 6 h) >5000 mg/kg 6,190 mg/kg In vitro: No data available. In vivo: No data available. No data available. No data available. No data available. Skin Sensitization:, (Guinea Pig) - non-sensitizing (Rabbit): very slight Specified substance(s) 2-methoxy-1-methylethyl acetate (Rabbit, 4 h): none (Rabbit, 24
Carcinogenicity LC50 - Inhalation Rat LD50 - Dermal - Rabbit LD50 - Oral - Rat Mutagenicity Other adverse effects Repeated dose toxicity Reproductive toxicity. Respiratory or skin sensitization Serious eye damage/eye irritation Skin corrosion/irritation Specific target organ toxicity - repeated exposure Specific target organ toxicity -	No data available. >4345 ppm (Rat, 6 h) >5000 mg/kg 6,190 mg/kg In vitro: No data available. In vivo: No data available. No data available. No data available. No data available. Skin Sensitization:, (Guinea Pig) - non-sensitizing (Rabbit): very slight Specified substance(s) 2-methoxy-1-methylethyl acetate (Rabbit, 4 h): none (Rabbit, 24 h): none.
Carcinogenicity LC50 - Inhalation Rat LD50 - Dermal - Rabbit LD50 - Oral - Rat Mutagenicity Other adverse effects Repeated dose toxicity Reproductive toxicity. Respiratory or skin sensitization Serious eye damage/eye irritation Skin corrosion/irritation Specific target organ toxicity - repeated exposure Specific target organ toxicity - single exposure	No data available. >4345 ppm (Rat, 6 h) >5000 mg/kg 6,190 mg/kg In vitro: No data available. In vivo: No data available. No data available. No data available. No data available. Skin Sensitization:, (Guinea Pig) - non-sensitizing (Rabbit): very slight Specified substance(s) 2-methoxy-1-methylethyl acetate (Rabbit, 4 h): none (Rabbit, 24 h): none. No data available. No data available.
Carcinogenicity LC50 - Inhalation Rat LD50 - Dermal - Rabbit LD50 - Oral - Rat Mutagenicity Other adverse effects Repeated dose toxicity Reproductive toxicity. Respiratory or skin sensitization Serious eye damage/eye irritation Skin corrosion/irritation Specific target organ toxicity - repeated exposure Specific target organ toxicity - single exposure Parachlorobenzotrifluoride(98-5	No data available. >4345 ppm (Rat, 6 h) >5000 mg/kg 6,190 mg/kg In vitro: No data available. In vivo: No data available. No data available. No data available. No data available. Skin Sensitization:, (Guinea Pig) - non-sensitizing (Rabbit): very slight Specified substance(s) 2-methoxy-1-methylethyl acetate (Rabbit, 4 h): none (Rabbit, 24 h): none. No data available. No data available.
Carcinogenicity LC50 - Inhalation Rat LD50 - Dermal - Rabbit LD50 - Oral - Rat Mutagenicity Other adverse effects Repeated dose toxicity Reproductive toxicity. Respiratory or skin sensitization Serious eye damage/eye irritation Skin corrosion/irritation Specific target organ toxicity - repeated exposure Specific target organ toxicity - single exposure Parachlorobenzotrifluoride(98-5 Additional Information	No data available. >4345 ppm (Rat, 6 h) >5000 mg/kg 6,190 mg/kg In vitro: No data available. In vivo: No data available. No data available. No data available. No data available. Skin Sensitization:, (Guinea Pig) - non-sensitizing (Rabbit): very slight Specified substance(s) 2-methoxy-1-methylethyl acetate (Rabbit, 4 h): none (Rabbit, 24 h): none. No data available. No data available. No data available. RTECS: XS9145000 To the best of our knowledge, the chemical, physical, and toxicological properties have not been thoroughly investigated.
Carcinogenicity LC50 - Inhalation Rat LD50 - Dermal - Rabbit LD50 - Oral - Rat Mutagenicity Other adverse effects Repeated dose toxicity Reproductive toxicity. Respiratory or skin sensitization Serious eye damage/eye irritation Skin corrosion/irritation Specific target organ toxicity - repeated exposure Specific target organ toxicity - single exposure Parachlorobenzotrifluoride(98-5 Additional Information	No data available. >4345 ppm (Rat, 6 h) >5000 mg/kg 6,190 mg/kg In vitro: No data available. In vivo: No data available. No data available. No data available. No data available. Skin Sensitization:, (Guinea Pig) - non-sensitizing (Rabbit): very slight Specified substance(s) 2-methoxy-1-methylethyl acetate (Rabbit, 4 h): none (Rabbit, 24 h): none. No data available. No data available. Sin Sensitization: (Rabbit, 4 h): none (Rabbit, 24 h): none. No data available. No data available.
Carcinogenicity LC50 - Inhalation Rat LD50 - Dermal - Rabbit LD50 - Oral - Rat Mutagenicity Other adverse effects Repeated dose toxicity Reproductive toxicity. Respiratory or skin sensitization Serious eye damage/eye irritation Skin corrosion/irritation Specific target organ toxicity - repeated exposure Specific target organ toxicity - single exposure Parachlorobenzotrifluoride(98-5 Additional Information	No data available. >4345 ppm (Rat, 6 h) >5000 mg/kg 6,190 mg/kg In vitro: No data available. In vivo: No data available. No data available. No data available. No data available. Skin Sensitization:, (Guinea Pig) - non-sensitizing (Rabbit): very slight Specified substance(s) 2-methoxy-1-methylethyl acetate (Rabbit, 4 h): none (Rabbit, 24 h): none. No data available. No data available. No data available. No data available. IARCS: XS9145000 To the best of our knowledge, the chemical, physical, and toxicological properties have not been thoroughly investigated. No data available. IARC: No component of this product present at levels greater than or equal to 0.1% is identified as probable, possible or confirmed human carcinogen by IARC. ACGIH: No component of this product present at levels greater than or equal to 0.1% is identified as a carcinogen or potential carcinogen by ACGIH. NTP: No component of this product present at levels greater than or anticipated carcinogen by NTP. OSHA: No component of this product present at levels greater than or
Carcinogenicity LC50 - Inhalation Rat LD50 - Dermal - Rabbit LD50 - Oral - Rat Mutagenicity Other adverse effects Repeated dose toxicity Reproductive toxicity. Respiratory or skin sensitization Serious eye damage/eye irritation Skin corrosion/irritation Specific target organ toxicity - repeated exposure Specific target organ toxicity - single exposure Parachlorobenzotrifluoride(98-5) Additional Information Aspiration hazard Carcinogenicity	No data available. >4345 ppm (Rat, 6 h) >5000 mg/kg 6,190 mg/kg In vitro: No data available. In vivo: No data available. No data available. No data available. No data available. Skin Sensitization:, (Guinea Pig) - non-sensitizing (Rabbit): very slight Specified substance(s) 2-methoxy-1-methylethyl acetate (Rabbit, 4 h): none (Rabbit, 24 h): none. No data available. No data available. No data available. No data available. IARCS: XS9145000 To the best of our knowledge, the chemical, physical, and toxicological properties have not been thoroughly investigated. No data available. IARC: No component of this product present at levels greater than or equal to 0.1% is identified as probable, possible or confirmed human carcinogen by IARC. ACGIH: No component of this product present at levels greater than or equal to 0.1% is identified as a carcinogen or potential carcinogen by ACGIH. NTP: No component of this product present at levels greater than or equal to 0.1% is identified as a carcinogen by NTP. OSHA: No component of this product present at levels greater than or equal to 0.1% is identified as a carcinogen by NTP. OSHA: No component of this product present at levels greater than or equal to 0.1% is identified as a carcinogen by OSHA.
Carcinogenicity LC50 - Inhalation Rat LD50 - Dermal - Rabbit LD50 - Oral - Rat Mutagenicity Other adverse effects Repeated dose toxicity Reproductive toxicity. Respiratory or skin sensitization Serious eye damage/eye irritation Skin corrosion/irritation Specific target organ toxicity - repeated exposure Specific target organ toxicity - single exposure Parachlorobenzotrifluoride(98-5) Additional Information Aspiration hazard Carcinogenicity Germ cell mutagenicity	No data available. >4345 ppm (Rat, 6 h) >5000 mg/kg 6,190 mg/kg In vitro: No data available. In vivo: No data available. No data available. No data available. No data available. Skin Sensitization:, (Guinea Pig) - non-sensitizing (Rabbit): very slight Specified substance(s) 2-methoxy-1-methylethyl acetate (Rabbit, 4 h): none (Rabbit, 24 h): none. No data available. No data available. No data available. 66-6) RTECS: XS9145000 To the best of our knowledge, the chemical, physical, and toxicological properties have not been thoroughly investigated. No data available. IARC: No component of this product present at levels greater than or equal to 0.1% is identified as probable, possible or confirmed human carcinogen by IARC. ACGIH: No component of this product present at levels greater than or equal to 0.1% is identified as a carcinogen or potential carcinogen by ACGIH. NTP: No component of this product present at levels greater than or equal to 0.1% is identified as a known or anticipated carcinogen by NTP. OSHA: No component of this product present at levels greater than or equal to 0.1% is identified as a 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 or potential carcinogen by OSHA. Human Embryo Unscheduled DNA synthesis.
Carcinogenicity LC50 - Inhalation Rat LD50 - Dermal - Rabbit LD50 - Oral - Rat Mutagenicity Other adverse effects Repeated dose toxicity Reproductive toxicity. Respiratory or skin sensitization Serious eye damage/eye irritation Skin corrosion/irritation Specific target organ toxicity - repeated exposure Specific target organ toxicity - single exposure Parachlorobenzotrifluoride(98-5) Additional Information Aspiration hazard Carcinogenicity Germ cell mutagenicity LD50 Oral - Rat	No data available. >4345 ppm (Rat, 6 h) >5000 mg/kg 6,190 mg/kg In vitro: No data available. In vivo: No data available. No data available. No data available. No data available. Skin Sensitization:, (Guinea Pig) - non-sensitizing (Rabbit): very slight Specified substance(s) 2-methoxy-1-methylethyl acetate (Rabbit, 4 h): none (Rabbit, 24 h): none. No data available. No data available. No data available. 6-6) RTECS: XS9145000 To the best of our knowledge, the chemical, physical, and toxicological properties have not been thoroughly investigated. No data available. IARC: No component of this product present at levels greater than or equal to 0.1% is identified as probable, possible or confirmed human carcinogen by IARC. ACGIH: No component of this product present at levels greater than or equal to 0.1% is identified as a carcinogen or potential carcinogen by ACGIH. NTP: No component of this product present at levels greater than or equal to 0.1% is identified as a carcinogen by NTP. OSHA: 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 by NTP. OSHA: No component of this product present at levels greater than or equal to 0.1% is identified as a carcinogen by NTP. OSHA: No component of this product present at levels greater than or equal to 0.1% is identified as a carcinogen by OSHA. Human Embryo Unscheduled DNA synthesis. 13,000 mg/kg Dermal: No data available.
Carcinogenicity LC50 - Inhalation Rat LD50 - Dermal - Rabbit LD50 - Oral - Rat Mutagenicity Other adverse effects Repeated dose toxicity Reproductive toxicity. Respiratory or skin sensitization Serious eye damage/eye irritation Skin corrosion/irritation Specific target organ toxicity - repeated exposure Specific target organ toxicity - single exposure Parachlorobenzotrifluoride(98-5) Additional Information Aspiration hazard Carcinogenicity Germ cell mutagenicity	No data available. >4345 ppm (Rat, 6 h) >5000 mg/kg 6,190 mg/kg In vitro: No data available. In vivo: No data available. No data available. No data available. No data available. Skin Sensitization:, (Guinea Pig) - non-sensitizing (Rabbit): very slight Specified substance(s) 2-methoxy-1-methylethyl acetate (Rabbit, 4 h): none (Rabbit, 24 h): none. No data available. No data available. No data available. 66-6) RTECS: XS9145000 To the best of our knowledge, the chemical, physical, and toxicological properties have not been thoroughly investigated. No data available. IARC: No component of this product present at levels greater than or equal to 0.1% is identified as probable, possible or confirmed human carcinogen by IARC. ACGIH: No component of this product present at levels greater than or equal to 0.1% is identified as a carcinogen or potential carcinogen by ACGIH. NTP: No component of this product present at levels greater than or equal to 0.1% is identified as a known or anticipated carcinogen by NTP. OSHA: No component of this product present at levels greater than or equal to 0.1% is identified as a 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 or potential carcinogen by OSHA. Human Embryo Unscheduled DNA synthesis.

sensitization	
	No data available.
Serious eye damage/eye irritation	No data available.
	No data continue
Skin corrosion/irritation	No data available.
Specific target organ toxicity -	No data available.
repeated exposure	
Specific target organ toxicity -	Inhalation - May cause respiratory irritation.
single exposure	
Styrene(100-42-5)	
Irritation / corrosion - Eye	Species: Rabbit; Result: non-irritant; Method: BASF - Test
Irritation / corrosion -	Species: Guinea pig; Result: non-sensitization; Method: OECD Guideline 406.
Sensitization	
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
TALC(14807-96-6)	
Acute toxicity - Dermal	No data available.
Acute toxicity - Inhalation	No data available.
Additional Information	RTECS: WW2710000 Prolonged inhalation of crystalline silica may result in silicosis, a
	disabling pulmonary fibrosis characterized by fibrotic changes and miliary 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.
	To the best of our knowledge, the chemical, physical, and toxicological properties have
	not been thoroughly investigated. Stomach - Irregularities - Based on Human Evidence
	Liver - Irregularities - Based on Human Evidence Stomach - Irregularities - Based on
	Human Evidence Liver - Irregularities - Based on Human Evidence (Quartz).
Aspiration hazard	No data available.
Carcinogenicity	Carcinogenicity - Rat - Inhalation Tumorigenic: Equivocal tumorigenic agent by RTECS
Carcinogenicity	criteria. Lungs, Thorax, or Respiration: Tumors. IARC: 1 - Group 1: Carcinogenic to
	humans (Quartz) IARC: 3 - Group 3: Not classifiable as to its carcinogenicity to humans
	(Hydrous magnesium silicate) 3 - Group 3: Not classifiable as to its carcinogenicity to
	humans (Hydrous magnesium silicate) 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 mutagenicity	No data available.
	No data available.
Reproductive toxicity Respiratory or skin	No data available.
sensitization	No data available.
Serious eye damage/eye	No data quallalla
, , ,	No data available.
irritation	Skin - Human Result: Mild skin irritation - 3 h
Skin corrosion/irritation	
Specific target organ toxicity -	No data available.
repeated exposure	N. J. L. William
Specific target organ toxicity -	No data available.
single exposure	
Titanium Dioxide(13463-67-7)	T 100 11 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
Carcinogenicity	In lifetime inhalation studies rats were exposed for 2 years to respectively 10, 50, 250
	mg/m3 of respirable Ti02.
Dermal ALD (rabbit)	>10000 mg/m3
Eye irritation	slight irritation
Inhalation 4 h ALC	>6.82 mg/l
ORAL ALD (rat)	>2400 mg/kg
Sensitsation	Did not cause sensitsation on laboratory animals.
Skin irritation	slight irritation

12. ECOLOGICAL INFORMATION

Acetone(67-64-1)	
Bioacculative potential	Parition coefficient: n-octanol/water: log Pow: -0.24
EC50 (Daphnia magna (Water	7,630 mg/l (Exposure time 48 h); Test substance: Acetone
flea))	
LC50 (Oncorhynchus mykiss	6,100 mg/l (Exposure time: 48 h)
(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 degrability	Biodegrability: Remarks: No data available
Toxicity to algae Aluminum Hydroxide(21645-51	Remarks: No data available
Bioaccumulative potential	Inert material.
EC50 - Daphnia - Toxicity to	>10,000 mg/l, Daphnia magna (Water flea) (OECD Test Guideline 202)
daphnia and other aquatic	- 10/000 mg/, 2 aprilla magna (mater mea) (0200 mea eataeille 202)
invertebrates	
EC50 - Fish - Toxicity to fish	>10,000 mg/l, Fish
Mobility in soil	Inert material.
NOEC - Toxicity to algae Other adverse effects	>0.004 mg/l, 72 h, Pseudokirchneriella subcapitata (algae) - (OECD Test Guideline 201) None known.
Persistence and degradability	Non-degradable
Amorphous Silica(7631-86-9)	Their degradable
Additional ecological	General notes: Do not allow product to reach ground water, water course or sewage
information	system.
Bioaccumulative potential	No further revelent 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 Mobility in soil	10000 mg/l (zebra fish) (96 h) (static) (OCED203) No further revelent information available.
Persistence and degrability	The product is chemically and biologically inert. By the insolubility in water there is a
reisisterice and degrability	separation at every filtration and sedimentation process.
Carbon Black(1333-86-4)	separation at every intration and seamentation processi
Behavior in water treatment	Activated sludge, EC0 (3 h) > 800 mg/L. DEV L3 (TTC test)
plants	
Bioaccumulation Potential	Potential bioaccumulation is not expected because of the physicochemical properties of
EC50 (Scenedesmus	the substance > 10,000 mg/L, OECD (Guideline 201)
subspicatus)	> 10,000 mg/L, OLCD (Guideline 201)
EC50 Daphnia magna	>5600 mg/l (24 h) OECD (Guideline 202)
(waterflea)	3, (, , : , : , ; ,
Environmental fate	Carbon black is an inert solid, stable and insoluble in water or organic solvents. Its vapour
	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 (Scenedesmus	> 10,000 mg/L, OECD (Guideline 201)
subspicatus) Cyclohexanone(108-94-1)	
Bioaccumulative potential	No data available.
EC50 - (Pimephales	527-732 mg/l, (Pimephales promelas (fathead minnow)) Exposure time: 96 h, Test types:
promelas)- Toxicity to fish	flow-through test.
EC50 - Daphnia magna -	>100 mg/l, exposure time 48 h, Test Type: static test, Method: OECD Test Guideline 202,
Toxicity to daphnia and other	GLP: yes.
aquatic invertebrates	> 100 mg/l (Desmadesmus subspicientus (Scanadesmus subspicatus)), and points Crowth
EC50 - Toxicity to algae	>100 mg/l (Desmodesmus subspicicatus (Scenedesmus subspicatus)), end point: Growth rate, Exposure time: 72 h, Test Type: static test, Analytical monitoring: yes, Method
	Guideline 201, GLP: yes.
Mobility in soil	No data available.
Other adverse effects	No data available.
Persistence and degradability	Biodegradation: >60%, Remarks: Readily biodegradable.
Results of PBT and vPvB	PBT/vPvB assessment not available as chemical safety assessment not required/not
assessment Dibutyltin Dilaurate(77-58-7)	conducted.
Aquatic toxicity	No data is available on the product itself.
Bioaccumulation	No data is available on the product itself.
EC50 - Daphnia	2.28 mg/l, Species : Daphnia magna.
LC50 - Fish	2 mg/l, Species : Fish.
Mobility	No data available.
Persistence and degradability	Biodegradability: No data is available on the product itself.
Toxicity to other organisms	Biodegradability: No data is available on the product itself. No data available.
Toxicity to other organisms Formaldehyde(50-00-0)	No data available.
Toxicity to other organisms	

LC50 Oncorhynchus - Toxicity to fish	100-136 mg/l, (96 h), Oncorhynchus mykiss	
Toxicity to Algae	Not available.	
Isobutyl Alcohol(78-83-1)		
Chronic	No data available.	
Degradability / Persistence; Biological / A biological Degradation	Evaluation: Not readily biodegradable (by OECD criteria).	
EC50 - Aquatic Plants	>100 mg/l (72 h) The product has not been tested. The statement has been derived from properties of the individual components.	
EC50 - Daphnia - Acute	>100 mg/l (48 h) The product has not been tested. The statement has been derived from properties of the individual components.	
LC50 - Fish - Acute	>100 mg/l (96 h) The product has not been tested. The statement has been derived from properties of the individual components.	
Microorganisms	Toxicity to microorganisms: bacteria EC10 (17 h): >750 mg/l. The product has not been tested. The statement has been derived from properties of the individual components.	
Methyl Amyl Ketone(110-43-0)		
Aquatic invertebrates	No data available.	
Bioaccumulative potential	No data available.	
Chronic Toxicity (Fish)	No data available.	
ErC50 (Selenastrum capricornutum)	98.2 mg/l, 72 h	
LC50 (Fathead Minnow) Acute toxicity	131 mg/l , (96 h)	
Mobility in soil	No data available.	
Persistence and degradability	69 % (28 d, Ready Biodegradability - CO2 in Sealed Vessels (Headspace Test)). Biological Oxygen Demand BOD-5: 1,770 mg/g BOD-20: 2,000 mg/g , Chemical Oxygen Demand:	
Results of PBT and vPvB	2,420 mg/g, BOD/COD ratio No data available. No data available.	
assessment		
Methyl Ethyl Ketone(78-93-3) Bioaccumulative potential	Partition coefficient: n-octanol/water: log Pow: 2.49	
EC50 (Algae)	2029 mg/l (48 h; Pseudokirchneriella subcapitata (Green Algae))	
EC50 (Algde)	308 mg/l (48 h; Daphnia magna (Water flea))	
LC50 (fish)	2993 mg/l (96 h; Pimephales promelas (Fathead minnow))	
Mobility in soil	No data available	
Other adverse effects	No data available	
Persistence and degradability	Biodegradability: Concentration: 2mg/l; Result: Readily biodegradation: 98%; Exposure 28 d;	
Product	Regulation: 40CFR Protection of Environment, Part 82 Protection of Stratospheric Ozone - CAA Section 602 Class 1 Substances:	
n-Butyl Acetate(123-86-4)		
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 Minnow) Acute Toxicity	18 mg/l, (96 h)	
LC-50 (Water Flea) Aquatic invertebrates	44 mg/l , (48 h)	
Mobility in soil	Known or predicted distribution to environmental compartments: No data available.	
Other adverse effects	No data available.	
Persistence and degradability	83 % (28 d), Biological Oxygen Demand:BOD-5: 730 mg/g, Chemical Oxygen Demand:1,010 mg/g, BOD/COD ratio:72 %.	
Results of PBT and vPvB assessment	No data available.	
P.M. Acetate(108-65-6)		
Aquatic invertebrates	NOEC (daphnia, 21 d): >= 100 mg/l EC-50 (daphnia, 21 d): > 100 mg/l	
Bioaccumulative potential	No data available.	
Biological Oxygen Demand	363 mg/g 1,050 mg/g	
Chemical Oxygen Demand Chronic Toxicity Fish	No data available. LC-50 (Oryzias latipes, 14 d): 63.5 mg/l NOEC (Oryzias latipes, 14 d): 47.5 mg/l	
LC50 - Daphnoid - Aquatic	408 mg/l (48 h)	
invertebrates LC50 - Fathead Minnow -	161 mg/l (96 h)	
Toxicity to Fish Mobility in soil	No data available.	
Other adverse effects	No data available. No data available.	
Persistence and degradability	Biodegradation - 90 % (28 d, Ready Biodegradability: CO2 Evolution Test) Readily	
	biodegradable Page 12 of 16	

Results of PBT and vPvB	No data available.	
assessment		
Toxicity to Aquatic Plants	EC-50 (Selenastrum capricornutum, 96 h): > 1,000 mg/l NOEC (Selenastrum capricornutum, 96 h): >= 1,000 mg/l	
Parachlorobenzotrifluoride(98-5		
Bioaccumulative potential	No data available.	
Mobility in soil	No data available.	
Other adverse effects	No data available.	
Persistence and degradability	No data available.	
Results of PBT and vPvB	PBT/vPvB assessment not available as chemical safety assessment not required/not	
assessment	conducted.	
Toxicity	No data available.	
Styrene(100-42-5)		
Bioaccumulation	At present state of knowledge, no negative ecological effects are expected.	
Chronic	No data available regarding toxicity to Daphnis.	
Chronic	No data available regarding toxicity to fish.	
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 idus)	>100 mg/l (96 h)	
Acute		
Microorganisms	Toxicity to microorganisms: The inhibition of the degradation activity sludge is not anticipated when introduced to biological treatment plants in appropriate low	
TALC(14907.06.6)	conceratrations.	
TALC(14807-96-6)	No data available	
Bioaccumulative potential	No data available. No data available.	
Mobility in soil Other adverse effects	No data available.	
Persistence and degradability	No data available.	
Results of PBT and vPvB	PBT/vPvB assessment not available as chemical safety assessment not required/not	
assessment	conducted	
Toxicity	No data available.	
Titanium Dioxide(13463-67-7)	ויט עמגמ מימוומטוכ.	
LC50 fish	Fathead minnow 96 h >1000 mg/l	
LCJU IISII	Tauleau Hillinow 50 H >1000 Hig/T	

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 incenerate. 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 RESTRICTIONS THAT MAY APPLY.

USDOT GROUND

DOT (DEPARTMENT OF TRANSPORTATION) PROPER SHIPPING NAME (DOT): Paint

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 HAZARDS CLASS : 3

UN/NA NUMBER: UN1263
PACKING GROUP: PG II

EMERGENCY RESPONSE GUIDE (ERG): 128

IMDG (OCEAN)

PROPER SHIPPING NAME: Paint

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, Fire hazard, Chronic health hazard, Moderate skin irritant, Moderate eye irritant,

Carcinogen.

EPCRA - Emergency

CERCLA REPORTABLE QUANTITY

This product contains:	Chemical CAS#
n-Butyl Acetate	123-86-4
Carbon Black	1333-86-4
Methyl Ethyl Ketone	78-93-3
Isobutyl Alcohol	78-83-1

SARA 304 Extremely Hazardous Substances Reportable Quantity: This material does not contain any components with a section 304 EHS RQ.

SARA TITLE III (SUPERFUND AMENDMENTS 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
Titanium Dioxide	13463-67-7
Parachlorobenzotrifluoride	98-56-6
P.M. Acetate	108-65-6
Amorphous Silica	7631-86-9
n-Butyl Acetate	123-86-4
Carbon Black	1333-86-4

CLEAN AIR ACT:

This product contains:	Chemical CAS#
Styrene	100-42-5
Formaldehyde	50-00-0

INTERNATIONAL REGULATIONS

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

Flam. Liq. Cat. 2; H226 Eye Irrit. Cat. 2; H319 STOT SE 3 H336

NATIONAL REGULATIONS

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

STATE REGULATIONS CALIFORNIA PROPOSITION 65

This product contains:	Chemical CAS#
*Talc	14807-96-6

- *This product contains (a) chemical (s) known to the State of California to cause cancer.
- #This product contains (a) chemical (s) known to the State of California to be carcinogenic.
 +This product contains (a) chemical (s) known to the State of California to cause birth defects or other reproductive harm.

Massachusetts Right to Know

This product contains	Chemical CAS#
Acetone	67-64-1
Parachlorobenzotrifluoride	98-56-6
n-Butyl Acetate	123-86-4
Silica Gel	112926-00-8
Talc	14807-96-6
Carbon Black	1333-86-4
Methyl Amyl Ketone	110-43-0
Methyl Ethyl Ketone	78-93-3
Isobutyl Alcohol	78-83-1
Cyclohexanone	108-94-1

Pennsylvania Right to Know

This product contains	Chemical CAS#
Acetone	67-64-1
Titanium Dioxide	13463-67-7
Parachlorobenzotrifluoride	98-56-6
P.M. Acetate	108-65-6
Amorphous Silica	7631-86-9
n-Butyl Acetate	123-86-4
Silica Gel	112926-00-8
Aluminum Hydroxide	21645-51-2
Talc	14807-96-6
Carbon Black	1333-86-4
Methyl Amyl Ketone	110-43-0
Methyl Ethyl Ketone	78-93-3
Isobutyl Alcohol	78-83-1
Dibutyltin Dilaurate	77-58-7
Cyclohexanone	108-94-1

New Jersey Right to Know

This product contains	Chemical CAS#
Acetone	67-64-1
Titanium Dioxide	13463-67-7
Parachlorobenzotrifluoride	98-56-6
P.M. Acetate	108-65-6
Amorphous Silica	7631-86-9
n-Butyl Acetate	123-86-4
Silica Gel	112926-00-8
Aluminum Hydroxide	21645-51-2

Tlac	14807-96-6
Carbon Black	1333-86-4
Methyl Amyl Ketone	110-43-0
Methyl Ethyl Ketone	78-93-3
Isobutyl Alcohol	78-83-1
Dibutyltin Dilaurate	77-58-7
Cyclohexanone	108-94-1

16. OTHER INFORMATION

Other Product Information

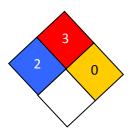
% Volatile by Volume: 46.76 % Volatile by Weight: 27.34 % Solids by volume: 53.24 % Solids by Weight: 72.66 % Exempt by Volume: 38.75 % Exempt by Weight: 22.46

VOC CONTENT: Excluding Exempt VOC: 120 Including Exempt VOC: 75

HMIS RATING

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

NFPA CODES



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