SAFETY DATA SHEET



 DATE ISSUED :
 8/10/2018

 SDS REF. No :
 2059-CLE18773

2059-CLE18773 VINYL NITRO SEALER GL.LAC. MUSICAL INSTRUMENTS

1. PRODUCT AND COMPANY IDENTIFICATION

PRODUCT NAME: 2059-CLE18773 VINYL NITRO SEALER GL.LAC. MUSICAL INSTRUMENTS

PRODUCT CODE: PRODUCT USE: 2059-CLE18773 Industrial Solventborne Paint

MANUFACTURER

Cardinal Industrial Finishes 1329 Potrero Ave

24 HR. EMERGENCY TELEPHONE NUMBER CHEMTREC (US Transportation): (800)424-9300 CHEMTREC (International : 1(202)483-7616 Transportation) WEB: WWW.CARDINALPAINT.COM

S. El Monte, CA, 626 444-9274

2. HAZARDS IDENTIFICATION

PICTOGRAMS



SIGNAL WORD : DANGER

HAZARD STATEMENTS :

H226 Flammable liquid and vapor.

H301+H331+H331 Toxic if swallowed, in contact with skin or inhaled.

H302+H332 Harmful if swallowed or inhaled.

H304 May be fatal if swallowed and enters airways.

- H312 Harmful if contact with skin.
- H315 Causes skin irritation.
- H319 Causes serious eye irritation.
- H335 May cause respiratory irritation.
- H336 May cause drowsiness or dizziness.
- H351 Suspected of causing cancer.

H360 May damage fertility or unborn child.

- H401 Toxic to aquatic life.
- H402 Harmful to aquatic life.

H412 Harmful to aquatic life with long lasting effects.

PRECAUTIONARY STATEMENTS : P210 Keep away from heat/sparks/open flames/hot surfaces. No smoking.

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
Ethyl Alcohol	35% - 40%	64-17-5
n-Butyl Acetate	15% - 20%	123-86-4
Aliphatic Solvent	10% - 15%	64742-47-8
Acetone	5% - 10%	67-64-1
Isopropyl Alcohol	5% - 10%	67-63-0
Pseudocumene	5% - 10%	95-63-6
Methyl Isobutyl Ketone	1% - 5%	108-10-1
Methyl Amyl Ketone	1% - 5%	110-43-0
Methyl Alcohol	1% - 5%	67-56-1
Dioctyl Phthalate	1% - 5%	117-81-7
n-Methyl-2-pyrrolidone	1% - 5%	872-50-4
Phenylethane	0.10% - 0.50%	100-41-4

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
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
Aliphatic Solvent(64742-47-8)		
USA ACGIH	ACGIH (TLV) TWA	200 mg/m3
USA NIOSH	NIOSH REL (ST)	10 mg/m3
USA NIOSH	NIOSH REL (TWA)	5 mg/m3
USA OSHA	OSHA OEL (TLV) TWA Table Z-1	500 ppm, 2,000 mg/m3
USA OSHA	OSHA OEL Table Z-1	5 mg/m3
Cumene(98-82-8)		
USA ACGIH	ACGIH (TLV) TWA	50 ppm
USA NIOSH	NIOSH (TWA) REL	50 ppm, 245 mg/m3
USA OSHA	OSHA (TWA) Table Z-1	50 ppm, 245 mg/m3
Dioctyl Phthalate(117-81-7)		
ACGIH TWA	TWA	5 mg/m3
NIOSH REL ST	ST	10 mg/m3
NIOSH REL TWA	TWA	5 mg/m3
OSHA P0 STEL	STEL	10 mg/m3
OSHA PO TWA	TWA	5mg/m3
OSHA Z-1 TWA	TWA	5 mg/m3
Ethyl Alcohol(64-17-5)		
USA ACGIH	ACGIH TWA (TLV)	1,000 ppm
USA NIOSH	NIOSH TWA	1,000 ppm, 1,900 mg/m3
USA OSHA	OSHA TWA (Table Z-1)	1,000 ppm, 1,900 mg/m3
Ethylene glycol mono butyl ether(111-76-2)		
USA ACGIH	ACGIH TWA (ppm)	20 ppm
USA NIOSH	NIOSH REL (ppm)	5 ppm
USA OSHA	OSHA PO TWA (ppm)	25 ppm
USA OSHA	OSHA TABLE Z-1 TWA (mg/m3)	50 ppm, 240 mg/m3

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USA NIOSH REL NIOSH TWA (ppm) 50 ppm USA OSHA OSHA TWA (ppm) 100 ppm n-Butyl Acetate(123-86-4) USA ACGIH ACGIH STEL 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) V/E USA OSHA OSHA TWA N/E USA ACGIH ACGIH PEL N/E USA OSHA OSHA TWA N/E USA ACGIH ACGIH STEL 125 ppm USA ACGIH ACGIH STEL 125 ppm	USA NIOSH REL	NIOSH STEL (ppm)	75 ppm	
USA OSHA OSHA TWA (ppm) 100 ppm 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 N/E USA OSHA OSHA TWA N/E USA ACGIH ACGIH PEL N/E USA OSHA OSHA TWA N/E USA ACGIH ACGIH STEL 125 ppm USA ACGIH ACGIH STEL 125 ppm	USA NIOSH REL	NIOSH TWA (ppm)	50 ppm	
n-Butyl Acetate(123-86-4) ACGIH STEL 200 ppm 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) V/E USA ACGIH ACGIH PEL N/E USA OSHA OSHA TWA N/E USA ACGIH ACGIH PEL N/E USA ACGIH ACGIH STEL 125 ppm USA ACGIH ACGIH STEL 125 ppm	USA OSHA	OSHA TWA (ppm)	100 ppm	
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 N/E USA ACGIH ACGIH PEL N/E USA OSHA OSHA TWA N/E USA ACGIH ACGIH STEL 125 ppm USA ACGIH ACGIH STEL 125 ppm	n-Butyl Acetate(123-86-4)			
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 N/E USA ACGIH ACGIH PEL N/E USA OSHA OSHA TWA N/E Phenylethane(100-41-4) USA ACGIH 125 ppm USA ACGIH ACGIH STEL 125 ppm	USA ACGIH	ACGIH STEL	200 ppm	
USA OSHA OSHA PEL (Table Z-1) 150 ppm, 710 mg/m3 n-Methyl-2-pyrrolidone(872-50-4) USA ACGIH N/E USA ACGIH ACGIH PEL N/E USA OSHA OSHA TWA N/E Phenylethane(100-41-4) ACGIH STEL 125 ppm USA ACGIH ACGIH TWA 20 npm	USA ACGIH	ACGIH TWA	150 ppm	
n-Methyl-2-pyrrolidone(872-50-4) N/E USA ACGIH ACGIH PEL N/E USA OSHA OSHA TWA N/E Phenylethane(100-41-4) USA ACGIH 125 ppm USA ACGIH ACGIH TWA 20 ppm	USA OSHA	OSHA PEL (Table Z-1)	150 ppm, 710 mg/m3	
USA ACGIH ACGIH PEL N/E USA OSHA OSHA TWA N/E Phenylethane(100-41-4) USA ACGIH ACGIH STEL USA ACGIH ACGIH TWA 20 ppm	n-Methyl-2-pyrrolidone(872-50-4)			
USA OSHA OSHA TWA N/E Phenylethane(100-41-4) USA ACGIH 125 ppm USA ACGIH ACGIH STEL 125 ppm	USA ACGIH	ACGIH PEL	N/E	
Phenylethane(100-41-4) ACGIH STEL 125 ppm USA ACGIH ACGIH TWA 20 ppm	USA OSHA	OSHA TWA	N/E	
USA ACGIH ACGIH STEL 125 ppm	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	100 ppm, 435 mg/m3	
USA NIOSH NIOSH REL (ST) 125 ppm, 545 mg/m3	USA NIOSH	NIOSH REL (ST)	125 ppm, 545 mg/m3	
USA OSHA OSHA STEL 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	USA OSHA	OSHA TWA (Table Z-1)	100 ppm, 435 mg/m3	
Phosphoric Acid(7664-38-2)	Phosphoric Acid(7664-38-2)			
USA ACGIH ACGIH (TLV) STEL 3 mg/m3	USA ACGIH	ACGIH (TLV) STEL	3 mg/m3	
USA ACGIH ACGIH (TLV) TWA 1 mg/m3	USA ACGIH	ACGIH (TLV) TWA	1 mg/m3	
USA NIOSH NIOSH (TWA) REL 1 mg/m3	USA NIOSH	NIOSH (TWÁ) REL	1 mg/m3	
USA NIOSH NIOSH (TWA) ST 3 mg/m3	USA NIOSH	NIOSH (TWA) ST	3 mg/m3	
USA OSHA OSHA (TWA) Table Z-1 1 mg/m3	USA OSHA	OSHA (TWA) Table Z-1	1 mg/m3	
Pseudocumene(95-63-6)	Pseudocumene(95-63-6)			
USA NIOSH NIOSH (TWA) REL 25 ppm, 125 ma/m3	USA NIOSH	NIOSH (TWA) REL	25 ppm, 125 mg/m3	
Xylene(1330-20-7)	Xylene(1330-20-7)			
USA ACGIH ACGIH STEL 150 ppm	USA ACGIH	ACGIH STEL	150 ppm	
USA ACGIH ACGIH TWA 100 ppm	USA ACGIH	ACGIH TWA	100 ppm	
USA OSHA QSHA TWA (Table Z-1) 100 PPM, 435 mg/m3	USA OSHA	OSHA TWA (Table Z-1)	100 PPM, 435 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

:	Liquid
:	Various colors depending on the pigmentation.
:	Characteristic. Sweet. Mint like.
:	No data available.
:	N/A – See Technical Data Sheet
:	Slower Than Ether
:	-94.7 C (-138.46 F)
:	No data available.
:	133.0 deg F TO 397.0 deg F
:	-4.00 deg F
:	.3
:	36.5
:	185 mm Hg
:	Heavier than air
:	No data available.
:	6.9792
:	No data available.
:	No data available.
:	No data available.
:	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 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 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.

440, 2200, 11,000 ppm; Frequency of Treatment: 7 days/week; General Toxicity NOAEC: 2,200 ppm; Tetragenicity: NOAEC: 2,200 ppm; Embryo-fetal toxicity:: No ppm; Result: No teratogenic potential. GLP: No data available.; Reproductive toxic	Dose: 0, Material: DAEC: 2,200
Assessment: Did not show teratogenic effects in animal experiments. Respiratory or skin Test type: Maximization test, Species: guinea pig, Assessment: Does not cause skin	in
sensitisation sensitization. Result: Did not cause sensitization on laboratory animals.	i an i
damage/eye irritation Irritating to eyes, Remarks: Eye irritation.	lon:
Skin Species: rabbit, Exposure time: 24 h, Classification: Not irritating to skin, Method corrosion/irritation Result: Mild irritation, Remarks: Repeated or prolonged contact with the mixture r removal natural fat from the skin resulting in desiccation of the skin.	In vivo, nay cause
STOT - single exposure Exposure routes: Inhalation (vapor); Assessment: May cause drowsiness or dizzin	ess.
STOT- repeated No data available. exposure	
Aliphatic Solvent(64742-47-8)	
Acute Dermal toxicity No data available.	
Acute Inhalation No data available. toxicity	
Acute toxicity No data available.	
Additional Information RTECS: Not available Prolonged or repeated exposure to skin causes defatting and dermatitis., To the best of our knowledge, the chemical, physical, and toxicologica have not been thoroughly investigated.	l properties
Aspiration hazard No data available.	
Carcinogenicity IARC: 3 - Group 3: Not classifiable as to its carcinogenicity to humans (Distillates (petroleum), hydrotrated light, kerosene - unspecified) NTP: No component of this present at levels greater than or equal to 0.1% is identified as a known or anticipa carcinogen by NTP. OSHA: No component of this product present at levels greater	product ted than or
Germ cell mutagenicity Reverse mutation assay S tynhimurium Result: negative	
Reproductive toxicity No data available.	
Respiratory or skin Draize Test - Guinea pig Result: Does not cause skin sensitization. sensitization Draize Test - Guinea pig Result: Does not cause skin sensitization.	
Serious eye Eyes - Rabbit Result: No eye irritation damage/eye irritation	
Skin Skin - Rabbit Result: No skin irritation - 4 h	
Specific target organ No data available. toxicity - repeated exposure	
Specific target organ toxicity - single	
Cumene(98-82-8)	
Additional Information RTECS: GR8575000	
Aspiration hazard No data available.	
Carcinogenicity Carcinogenicity IARC: 2B - Group 2B: Possibly carcinogenic to humans (Cumene) component of this product present at levels greater than or equal to 0.1% is identicarcinogen or potential carcinogen by ACGIH. NTP: No component of this product levels greater than or equal to 0.1% is identified as a known or anticipated carcinogen NTP. OSHA: No component of this product present at levels greater than or equal identified as a carcinogen or potential carcinogen by OSHA.	ACGIH: No ified as a present at ogen by to 0.1% is
Dermal No data available.	
Germ cell mutagenicity invitro assay, S. typhimurium, Result: negative	
Inhalation: No data available.	
LD50 Oral - Rat - 2,260 mg/kg, Acute toxicity	
Reproductive toxicity No data available.	
Respiratory or skin Guinea pig - Result: No skin irritation. (OECD Test Guideline 406) sensitization	
Serious eye Eyes - Rabbit Result: No skin irritation. (OECD Test Guideline 405) damage/eye irritation	
Skin Skin - Rabbit Result: No skin irritation. (OECD Test Guideline 404) corrosion/irritation	
Specific target organ No data available. toxicity - repeated exposure	

Specific target organ	No data available.
toxicity - single	
exposure	
Dioctyl Phthalate(117-8:	1-7)
^ - Vivo	Test Type: DNA Damage and/or repair. Test Species: Rat (male) Cell Type: Bone Marrow. Application Route: Oral. Exposure Time: 5 d. Dose: 0, 0.5, 1.7, 5 ml/kg. Result : Negative
Acute Inhalation Toxicity	Remarks: No Data Available
Aspiration Toxicity	No aspiration toxicity classification
Carcinogenicity	Mouse (Male of Female) Oral - Exposure time (104wk) Dose: 0, 500, 1500, 6000 ppm. NOAEL: 500 ppm. LOAEL: 1,500 ppm. Method: OECD Test Guideline 453. Result: Evidence of Carcinogenic Activity. Symptoms: Hepatocellular carcinoma and adenoma.
Carcinogenicity - Assessment	Suspected Human Carcinogens
Effects on fetal development	Rat - Application Route: Oral. Dose: 0, 357, 666, 586, 1055 mg/kg Duration of Single Treatment: 21 d. General Toxicity Maternal: NOAEL: 357 mg/kg body weight. Teratogenicity: NOAEL: 1,055 mg/kg body weight. Developmental Toxicity: NOAEL: 357 mg/kg body weight. Symptoms: Maternal toxicity, reduced number of viable fetuses, Preimplantation loss. Result: Embryotoxicity effects and adverse effects on the offspring were detected. Species: Rat. Application Route: Oral. Dose: 0, 40, 200, 1000 mg/kg bw/d. Duration of Single Treatment: 10 d. General Toxicity Maternal: NOAEL: 200 mg/kg bw. Teratogenicity: NOAEL: 200 mg/kg
	bw. Developmental Toxicity: NOAEL: 200 mg/kg bw. Symptoms: Maternal toxicity, skeletal malformations, visceral malformations. Result: Teratogenic effects.
Germ Cell Mutagenicity - Assessment	Tests on bacterial or mammalian cell cultures did not show mutagenic effects
Germ Cell Mutagenicity - Vitro	Test Type: Chromosome aberration test n vitro. Test Species: Chinese hamster ovary (CHO) Metabolic Activation With and without metabolic activation. Result: Negative. Test Type: Chromosome aberration
LD50 Acute Dermal Toxicity - Rabbit	25,000 mg/kg
LD50 Acute Oral Toxicity - Bat	5,000 mg/kg
Repeated Dose Toxicity	Rat (male and female) Oral - 104 wks Number of Exposures: 7d/wk. Dose: 0, 100, 500, 2500, 12500 ppm. OECD Test Guideline 453. GLP: Yes. Symptoms: Liver effects and increased liver weight. Rat (male and female) NOAEL: 50. Inhalation (dust/mist/fume) 4wks. Number of Exposures: 6h/d, 5d/wk. Dose: 0, 10, 50, 1000 mg/3. OECD Test Guideline 412. GLP: Yes. Symptoms: Lung effects and increased liver weight
Reproductive Toxicity - Assessment	Presumed human reproductive toxicant
Reproductive Toxicity - Effects on Fertility	Rat (Male and Female) Multigenerational Study - Application Route: Oral. Dose: 1.5 - 10,000 ppm. General Toxicity - Parent: NOAEL: 300 ppm. General ToxicityF1: NOAEL: 300 ppm. Fertility: NOAEL: 1,000 ppm. Early Embryonic effects and adverse effects development: NOAEL: 100 ppm. Results: Embryotoxicity Effects and adverse effects on the offspring were detected. GLP: Yes. Symptoms: Reduced fertility Testicular effects.
Respiratory or Skin	Guinea Pig - Maximization Test - Results: Did not cause sensitization on laboratory animals
Serious Eye	Rabbit - Exposure time (72h) OECD Test Guideline 405 - Results: Mild Eye Irritation
Skin	Rabbit - Exposure time (4h) OECD Test Guideline 404 - Results: Mild Skin Irritation
STOT - Repeated	No Data Available
STOT - Single	No Data Available
Ethyl Alcohol(64-17-5)	
Additional Information	RTECS: KQ6300000 Central nervous system depression, narcosis. Damage to the heart To
	the best of our knowledge, the chemical, physical, and toxicological properties have not been thoroughly investigated. Heart - Irregularities - Based on Human Evidence Stomach - Irregularities - Based on Human Evidence
Aspiration hazard	No data available.
Carcinogenicity - Mouse - Oral	Tumorigenic: Equivocal tumorigenic agent by RTECS criteria. Liver: Tumors. Blood: Lymphomas including Hodgkin's disease. 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. NTP: No components 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 components 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	No data available.
LC50 Inhalation - Rat	20000 ppm, (10 h)

LD50 Oral - Rat	7,060 mg/kg , Remarks: Lungs, Thorax, or Respiration: Other changes.
Reproductive toxicity	No data available. Reproductive toxicity - Human - female - Oral Effects on Newborn: Apgar
	score (human only). Effects on Newborn: Other neonatal measures or effects. Effects on
	Newborn: Drug dependence.
Respiratory or skin	No data available.
Serious ave	Results Mild ave irritation 24 h (OECD Test Cuideline 40E)
damage/eve irritation	Result: Mild eye initiation - 24 if (OECD Test Guideline 405)
Eves Rabbit	
Skin	Result: No skin irritation - 24 h (OECD Test Guideline 404)
corrosion/irritation	
Skin - Rabbit	
Specific target organ	No data available.
toxicity - repeated	
exposure	
Specific target organ	No data available.
toxicity - single	
Ethylong glycol mong by	the other (111, 76, 2)
Aspiration toxicity	Romarks: No data available
Carcinogenicity	Remarks, No usite Available.
carcinogenicity	Frequency of Treatment' 5 days/week NAOFI ' 125 ppm Result' Limited evidence of
	carcinogenic effects with no relevance to humans., Carcinogenicity-Assement: Not evidence of
	carcinogenicity in animal studies.
Further information	Product Remarks: Symptoms of overexposure may be headache, dizziness, tiredness, nausea
	and vomiting.,
Germ cell mutagenicity	Genotoxicity in vitro: Test Type: Mammalian cell gene mutation assay; Test species: Chinese
	hamster (CHO), Metabolic activation: with and without metabolic activation. Result: negative.,
	Genotoxicity in vivo: Test Type: In vivo micronucleus test., Test species:: mouse (male),
	application Route: Intraperitoneal, Result: negative, Germ cell mutagenicity Assessment:
LCEQ (rat) inhalation	Acute inhelation toxicity: 500 ppm. Expecting 4 by Accessment: the component/mixture
	is moderately toxic after short term inhalation
LC50 (rat) Oral	Acute toxicity estimate: 500 mg/kg: Method: Expert judgment.: Assessment: the
	component/mixture is moderately toxic after single ingestion.
LD50 (rat) dermal	Acute toxicity estimate: 1,1000 mg/kg; Method: Expert judgment; Assessment: the
	component/mixture is moderately toxic after single contact with skin.
Repeated dose toxicity	Species: rat NOAEL: 30, Application Route: Inhalation Exposure time: 14 wk Number of
	exposures: 6 h/d, 5 d/wk.
Reproductive toxicity	Effects on fertility : lest Type: Two-generation study Species: mouse Application Route: oral
	fertility at maternally toxic doses Effects on fetal development - Test Type: Embryo-fetal
	development Species: rat Application Boute: Inhalation Duration of Single Treatment: 10 d
	Frequency of Treatment: 6 hr/day Developmental Toxicity: Lowest observed adverse effect
	level: 100 ppm Result: Developmental toxicity occurred at maternal toxicity dose levels
	Reproductive toxicity - Assessment : No evidence of adverse effects on sexual function and
	fertility, and on development, based on animal experiments
Respiratory or skin	Test Type: Maximization test, Species guinea pig, Result: Did not cause sensitization on
Sensitivation	laboratory animals.
Serious eye damage/	Species rappin, exposure time 24 n, Result: Irritating to eyes.
Skin	Remarks: Moderate skin irritation in suscentible persons. Species rabbit. Exposure time 24 b
corrosion/irritation	Result: Mild skin irritation
STOT - repeated	No data available.
exposure	
STOT - single exposure	No data available.
Isobutyl Alcohol(78-83-1	<u>()</u>
Carcinogenicity Data:	The ingredient(s) of this product is (are) not classified as carcinogenic by ACGIH, IARC, OSHA
LCED Inhalation Dat	00 nnm: (4 h)
LCOU IIIIdidUUII - KAL	3400 mg/kg
LD50 Derinai - Kabbil	2460 mg/kg
Toxicity)	
Mutagenicity Data:	No adverse mutagenicity effects are anticipated.
Reproductive Data:	No adverse reproductive effects are anticipated.
Respiratory / Skin	None known.
Sensitization Data:	
Synergistic Materials:	Alcohols may interact synergistically with chlorinated solvents (example - carbon
	tetrachloride, chloroform, bromotrichloromethane), dithiocarbamates (example - disulfiram),
	dimethyInitrosamine and thioacetamide.

Tetragenicity Data:	No adverse Tetragenicity effects are anticipated.
Isopropyl Alcohol(67-63	-0)
Aspiration hazard	Based on physico-chemical values or lack of human evidence, not classified.
Carcinogenicity	Not classified.
Effects on	Not classified.
Development	
Germ cell mutagenicity	Not classified No adverse effect observed.
LC50 (Rat)	46.6 mg/l; Exposure time: 8 h, Acute inhalation toxicity: Based on acute toxicity values, not
	classified. High vapor concentrations may cause irritation of the eyes, nose, and/or throat,
	changes to the liver, lung, spieen, and brain, and central nervous system depression (ataxia, diazing participation with requirement and death in cases of
	uiziness, narcosis, and muscle relaxation, with respiratory arrest and death in cases of
LD50 (Rabbit)	12 870 mg/kg
1D50 (Rabbit)	4 396 mg/kg. Acute oral toxicity: Based on acute toxicity values not classified. Ingestion may
	cause gastrointestinal effects (pain, nausea, vomiting, and hemorrhage), hypothermia.
	cardiac effects (low blood pressure, shock and cardiac arrest), liver changes, kidney damage,
	and CNS effects (headache, dizziness, sleepiness, coma and death).
Reproductive toxicity	Effects on fertility / Effects on or via lactation: Not classified.
Respiratory or skin	Not classified No adverse effect observed.
sensitization	
Serious eye	Classified Causes serious eye irritation.
aamage/eye irritation	Describer altig instantion and an anti-stantification of the state of the state of the state of the state of the
Skin	Based on skin irritation values, not classified. Liquid may cause slight skin irritation. Exposure
Target Organ Systemic	Based on repeated exposure toxicity values not classified
Toyicant - Repeated	based on repeated exposure toxicity values, not classified.
exposure	
Target Organ Systemic	Routes of exposure: Ingestion, Inhalation Target Organs: Central pervous system Classified.
Toxicant - Single	May cause drowsiness or dizziness.
exposure	
Methyl Alcohol(67-56-1)	
Additional Information	RTECS: PC1400000 Methyl alcohol may be fatal or cause blindness if swallowed. Effects due
	to ingestion may include:, Headache, Dizziness, Drowsiness, metabolic acidosis, Coma,
	Seizures. Symptoms may be delayed., Damage of the:, Liver, Kidney Central nervous system
	- Breathing difficulties - Based on Human Evidence Stomach - Irregularities - Based on
	Human Evidence.
Aspiration hazard	No aspiration toxicity classification
Carcinogenicity	IARC: No component of this product present at levels greater than or equal to 0.1% is
	component of this product present at levels greater than or equal to 0.1% is identified as a
	carcinogen or notential 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. Reproductive toxicity Damage to
	fetus not classifiable Fertility classification not possible from current data. Specific target
	organ toxicity - single exposure Causes damage to organs.
Germ cell mutagenicity	Ames test S. typhimurium Result: negative in vitro assay fibroblast Result: negative Mutation
	in mammalian somatic cells. Mutagenicity (in vivo mammalian bone-marrow cytogenetic test,
ICEO Inhalation Bat	
I D50 Dermal - Pabbit	300 mg/kg
1D50 Oral - Rat Acute	100 mg/kg
Toxicity	
Reproductive toxicity	Damage to fetus not classifiable Fertility classification not possible from current data.
Respiratory or skin	Maximization Test (GPMT) - Guinea pig Does not cause skin sensitization. (OECD Test
sensitization	Guideline 406)
Serious eye	Eyes - Rabbit Result: No eye irritation
damage/eye irritation	
Skin	Skin - Rabbit Result: No skin irritation
corrosion/irritation	
Specific target organ	The substance or mixture is not classified as specific target organ toxicant, repeated
toxicity - repeated	exposure.
Exposure Specific target organ	Causes damage to organs
toxicity - single	Causes uamaye to organs.
exposure	
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	Skin Sensitization:, (Mouse) - non-sensitizing.
Sensitization	
damage/ove irritation	(Rabbit, 24 ft): slight.
	(Pabhit 24 h); modorato
corrosion/irritation	(Nabbit, 24 II). Inderate.
Specific target organ	No data available
toxicity - repeated	
exposure	
Specific target organ	No data available.
toxicity - single	
exposure	
Methyl Isobutyl Ketone(108-10-1)
Carcinogenicity Data	Methyl Isobutyl Ketone: Possibly carcinogenic to humans. (IARC-2B)
LC50 (Rat, 4)	8.2 - 16.4 mg/l
	> 1 C00 mg/l/g
LD50 (Rabbit) Dermal	>1 000 11g/kg
	2 000 - 4 000 IIIY/KY Mutagonicity tosts in animals have been negative as inconclusive. See "Other Studies Delevent
Mutagenicity Data	to Material"
Other Studies Revelant	According to the International Agency for Research on Cancer (IARC) methyl isobutyl ketone
Material	is possibly carcinogenic to humans. (IARC-2B) MIBK was not teratogenic, embryotoxic or
	fetotoxic following exposures that did not produce maternal toxicity. Rats and mice were
	exposed to 300, 1000 or 3000 ppm MIBK on days 6-15 of pregnancy. Exposures to 3000 ppm
	produced maternal and fetal toxicity, but no teratogenicity. There was no maternal toxicity,
	embryotoxicity or teratogenicity at 300 or 1000 ppm. Findings of fetotoxicity at 300 ppm were
	complicated by abnormal litter sizes and were determined not to be treatment related. (4)
	MIBK produced negative results in the micronucleus cryptogenic assay in mice in vivo. Most
Banroductiva Data	No adverse repreductive effects are anticipated
Reproductive Data	Nona known
Sensitization Data	None known.
Synergistic Materials	In studies with mice. MIBK prolonged the loss of righting reflex induced by ethanol. In animal
	studies, MIBK has been shown to potentiate the hepatotoxicity of haloalkanes, such as
	chloroform, carbon tetrachloride and 1,2-dichlorobenzene. Combined exposure to methyl
	ethyl ketone and MIBK caused increased behavioral responses in baboons.
Teratogenicity Data	No adverse teratogenic effects are anticipated. See "Other Studies Relevant to Material".
n-Butyl Acetate(123-86-	4)
Aspiration hazard	No data available.
Carcinogenicity	No data available.
Inhalation	No data available.
LD-50 Dermal -	> 16ml/kg
(Rabbil)	14 130 mg/kg
LD-50 Oldi - (Rdl)	14,150 Mg/Kg 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.
sensitization	
Serious eye	(Rabbit, 24 h): none
damage/eye irritation	
Skin	(Rabbit, 24 h): none
corrosion/irritation	
Specific target organ	No data available.
toxicity - repeated	
exposure	Nexastia offect
specific target organ	
exposure	
n-Methyl-2-nyrrolidone(872-50-4)
Aspiration Hazard	Not Applicable.

Assessment other	Assessment of STOT single: Causes temporary irritation of the respiratory tract. Irritation /
Assessment other	correction Accossment of irritating offacts: Eva contact causes irritation. Skip contact causes
acute effects	Controlling Assessment of initiating effects. Lye contact causes initiation. Skin Contact causes
	rability address temporary initiation of the respiratory tract. Eo-classification Skin Species.
	has be added this evolution with the training to align (120). For Charles, which has whether a subtraining the strike the strike the second strike the secon
	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 evoluted 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 accorsable provides no indication of a carcinogonic effect
Constin toxicity	Assessment of mutagenicity. The substance was not mutagenic in betaging the mutagenic
Genetic toxicity	Assessment of mutagenicity. The substance was not mutagenic in bacteria, no mutagenic
	enect was found in various tests with mammalian cell culture and mammals.
LC50 Inhalation - Rat	> 5.1 mg/I (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)
	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
Reproductive toxicity	damage to the testes after repeated high exposures that cause other toxic effects
	damage to the testes after repeated might exposite 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.
Tetragenicity	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
	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 neoplasms, increase incidence of
	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity
	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity classification not possible from current data.
Germ cell mutagenicity	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity classification not possible from current data.
Germ cell mutagenicity	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity classification not possible from current data. Genotoxicity in vitro, Test Type: Chromosome aberration test in vitro Test species: Chinese hamster ovary (CHO) Metabolic activation: with and without metabolic activation Method:
Germ cell mutagenicity	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity classification not possible from current data. 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: penative GLP: no : Test Type: Mammalian cell gene
Germ cell mutagenicity	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity classification not possible from current data. 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 accay Tact capaciac: mouse lymphoma colls Metabolic activation; with and without
Germ cell mutagenicity	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity classification not possible from current data. 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 0 FCP Test Cuideline 476 Deputy Provide La Participa Classification (Classification) (Classificat
Germ cell mutagenicity	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity classification not possible from current data. 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
Germ cell mutagenicity	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity classification not possible from current data. 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 Test species: mouse (male)
Germ cell mutagenicity	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity classification not possible from current data. 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 Test species: mouse (male) Application Route: Oral Method: OECD Test Guideline 474 Result: negative GLP: yes Test
Germ cell mutagenicity	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity classification not possible from current data. 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 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:
Germ cell mutagenicity	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity classification not possible from current data. 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 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
Germ cell mutagenicity	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity classification not possible from current data. 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 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
Germ cell mutagenicity	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity classification not possible from current data. 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 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 10 mg/l Assessment: The component/mixture is moderately toxic after short term inhalation.
Germ cell mutagenicity LC50 (Mouse, Male) LD50 (rabbit)	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity classification not possible from current data. 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 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 10 mg/l Assessment: The component/mixture is moderately toxic after short term inhalation. 15,433 mg/kg
Germ cell mutagenicity LC50 (Mouse, Male) LD50 (rabbit) Repeated dose toxicity	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity classification not possible from current data. 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 Test species: mouse (male) Application Route: Oral Method: OECD Test Guideline 474 Result: negative GLP: yes 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 10 mg/I Assessment: The component/mixture is moderately toxic after short term inhalation. 15,433 mg/kg Species: rat, male and female NOAEL: 75 mg/kg Application Route: Oral Exposure time: 28 d
Germ cell mutagenicity LC50 (Mouse, Male) LD50 (rabbit) Repeated dose toxicity	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity classification not possible from current data. 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 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 10 mg/l Assessment: The component/mixture is moderately toxic after short term inhalation. 15,433 mg/kg 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
Germ cell mutagenicity LC50 (Mouse, Male) LD50 (rabbit) Repeated dose toxicity	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity classification not possible from current data. 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 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 10 mg/l Assessment: The component/mixture is moderately toxic after short term inhalation. 15,433 mg/kg 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
Germ cell mutagenicity LC50 (Mouse, Male) LD50 (rabbit) Repeated dose toxicity	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity classification not possible from current data. 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 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 10 mg/l Assessment: The component/mixture is moderately toxic after short term inhalation. 15,433 mg/kg 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
Germ cell mutagenicity LC50 (Mouse, Male) LD50 (rabbit) Repeated dose toxicity Reproductive toxicity	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity classification not possible from current data. 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 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 10 mg/l Assessment: The component/mixture is moderately toxic after short term inhalation. 15,433 mg/kg 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 npm Duration of Single Treatment:
Germ cell mutagenicity LC50 (Mouse, Male) LD50 (rabbit) Repeated dose toxicity Reproductive toxicity	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity classification not possible from current data. 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 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 10 mg/l Assessment: The component/mixture is moderately toxic after short term inhalation. 15,433 mg/kg 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 npm General Toxicity E1: NOAEC: 100 npm
Germ cell mutagenicity LC50 (Mouse, Male) LD50 (rabbit) Repeated dose toxicity Reproductive toxicity	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity classification not possible from current data. 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 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 10 mg/l Assessment: The component/mixture is moderately toxic after short term inhalation. 15,433 mg/kg 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 Resured offering weight gain. Method: OECD Test
Germ cell mutagenicity LC50 (Mouse, Male) LD50 (rabbit) Repeated dose toxicity Reproductive toxicity	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity classification not possible from current data. 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 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 10 mg/l Assessment: The component/mixture is moderately toxic after short term inhalation. 15,433 mg/kg 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 Cl P: ves Effects on fetal development :
Germ cell mutagenicity LC50 (Mouse, Male) LD50 (rabbit) Repeated dose toxicity Reproductive toxicity	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity classification not possible from current data. 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 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 10 mg/l Assessment: The component/mixture is moderately toxic after short term inhalation. 15,433 mg/kg 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 Amelication Route: Inhalation Dose: 0, 100, 500 and 1000 ppm Duration of Single Treatment: Species: rat Amelication Route: Inhalation Dose: 0, 100, 500 and 1000 ppm Duration for fetal development : Species: rat Amelication Route: I
Germ cell mutagenicity LC50 (Mouse, Male) LD50 (rabbit) Repeated dose toxicity Reproductive toxicity	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity classification not possible from current data. 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 Test species: mouse (male) Application Route: Oral Method: OECD Test Guideline 476 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 10 mg/l Assessment: The component/mixture is moderately toxic after short term inhalation. 15,433 mg/kg 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 Concert Toxicity - F10 Application of Single Repaired F1 Method: Repaired Method: Concert Toxicity F1
Germ cell mutagenicity LC50 (Mouse, Male) LD50 (rabbit) Repeated dose toxicity Reproductive toxicity	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity classification not possible from current data. 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 Test species: mouse (male) Application Route: Oral Method: OECD Test Guideline 474 Result: negative GLP: yes 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 10 mg/l Assessment: The component/mixture is moderately toxic after short term inhalation. 15,433 mg/kg 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:
Germ cell mutagenicity LC50 (Mouse, Male) LD50 (rabbit) Repeated dose toxicity Reproductive toxicity	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity classification not possible from current data. 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 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 10 mg/l Assessment: The component/mixture is moderately toxic after short term inhalation. 15,433 mg/kg 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 b
Germ cell mutagenicity LC50 (Mouse, Male) LD50 (rabbit) Repeated dose toxicity Reproductive toxicity	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity classification not possible from current data. 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 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 10 mg/l Assessment: The component/mixture is moderately toxic after short term inhalation. 15,433 mg/kg 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 NAEC: 500 ppm Teratogenicity: NOAEC: 2,000 ppm Developmental Toxicity: NOAEC: 500 ppm Symptoms: Reduced body weight
Germ cell mutagenicity LC50 (Mouse, Male) LD50 (rabbit) Repeated dose toxicity Reproductive toxicity	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity classification not possible from current data. 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 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 10 mg/l Assessment: The component/mixture is moderately toxic after short term inhalation. 15,433 mg/kg 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
Germ cell mutagenicity LC50 (Mouse, Male) LD50 (rabbit) Repeated dose toxicity Reproductive toxicity	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity classification not possible from current data. 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 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 10 mg/l Assessment : The component/mixture is moderately toxic after short term inhalation. 15,433 mg/kg 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 Maternal: NOAEC: 500 ppm Teratoge
Germ cell mutagenicity LC50 (Mouse, Male) LD50 (rabbit) Repeated dose toxicity Reproductive toxicity Reproductive toxicity	Symptoms: increased incidences of alveolar/bronchiolar neoplasms, increase incidence of hepatocellular carcinomas GLP: yes Carcinogenicity - Assessment : Carcinogenicity classification not possible from current data. 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 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 10 mg/l Assessment: The component/mixture is moderately toxic after short term inhalation. 15,433 mg/kg 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

Serious eye	Species: rabbit Result: Mild eye irritation Remarks: No data available
Skin	Species: rabbit Result: Mild skin irritation
corrosion/irritation	
STOT - repeated	Target Organs: Auditory system Assessment: May cause damage to organs through prolonged
exposure	or repeated exposure, The substance or mixture is classified as specific target organ toxicant, repeated exposure, category 2
STOT - single exposure	No data available.
Phosphoric Acid(7664-38	3-2)
Additional Information	RTECS: TB6300000 burning sensation, Cough, wheezing, laryngitis, Shortness of breath,
	Headache, Nausea, Vomiting, May cause cyanosis. Stomach - Irregularities - Based on Human
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
	component of this product present at levels greater than or equal to 0.1% is identified as a
	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	No data available.
I D50 Dermal - Rabbit	2.740 mg/kg Remarks: Behavioral: Somnolence (general depressed activity). Behavioral:
	Excitement. No data available.
LD50 Oral - Rat -	> 5,000 mg/kg, (OECD Test Guideline 423),
Reproductive toxicity	No data available.
Respiratory or skin	No data available.
sensitization	
Serious eye	Eyes - Rabbit Result: Corrosive
Skin	Skin - Rabbit Result: Causes burns 24 h
corrosion/irritation	
Specific target organ	No data available.
toxicity - repeated	
Specific target organ	No data available.
toxicity - single	
exposure	
Additional Information	RTECS: DC3325000 prolonged or repeated exposure can cause: parcosis. Bronchitis
	Symptoms and signs include headache, dizziness, fatigue, muscular weakness, drowsiness
	and in extreme cases, loss of consciousness., To the best of our knowledge, the chemical,
	physical, and toxicological properties have not been thoroughly investigated. Central nervous
Carcinogenicity	IARC: No component of this product present at levels greater than or equal to 0.1% is
curentogeneity	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	in vitro assay S. typhimurium Result: negative Mutagenicity (micronucleus test) Rat - male
Inhalation:	No data available.
LD50 Oral - Rat -	6,000 mg/kg, Rat - male.
Acute toxicity	
Reproductive toxicity	No data available.
sensitization	
Serious eye	No data available.
damage/eye irritation	
SKIN corrosion/irritation	NO DATA AVAIIADIE
Specific target organ	No data available.
toxicity - repeated	
exposure	

Specific target organ	No data available.	
exposure		
Xylene(1330-20-7)		
Acute dermal toxicity	Acute toxicity estimate : 1 100 mg/kg Method: Expert judgment	
Acute inhalation	Acute toxicity estimate 4631 npm 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	Test Type: Chromosome aberration test in virto. Test Species: Chinese hamster ovary (CHO) Metabolic Activation: With and without metabolic activation. Method Mutagenicity (in vitro mammalian cytogenetic test) Result: Negative. Test Type: Sistrer chromatic exchange assay in mammalian cells.	
Germ cell mutagenicity Assessment	Animal testing did not show any mutagenic effects.	
LC50 (rat, male)	6700 ppm Exposure time: 4 h Method: Directive 67/548/EEC, Annex V, B.2. GLP: No data	
Inhalation	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	Species: rabbit Result: Mild eye irritation	
damage/eye irritation	· · · · · ·	
Skin	Species: rabbit Exposure time: 24 h Result: Irritating to skin Remarks: Skin irritation,	
corrosion/irritation	Category 2	
STOT - repeated	Target Organs: Liver, Kidney, Central nervous system Assessment: May cause damage to	
exposure	organs through prolonged or repeated exposure.	
STOT - single exposure	No data available.	

12. ECOLOGICAL INFORMATION

Acetone(67-64-1)	
Bioacculative potential	Parition coefficient: n-octanol/water: log Pow: -0.24
EC50 (Daphnia magna (Water flea))	7,630 mg/l (Exposure time 48 h); Test substance: Acetone
LC50 (Oncorhynchus mykiss (rainbow trout))	6,100 mg/l (Exposure time: 48 h)
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	Remarks: No data available
Aliphatic Solvent(64742	-47-8)
Bioaccumulative potential	No data available.

EC50 (Daphnia Magna)	1.4 mg/l - 48 h - Daphnia magna (Water flea) (OECD Test Guideline 202)	
	1.7 mg/r - 70 m, - Dapinina magna (water nea), (OLCD Test Guideline 202)	
i oxicity to daphnia and		
other aquatic		
invertebrates		
LC50 (Rainbow trout)	2.9 mg/l - 96 h. Oncorhynchus mykiss (rainhow trout)	
Toxicity to fish		
MODILITY IN SOIL	No data available.	
Other adverse effects	An environmental hazard cannot be excluded in the event of unprofessional handling or	
	disposal. Toxic to aquatic life. No data available.	
Persistence and	No data available	
de ave de bility		
degradability		
Results of PBT and	PBT/vPvB assessment not available as chemical safety assessment not required/not	
vPvB assessment	conducted.	
$C_{\rm umene}(98-82-8)$		
Bioaccumulative	No data avallable.	
potential		
EC50 - Daphnia (water	2.14 mg/l - 48 h (OECD Test Guideline 202), Daphnia (water flea)	
flea) - Toxicity to		
daphnia and other		
aquatic invertebrates		
EC50 -	2.60 mg/l - 72 h, Pseudokirchneriella subcapitata (green algae)	
Pseudokirchneriella		
subcapitata (groop		
algae) - Toxicity to		
algae		
LC50 - Oncorhynchus	4.8 mg/l - 96 h, Oncorhynchus mykiss (rainbow trout)	
mykiss (rainbow trout)		
i oxicity to fish		
Mobility in soil	No data available.	
Other adverse effects	An environmental hazard cannot be excluded in the event of upprofessional handling or	
	disposal. Toxic to aquiatic life with long lasting offects	
	disposal. Toxic to aquatic me with ong lasting effects.	
Persistence and	Biodegradability Result: - According to the results of tests of biodegradability this product is	
degradability	not readily biodegradable.	
Results of PBT and	PBT/vPvB assessment not available as chemical safety assessment not required/not	
VPVB assossment	conductod	
Dioctyl Phthalate(117-81	1-7)	
Dioctyl Phthalate(117-81 Bioaccumulative	1-7) Oncorhynchus mykiss (rainbow trout) BFC: 113. Exposure time: 100 d.	
Dioctyl Phthalate(117-81 Bioaccumulative Potential -	1-7) Oncorhynchus mykiss (rainbow trout) BFC: 113. Exposure time: 100 d.	
Dioctyl Phthalate(117-81 Bioaccumulative Potential - Bioaccumulation	0ncorhynchus mykiss (rainbow trout) BFC: 113. Exposure time: 100 d.	
Dioctyl Phthalate(117-81 Bioaccumulative Potential - Bioaccumulation	Oncorhynchus mykiss (rainbow trout) BFC: 113. Exposure time: 100 d.	
Dioctyl Phthalate(117-81 Bioaccumulative Potential - Bioaccumulation Chronic Aquatic	1-7) Oncorhynchus mykiss (rainbow trout) BFC: 113. Exposure time: 100 d. Very toxic to aquatic life with long lasting effects	
Dioctyl Phthalate(117-81 Bioaccumulative Potential - Bioaccumulation Chronic Aquatic Toxicity	Oncorhynchus mykiss (rainbow trout) BFC: 113. Exposure time: 100 d. Very toxic to aquatic life with long lasting effects	
Dioctyl Phthalate(117-81 Bioaccumulative Potential - Bioaccumulation Chronic Aquatic Toxicity EC50 Daphnia Magna -	1-7) Oncorhynchus mykiss (rainbow trout) BFC: 113. Exposure time: 100 d. Very toxic to aquatic life with long lasting effects 0.16 mg/l. 48 h. Immobilization	
Dioctyl Phthalate(117-81 Bioaccumulative Potential - Bioaccumulation Chronic Aquatic Toxicity EC50 Daphnia Magna - Toxicity to daphnia and	1-7) Oncorhynchus mykiss (rainbow trout) BFC: 113. Exposure time: 100 d. Very toxic to aquatic life with long lasting effects 0.16 mg/l. 48 h. Immobilization	
Dioctyl Phthalate(117-81 Bioaccumulative Potential - Bioaccumulation Chronic Aquatic Toxicity EC50 Daphnia Magna - Toxicity to daphnia and other aquatic	1-7) Oncorhynchus mykiss (rainbow trout) BFC: 113. Exposure time: 100 d. Very toxic to aquatic life with long lasting effects 0.16 mg/l. 48 h. Immobilization	
Dioctyl Phthalate(117-81 Bioaccumulative Potential - Bioaccumulation Chronic Aquatic Toxicity EC50 Daphnia Magna - Toxicity to daphnia and other aquatic	1-7) Oncorhynchus mykiss (rainbow trout) BFC: 113. Exposure time: 100 d. Very toxic to aquatic life with long lasting effects 0.16 mg/l. 48 h. Immobilization	
Dioctyl Phthalate(117-81 Bioaccumulative Potential - Bioaccumulation Chronic Aquatic Toxicity EC50 Daphnia Magna - Toxicity to daphnia and other aquatic invertebrates	1-7) Oncorhynchus mykiss (rainbow trout) BFC: 113. Exposure time: 100 d. Very toxic to aquatic life with long lasting effects 0.16 mg/l. 48 h. Immobilization	
Dioctyl Phthalate(117-81 Bioaccumulative Potential - Bioaccumulation Chronic Aquatic Toxicity EC50 Daphnia Magna - Toxicity to daphnia and other aquatic invertebrates EC50	1-7) Oncorhynchus mykiss (rainbow trout) BFC: 113. Exposure time: 100 d. Very toxic to aquatic life with long lasting effects 0.16 mg/l. 48 h. Immobilization 0.003 mg/l. Biomass 72 h. ISO 8692.	
Dioctyl Phthalate(117-81 Bioaccumulative Potential - Bioaccumulation Chronic Aquatic Toxicity EC50 Daphnia Magna - Toxicity to daphnia and other aquatic invertebrates EC50 Pseudokirchneriella	1-7) Oncorhynchus mykiss (rainbow trout) BFC: 113. Exposure time: 100 d. Very toxic to aquatic life with long lasting effects 0.16 mg/l. 48 h. Immobilization 0.003 mg/l. Biomass 72 h. ISO 8692.	
Dioctyl Phthalate(117-81 Bioaccumulative Potential - Bioaccumulation Chronic Aquatic Toxicity EC50 Daphnia Magna - Toxicity to daphnia and other aquatic invertebrates EC50 Pseudokirchneriella Subcapitata - Toxicity	1-7) Oncorhynchus mykiss (rainbow trout) BFC: 113. Exposure time: 100 d. Very toxic to aquatic life with long lasting effects 0.16 mg/l. 48 h. Immobilization 0.003 mg/l. Biomass 72 h. ISO 8692.	
Dioctyl Phthalate(117-81 Bioaccumulative Potential - Bioaccumulation Chronic Aquatic Toxicity EC50 Daphnia Magna - Toxicity to daphnia and other aquatic invertebrates EC50 Pseudokirchneriella Subcapitata - Toxicity to Algae	1-7) Oncorhynchus mykiss (rainbow trout) BFC: 113. Exposure time: 100 d. Very toxic to aquatic life with long lasting effects 0.16 mg/l. 48 h. Immobilization 0.003 mg/l. Biomass 72 h. ISO 8692.	
Dioctyl Phthalate(117-81 Bioaccumulative Potential - Bioaccumulation Chronic Aquatic Toxicity EC50 Daphnia Magna - Toxicity to daphnia and other aquatic invertebrates EC50 Pseudokirchneriella Subcapitata - Toxicity to Algae	1-7) Oncorhynchus mykiss (rainbow trout) BFC: 113. Exposure time: 100 d. Very toxic to aquatic life with long lasting effects 0.16 mg/l. 48 h. Immobilization 0.003 mg/l. Biomass 72 h. ISO 8692.	
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Ethyl Alcohol(64-17-5)	
Bioaccumulative	No data available.
potential	
Mobility in soil	No data available.
Other adverse effects	No data available.
Persistence and	No data available.
degradability	
Results of PBT and	PBT/vPvB assessment not available as chemical safety assessment not required/not
VPvB assessment	conducted
loxicity	No data available.
Ethylene glycol mono bu	tyl etner(111-/6-2)
potential	Partition coefficient: n-octanol/water: log Pow: 0.83
EC50 (Algee)	911 mg/l End point: Biomass Exposure time: 72 h Test Type: static test Analytical
EC50 (Daphnia)	1,800 mg/l(48 h; Daphnia magna (Water flea)): Exposure time: 48 h Test Type: static test Method: OECD Test Guideline 202 GLP: no
LC50 (fish)	1,474 mg/l Pimephales promelas (Fathead minnow))Exposure time: 96 h Test Type: static test, Method: OECD Test Guideline 203 GLP: no
Mobility in soil	No data available
Other adverse effects	No data available
Persistence and degradability	aerobic Inoculum: Activated sludge, domestic, adaption not specified, Result: Readily biodegradable. Biodegradation: 90.4 % Exposure time: 28 d Method: OECD Test Guideline 301B GLP: no
Product	Regulation: 40CFR Protection of Environment, Part 82 Protection of Stratospheric Ozone - CAA Section 602 Class 1 Substances:
Isobutyl Alcohol(78-83-1	
Chronic	No data available.
Degradability /	Evaluation: Not readily biodegradable (by OECD criteria).
Persistence; Biological	, 5 (, ,
/ A biological	
Degradation	
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.
Isopropyl Alcohol(67-63-	-0)
Bioaccumulative	Bioaccumulation : Bioconcentration factor (BCF): 3.16 this material is not expected to
potential	bioaccumulate.
Ecotoxicology Assessment	Acute aquatic toxicity: Based on acute aquatic toxicity values, not classified. Chronic aquatic toxicity: Not classified, based on readily biodegradability and low acute toxicity.
Mobility in soil	Distribution among environmental compartments: Stability in water initially partitioning
	mainly to water and air. Stability in soil Volatilization from water or soil surfaces is expected to be limited. Additional advice Environmental fate and pathways : No additional information
Other adverse effects	No additional information available.
Additional ecological	
information	
Persistence and	Biodegradability : 86 - 94 % Rapidly degradable. (After two weeks in a ready biodegradability
	Not applicable
VPVB assessment	
Toxicity to algae	Acute toxicity to aquatic plants very low.
Toxicity to bacteria	Low toxicity to sewage microbes.
Toxicity to daphnia and	Acute toxicity to freshwater and marine invertebrates is very low.
other aquatic	, ,
Toxicity to danhnia and	Chronic toxicity expected to be low
other aquatic	
invertebrates (Chronic	
toxicity)	
Toxicity to fish	Acute toxicity to fish is very low.
Toxicity to fish	Chronic toxicity to fish is expected to be low.
(Chronic toxicity)	
Methyl Alcohol(67-56-1)	

Bioaccumulative	Bioaccumulation Cyprinus carpio (Carp) - 72 d at 20 °C - 5 mg/l Bioconcentration factor	
EC50 - Daphnia magna	> 10,000.00 mg/l - 48 h Toxicity to daphnia and other aquatic invertebrates, Daphnia magna	
-	(Water flea)	
EC50 - Scenedesmus	22,000.0 mg/l - 96 h, Scenedesmus capricornutum (fresh water algae)	
Toxicity to algae		
IC50 Activated sludge -	>1,000 mg/l, Exposure 3 h, Test type Static, Method OECD Test Guideline 209.	
Toxicity to bacteria	15,400,0 mg/l 06 h Lonomic macrochirus (Pluagill)	
macrochirus - Toxicity	13,400.0 mg/1 - 90 m, Leponnis macrochinus (bluegin)	
to Fish		
Mobility in soil	Will not adsorb on soil.	
Persistence and	Biodegradability aerobic - Exposure time 5 d Result: 72 % - rapidly biodegradable	
degradability	Biochemical Oxygen Demand (BOD) 600 - 1,120 mg/g Chemical Oxygen Demand (COD) 1,420 mg/g Theoretical oxygen demand 1,500 mg/g	
Methyl Amyl Ketone(110	-43-0)	
Aquatic invertebrates	No data available.	
Bioaccumulative	No data available.	
Chronic Toxicity (Fish)	No data available.	
ErC50 (Selenastrum	98.2 mg/l, 72 h	
LC50 (Fathead	131 mg/L (96 h)	
Minnow) Acute toxicity		
Mobility in soil	No data available.	
Persistence and	69 % (28 d, Ready Biodegradability - CO2 in Sealed Vessels (Headspace Test)). Biological	
degradability	2,420 mg/g, BOD/COD ratio No data available.	
Results of PBT and	No data available.	
vPvB assessment		
Deactivating	None required	
Chemicals: None		
required.		
Disposal of Packaging	Empty containers retain product residue (liquid and/or vapour) and can be dangerous. Empty	
	reconditioner. Do not expose such containers to heat, flame, sparks, static electricity, or other	
	sources of ignition; they may explode and cause injury or death. Do not dispose of package	
	until thoroughly washed out.	
EC50 (Daphnia Magna)	>200 mg/l (48 h)	
Ecoloxicity Environmental Fate	Can be dangerous if allowed to enter drinking water intakes. Do not contaminate domestic or	
	irrigation water supplies, lakes, streams, ponds, or rivers. Methyl Isobutyl Ketone: This	
	product is biodegradable. This product does not bioaccumulate in aquatic or terrestrial food	
LC50 (Fathead	>179 ma/l (96 h)	
Minnow)		
Safe Handling of Residues	See "Waste Disposal Methods"	
Waste Disposal	. Reevaluation of the product may be required by the user at the time of disposal since the	
Methous	Dispose of waste material at an approved (hazardous) waste treatment/disposal facility in	
	accordance with applicable local, provincial and federal regulations. Do not dispose of waste	
	with normal garbage, or to sewer systems.	
Bioaccumulative	4) No data available	
potential		
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)	44 mg/l , (48 h)	
Aquatic invertebrates	Known or prodicted distribution to environmental compartmenter No data available	
Other adverse effects	No data available.	
Persistence and	22 W (22 d) Biological Ovygon Domandy BOD E: 720 mg/g Chamical Ovygon Domandy 1 010	
i di biblici de alla	os % (26 d), biológical Oxygen Demand:BOD-5: 750 mg/g, Chemical Oxygen Demand:1,010 j	

Results of PBT and	No data available.	
vPvB assessment		
n-Methyl-2-pyrrolidone(8	372-50-4)	
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 potential	Assessment bioaccumulation potential Because of the n-octanol/water distribution coefficient (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 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 the nominal concentration.	
Microorganisms/Effect on activated sludge	Toxicity to microorganisms DIN EN ISO 8192 aquatic activated sludge, industrial/EC50 (0.5 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 expected.	
Persistence and degradability	Assessment biodegradation and elimination (H2O) Readily biodegradable (according to OECD 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 contact with water the substance will hydrolyze slowly.	
Phenylethane(100-41-4)		
Bioaccumulative potential	Partition coefficient: noctanol/water : log Pow: 2.92	
EC50 (Daphnia magna (Water flea))	1.8 mg/l Exposure time: 48 h Test Type: static test	
EC50 (Pseudokirchneriella subcapitata)	5.4 mg/l Exposure time: 72 h Test Type: static test Analytical monitoring: yes Method: Static GLP: yes	
LC50 (Oncorhynchus mykiss (rainbow trout))	4.2 mg/l Exposure time: 96 h Test Type: semi-static test	
Mobility in soil	No data available.	
Other adverse effects	Results of PBT and vPvB assessment : This substance is not considered to be persistent, bioaccumulating nor toxic (PBT). This substance is not considered to be very persistent nor very bioaccumulating (vPvB).	
Persistence and degradability	Biodegradability : Inoculum: activated sludge Concentration: 22 mg/l Result: Readily biodegradable. Biodegradation: 70 % Exposure time: 28 d GLP: yes	
Toxicity to daphnia and other aquatic invertebrates (Chronic toxicity)	(Daphnia): 3.6 mg/l Toxicity to bacteria : GLP: Remarks: No data available Ecotoxicology Assessment Chronic aquatic toxicity : Harmful to aquatic life with long lasting effects.	
Phosphoric Acid(7664-38	3-2)	
Bioaccumulative potential	No data available.	
Mobility in soil	No data available.	
Other adverse effects	May be harmful to aquatic organisms due to the shift of the pH.	
Persistence and degradability	No data available.	
Results of PBT and	PBT/vPvB assessment not available as chemical safety assessment not required/not	
	No data available	
Pseudocumene(95-63-6)		
Bioaccumulative	No data available.	
potential		
EC50 - Daphnia magna (Water flea) - Toxicity	3.6 mg/l - 48 h (OECD Test Guideline 202), Daphnia magna (Water flea)	
to daphnia and other aquatic invertebrates		
LC50 - Pimephales	7.72 mg/l - 96.0 h, Pimephales promelas (fathead minnow)	
promelas (fathead		
minnow) - Toxicity to		
TISN Mobility in soil	No data available	

Other adverse effects	An environmental hazard cannot be excluded in the event of unprofessional handling or disposal. Toxic to aquatic life with long lasting effects.	
Persistence and degradability	No data available.	
Results of PBT and vPvB assessment	PBT/vPvB assessment not available as chemical safety assessment not required/not conducted	
Xylene(1330-20-7)		
Bioaccumulative potential	Partition coefficient: noctanol/water : log Pow: 2.77 - 3.15	
EC50 (Pseudokirchneriella subcapitata)	4.36 mg/l End point: Growth rate Exposure time: 73 h Test Type: static test Analytical monitoring: yes	
IC50 (Daphnia magna (Water flea))	1 mg/l Exposure time: 24 h Test Type: static test substance: Information given is based on data obtained from similar substances. Method: OECD Test Guideline 202 GLP	
LC50 (Oncorhynchus mykiss (rainbow trout))	2.6 mg/l Exposure time: 96 h Test substance: Information given is based on data obtained from similar substances. Method: OECD Test Guideline 203 GLP: No data available	
Mobility in soil	No data available.	
Persistence and degradability	Biodegradability : Inoculum: activated sludge Result: Readily biodegradable. Biodegradation: 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 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, Moderate skin irritant, Moderate eye irritant, Carcinogen.

EPCRA - Emergency CERCLA REPORTABLE QUANTITY

This product contains:	Chemical CAS#
Ethyl Alcohol	64-17-5
n-Butyl Acetate	123-86-4
Ethylene glycol mono butyl ether	111-76-2
Xylene	1330-20-7
Isobutyl Alcohol	78-83-1
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 AMENDMENTS AND REAUTHORIZATION ACT)

SARA 311/312 Hazards : Fire Hazard, Acute Health Hazard, Chronic Health Hazard SARA 313 :

This product contains:	Chemical CAS#
Ethyl Alcohol	64-17-5
n-Butyl Acetate	123-86-4
Aliphatic Solvent	64742-47-8
Acetone	67-64-1
Isopropyl Alcohol	67-63-0
Pseudocumene	95-63-6
Methyl Isobutyl Ketone	108-10-1
Methyl Amyl Ketone	110-43-0
Methyl Alcohol	67-56-1
Dioctyl Phthalate	117-81-7
n-Methyl-2-pyrrolidone	872-50-4
Phenylethane	100-41-4

CLEAN AIR ACT :

This product contains:	Chemical CAS#
Methyl Isobutyl Ketone	108-10-1
Methyl Alcohol	67-56-1
Cumene	98-82-8
Phenylethane	100-41-4

INTERNATIONAL REGULATIONS

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

Flam. Liq. Cat 2;	H226
Acute Tox. Cat 4;	H302
Aspir. Haz. Cat. 1;	H304
Acute Tox. Cat. 4;	H312
Skin Irrit. Cat. 2;	H315
Eye Irrit. Cat. 2A;	H319
Acute Toc. Cat. 4;	H332
STOT SE Resp. Cat. 3;	H335
STOT SE, Inhalation, Cat. 3;	H336
Carc. 2;	H351
STOT RE Cat.2;	H373
Aquatic Acute 2;	H401
Aquatic Tox. 3; Cat. 3;	H402

NATIONAL REGULATIONS

This product contains:	Chemical CAS#
#Methyl Isobutyl Ketone	108-10-1
#Phenylethane	100-41-4

Indicates a chemical listed by IARC as a possible carcinogen.

STATE REGULATIONS CALIFORNIA PROPOSITION 65

This product contains:	Chemical CAS#	
*Aliphatic Solvent	64742-47-8	
#Methyl Isobutyl Ketone	108-10-1	
+Methyl Alcohol	67-56-1	
*Dioctyl Phthalate	117-81-7	
+n-Methyl-2-pyrrolidone	872-50-4	
*Cumene	98-82-8	
*Phenylethane	100-41-4	

*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.

This product contains	Chemical CAS#
n-Butyl Acetate	123-86-4
Aliphatic Solvent	64742-47-8
Acetone	67-64-1
Pseudocumene	95-63-6
Methyl Amyl Ketone	110-43-0
Methyl Alcohol	67-56-1
Dioctyl Phthalate	117-81-7
n-Methyl-2-pyrrolidone	872-50-4
Ethylene glycol mono butyl ether	111-76-2
Xylene	1330-20-7
Cumene	98-82-8
Isobutyl Alcohol	78-83-1
Phenylethane	100-41-4
Phosphoric Acid	7664-38-2

Massachusetts Right to Know

Pennsylvania Right to Know

This product contains	Chemical CAS#
Ethyl Alcohol	64-17-5
n-Butyl Acetate	123-86-4
Aliphatic Solvent	64742-47-8
Acetone	67-64-1
Pseudocumene	95-63-6
Methyl Amyl Ketone	110-43-0

Methyl Alcohol	67-56-1
Dioctyl Phthalate	117-81-7
n-Methyl-2-pyrrolidone	872-50-4
Ethylene glycol mono butyl ether	111-76-2
Xylene	1330-20-7
Cumene	98-82-8
Isobutyl Alcohol	78-83-1
Phenylethane	100-41-4
Phosphoric Acid	7664-38-2
Water	7732-18-5

New Jersey Right to Know

This product contains	Chemical CAS#
Ethyl Alcohol	64-17-5
n-Butyl Acetate	123-86-4
Aliphatic Solvent	64742-47-8
Acetone	67-64-1
Pseudocumene	95-63-6
Methyl Amyl Ketone	110-43-0
Methyl Alcohol	67-56-1
Dioctyl Phthalate	117-81-7
n-Methyl-2-pyrrolidone	872-50-4
Ethylene glycol mono butyl ether	111-76-2
Xylene	1330-20-7
Cumene	98-82-8
Isobutyl Alcohol	78-83-1
Phenylethane	100-41-4
Phosphoric Acid	7664-38-2
Water	7732-18-5

16. OTHER INFORMATION

Other Product Information

% Volatile by Volume: 93.88% Solids by volume: 6.12% Exempt by Volume: 7.57

% Volatile by Weight: 91.73 % Solids by Weight: 8.27 % Exempt by Weight: 7.16

VOC CONTENT:

Excluding Exempt VOC: 765 Including Exempt VOC: 707

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



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