

Ardex WPM 240 Primer (Aus) Ardex (Ardex Australia)

Chemwatch: 5460-02 Version No: 2.1.1.1 Safety Data Sheet according to WHS and ADG requirements

Chemwatch Hazard Alert Code: 3

Issue Date: 23/03/2021 Print Date: 23/03/2021 S.GHS.AUS.EN

SECTION 1 Identification of the substance / mixture and of the company / undertaking

Product Identifier

Product name	Ardex WPM 240 Primer (Aus)	
Chemical Name	Not Applicable	
Synonyms	Not Available	
Proper shipping name	PAINT (including paint, lacquer, enamel, stain, shellac, varnish, polish, liquid filler and liquid lacquer base) or PAINT RELATED MATERIAL (including paint thinning or reducing compound)	
Chemical formula	Not Applicable	
Other means of identification	Not Available	

Relevant identified uses of the substance or mixture and uses advised against

Relevant identified uses	Bituminous solvent primer for the building industry.
--------------------------	--

Details of the supplier of the safety data sheet

Registered company name	Ardex (Ardex Australia)	
Address	20 Powers Road Seven Hills NSW 2147 Australia	
Telephone	1800 224 070	
Fax	1300 780 102	
Website	Not Available	
Email	Not Available	

Emergency telephone number

Association / Organisation	Ardex (Ardex Australia)	
Emergency telephone numbers	1800 224 070 (Mon-Fri, 9am-5pm)	
Other emergency telephone numbers	Not Available	

SECTION 2 Hazards identification

Classification of the substance or mixture

HAZARDOUS CHEMICAL. DANGEROUS GOODS. According to the WHS Regulations and the ADG Code.

ChemWatch Hazard Ratings

		Min	Max	
Flammability	3			
Toxicity	2			0 = Minimum
Body Contact	2			1 = Low
Reactivity	1			2 = Moderate
Chronic	2			3 = High 4 = Extreme

Poisons Schedule	S6
Classification ^[1]	Flammable Liquid Category 2, Aspiration Hazard Category 1, Skin Corrosion/Irritation Category 2, Eye Irritation Category 2A, Specific target organ toxicity - single exposure Category 3 (narcotic effects), Carcinogenicity Category 2, Reproductive Toxicity Category 2, Specific target organ toxicity - repeated exposure Category 2, Acute Aquatic Hazard Category 3, Chronic Aquatic Hazard Category 3
Legend:	1. Classified by Chernwatch; 2. Classification drawn from HCIS; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI



Signal word Danger

Hazard statement(s)

H225	Highly flammable liquid and vapour.	
H304	May be fatal if swallowed and enters airways.	
H315	Causes skin irritation.	
H319	Causes serious eye irritation.	
H336	May cause drowsiness or dizziness.	
H351	Suspected of causing cancer.	
H361d	Suspected of damaging the unborn child.	
H373	May cause damage to organs through prolonged or repeated exposure.	
H412	Harmful to aquatic life with long lasting effects.	

Precautionary statement(s) Prevention

P201	Obtain special instructions before use.	
P210	P210 Keep away from heat, hot surfaces, sparks, open flames and other ignition sources. No smoking.	
P260	260 Do not breathe mist/vapours/spray.	
P271 Use only outdoors or in a well-ventilated area.		

Precautionary statement(s) Response

P301+P310	IF SWALLOWED: Immediately call a POISON CENTER/doctor/	
P308+P313	IF exposed or concerned: Get medical advice/attention.	
P331	Do NOT induce vomiting.	
P370+P378 In case of fire: Use alcohol resistant foam or normal protein foam to extinguish.		

Precautionary statement(s) Storage

P403+P235	Store in a well-ventilated place. Keep cool.	
P405	Store locked up.	

Precautionary statement(s) Disposal

P501 Dispose of contents/container to authorised hazardous or special waste collection point in accordance with any local regulation.

SECTION 3 Composition / information on ingredients

Substances

See section below for composition of Mixtures

Mixtures

CAS No	%[weight]	Name
1330-20-7	10-40	xylene
108-88-3	10-20	toluene
64742-95-6.	3-10	naphtha petroleum, light aromatic solvent
123-86-4	3-10	n-butyl acetate
141-78-6	3-10	ethyl acetate
67-64-1	1-3	acetone
78-93-3	1-3	methyl ethyl ketone
100-42-5	1-3	styrene
100-41-4	1-3	ethylbenzene
108-10-1	1-3	methyl isobutyl ketone
108-67-8	1-2.5	1.3.5-trimethyl benzene
142-82-5	0.25-1	heptane
110-54-3	0.1-1	n-hexane

SECTION 4 First aid measures

Description of first aid measures

	 Immediately hold eyelids apart and flush the eye continuously with running water. Ensure complete irrigation of the eye by keeping eyelids apart and away from eye and moving the eyelids by occasionally lifting the upper and lower lids. Continue flushing until advised to stop by the Poisons Information Centre or a doctor, or for at least 15 minutes. Transport to hospital or doctor without delay. Removal of contact lenses after an eye injury should only be undertaken by skilled personnel.
Skin Contact	 If skin or hair contact occurs: Immediately flush body and clothes with large amounts of water, using safety shower if available. Quickly remove all contaminated clothing, including footwear. Wash skin and hair with running water. Continue flushing with water until advised to stop by the Poisons Information Centre. Transport to hospital, or doctor.
Inhalation	 If fumes or combustion products are inhaled remove from contaminated area. Lay patient down. Keep warm and rested. Prostheses such as false teeth, which may block airway, should be removed, where possible, prior to initiating first aid procedures. Apply artificial respiration if not breathing, preferably with a demand valve resuscitator, bag-valve mask device, or pocket mask as trained. Perform CPR if necessary. Transport to hospital, or doctor, without delay.
Ingestion	 Avoid giving milk or oils. Avoid giving alcohol. For advice, contact a Poisons Information Centre or a doctor at once. Urgent hospital treatment is likely to be needed. If swallowed do NOT induce vomiting. If vomiting occurs, lean patient forward or place on left side (head-down position, if possible) to maintain open airway and prevent aspiration. Observe the patient carefully. Never give liquid to a person showing signs of being sleepy or with reduced awareness; i.e. becoming unconscious. Give water to rinse out mouth, then provide liquid slowly and as much as casualty can comfortably drink. Transport to hospital or doctor without delay.

Indication of any immediate medical attention and special treatment needed

Any material aspirated during vomiting may produce lung injury. Therefore emesis should not be induced mechanically or pharmacologically. Mechanical means should be used if it is considered necessary to evacuate the stomach contents; these include gastric lavage after endotracheal intubation. If spontaneous vomiting has occurred after ingestion, the patient should be monitored for difficult breathing, as adverse effects of aspiration into the lungs may be delayed up to 48 hours. Treat symptomatically.

SECTION 5 Firefighting measures

Extinguishing media

- Alcohol stable foam.
- Dry chemical powder.
- BCF (where regulations permit).
- Carbon dioxide.

Special hazards arising from the substrate or mixture

Fire Incompatibility	Avoid contamination with oxidising agents i.e. nitrates, oxidising acids, chlorine bleaches, pool chlorine etc. as ignition may result
Advice for firefighters	
Fire Fighting	 Alert Fire Brigade and tell them location and nature of hazard. May be violently or explosively reactive. Wear breathing apparatus plus protective gloves in the event of a fire. Prevent, by any means available, spillage from entering drains or water course.
Fire/Explosion Hazard	 Liquid and vapour are highly flammable. Severe fire hazard when exposed to heat, flame and/or oxidisers. Vapour may travel a considerable distance to source of ignition. Heating may cause expansion or decomposition leading to violent rupture of containers. Combustion products include: carbon dioxide (CO2) other pyrolysis products typical of burning organic material. Contains low boiling substance: Closed containers may rupture due to pressure buildup under fire conditions.
HAZCHEM	•3YE

SECTION 6 Accidental release measures

Personal precautions, protective equipment and emergency procedures See section 8

Environmental precautions

See section 12

Methods and material for containment and cleaning up

Minor Spills	 Remove all ignition sources. Clean up all spills immediately. Avoid breathing vapours and contact with skin and eyes. Control personal contact with the substance, by using protective equipment.
Major Spills	 Clear area of personnel and move upwind. Alert Fire Brigade and tell them location and nature of hazard. May be violently or explosively reactive. Wear breathing apparatus plus protective gloves.

Personal Protective Equipment advice is contained in Section 8 of the SDS.

SECTION 7 Handling and storage

Precautions for safe handling	
Safe handling	 Containers, even those that have been emptied, may contain explosive vapours. Do NOT cut, drill, grind, weld or perform similar operations on or near containers. Contains low boiling substance: Storage in sealed containers may result in pressure buildup causing violent rupture of containers not rated appropriately. Check for bulging containers. Vent periodically Always release caps or seals slowly to ensure slow dissipation of vapours DO NOT allow clothing wet with material to stay in contact with skin Electrostatic discharge may be generated during pumping - this may result in fire. Ensure electrical continuity by bonding and grounding (earthing) all equipment. Restrict line velocity during pumping in order to avoid generation of electrostatic discharge (<=1 m/sec until fill pipe submerged to twice its diameter, then <= 7 m/sec). Avoid splash filling. Avoid splash filling. Avoid all personal contact, including inhalation. Wear protective clothing when risk of exposure occurs. Use in a well-ventilated area. Prevent concentration in hollows and sumps.
Other information	 Store in original containers in approved flame-proof area. No smoking, naked lights, heat or ignition sources. DO NOT store in pits, depressions, basements or areas where vapours may be trapped. Keep containers securely sealed.

Conditions for safe storage, including any incompatibilities

Suitable container	 Packing as supplied by manufacturer. Plastic containers may only be used if approved for flammable liquid. Check that containers are clearly labelled and free from leaks. For low viscosity materials (i) : Drums and jerry cans must be of the non-removable head type. (ii) : Where a can is to be used as an inner package, the can must have a screwed enclosure. For materials with a viscosity of at least 2680 cSt. (23 deg. C) For manufactured product having a viscosity of at least 250 cSt.
Storage incompatibility	 Avoid reaction with oxidising agents Avoid strong acids, acid chlorides, acid anhydrides and chloroformates.

SECTION 8 Exposure controls / personal protection

Control parameters

Occupational Exposure Limits (OEL)

INGREDIENT DATA

Source	Ingredient	Material name	TWA	STEL	Peak	Notes
Australia Exposure Standards	xylene	Xylene (o-, m-, p- isomers)	80 ppm / 350 mg/m3	655 mg/m3 / 150 ppm	Not Available	Not Available
Australia Exposure Standards	toluene	Toluene	50 ppm / 191 mg/m3	574 mg/m3 / 150 ppm	Not Available	Not Available
Australia Exposure Standards	n-butyl acetate	n-Butyl acetate	150 ppm / 713 mg/m3	950 mg/m3 / 200 ppm	Not Available	Not Available
Australia Exposure Standards	ethyl acetate	Ethyl acetate	200 ppm / 720 mg/m3	1440 mg/m3 / 400 ppm	Not Available	Not Available
Australia Exposure Standards	acetone	Acetone	500 ppm / 1185 mg/m3	2375 mg/m3 / 1000 ppm	Not Available	Not Available
Australia Exposure Standards	methyl ethyl ketone	Methyl ethyl ketone (MEK)	150 ppm / 445 mg/m3	890 mg/m3 / 300 ppm	Not Available	Not Available
Australia Exposure Standards	styrene	Styrene, monomer	50 ppm / 213 mg/m3	426 mg/m3 / 100 ppm	Not Available	Not Available
Australia Exposure Standards	ethylbenzene	Ethyl benzene	100 ppm / 434 mg/m3	543 mg/m3 / 125 ppm	Not Available	Not Available
Australia Exposure Standards	methyl isobutyl ketone	Methyl isobutyl ketone	50 ppm / 205 mg/m3	307 mg/m3 / 75 ppm	Not Available	Not Available
Australia Exposure Standards	heptane	Heptane (n-Heptane)	400 ppm / 1640 mg/m3	2050 mg/m3 / 500 ppm	Not Available	Not Available
Australia Exposure Standards	n-hexane	Hexane (n-Hexane)	20 ppm / 72 mg/m3	Not Available	Not Available	Not Available

Emergency Limits

Ingredient	TEEL-1	TEEL-2	TEEL-3
xylene	Not Available	Not Available	Not Available
toluene	Not Available	Not Available	Not Available
naphtha petroleum, light aromatic solvent	1,200 mg/m3	6,700 mg/m3	40,000 mg/m3

Chemwatch: **5460-02** Version No: **2.1.1.1**

Ardex WPM 240 Primer (Aus)

Ingredient	TEEL-1	TEEL-2		TEEL-3
n-butyl acetate	Not Available	Not Available		Not Available
ethyl acetate	1,200 ppm	1,700 ppm		10000** ppm
acetone	Not Available	Not Available		Not Available
methyl ethyl ketone	Not Available	Not Available		Not Available
styrene	Not Available	Not Available		Not Available
ethylbenzene	Not Available	Not Available		Not Available
methyl isobutyl ketone	75 ppm	500 ppm		3000* ppm
1,3,5-trimethyl benzene	Not Available	Not Available		480 ppm
heptane	500 ppm	830 ppm		5000* ppm
n-hexane	260 ppm	Not Available		Not Available
Ingredient	Original IDLH		Revised IDLH	
xylene	900 ppm		Not Available	
toluene naphtha petroleum, light aromatic solvent	500 ppm Not Available		Not Available	
n-butyl acetate	1,700 ppm		Not Available	
ethyl acetate	2,000 ppm		Not Available	
acetone	2,500 ppm		Not Available	
methyl ethyl ketone	3,000 ppm		Not Available	
styrene	700 ppm		Not Available	
ethylbenzene	800 ppm		Not Available	
methyl isobutyl ketone	500 ppm		Not Available	
1,3,5-trimethyl benzene	Not Available		Not Available	
heptane	750 ppm		Not Available	
n-hexane	1,100 ppm		Not Available	
Occupational Exposure Banding	9		-	
Ingredient	Occupational Exposure Band Rating		Occupational Expo	osure Band Limit
1,3,5-trimethyl benzene	E		≤ 0.1 ppm	
· ·	1			

Occupational exposure banding is a process of assigning chemicals into specific categories or bands based on a chemical's potency and the adverse health outcomes associated with exposure. The output of this process is an occupational exposure band (OEB), which corresponds to a range of exposure concentrations that are expected to protect worker health.

Exposure controls

Notes:

Appropriate engineering controls	CARE: Use of a quantity of this material in confined space or poorly ventilated area, where rapid build up of concentrated atmosphere may occur, could require increased ventilation and/or protective gear Engineering controls are used to remove a hazard or place a barrier between the worker and the hazard. Well-designed engineering controls can be highly effective in protecting workers and will typically be independent of worker interactions to provide this high level of protection. The basic types of engineering controls are: Process controls which involve changing the way a job activity or process is done to reduce the risk. Enclosure and/or isolation of emission source which keeps a selected hazard "physically" away from the worker and ventilation that strategically "adds" and "removes" air in the work environment.
Personal protection	
Eye and face protection	 Safety glasses with unperforated side shields may be used where continuous eye protection is desirable, as in laboratories; spectacles are not sufficient where complete eye protection is needed such as when handling bulk-quantities, where there is a danger of splashing, or if the material may be under pressure. Chemical goggles.whenever there is a danger of the material coming in contact with the eyes; goggles must be properly fitted. Full face shield (20 cm, 8 in minimum) may be required for supplementary but never for primary protection of eyes; these afford face protection. Alternatively a gas mask may replace splash goggles and face shields.
Skin protection	See Hand protection below
Hands/feet protection	 Elbow length PVC gloves For esters: Do NOT use natural rubber, butyl rubber, EPDM or polystyrene-containing materials. The selection of suitable gloves does not only depend on the material, but also on further marks of quality which vary from manufacturer to manufacturer. Where the chemical is a preparation of several substances, the resistance of the glove material can not be calculated in advance and has therefore to be checked prior to the application. The exact break through time for substances has to be obtained from the manufacturer of the protective gloves and has to be observed when making a final choice. Personal hygiene is a key element of effective hand care.
Body protection	See Other protection below
Other protection	 Overalls. PVC Apron. PVC protective suit may be required if exposure severe.
	Continued

- Evewash unit.
- Some plastic personal protective equipment (PPE) (e.g. gloves, aprons, overshoes) are not recommended as they may produce static electricity.
- ▶ For large scale or continuous use wear tight-weave non-static clothing (no metallic fasteners, cuffs or pockets).
- Non sparking safety or conductive footwear should be considered. Conductive footwear describes a boot or shoe with a sole made from a conductive compound chemically bound to the bottom components, for permanent control to electrically around the foot an shall dissipate static electricity from the body to reduce the possibility of ignition of volatile compounds.

Recommended material(s)

GLOVE SELECTION INDEX

Glove selection is based on a modified presentation of the:

"Forsberg Clothing Performance Index".

The effect(s) of the following substance(s) are taken into account in the $\ computer$ generated selection.

Ardex WPM 240 Primer (Aus)

Material	CPI
BUTYL	С
BUTYL/NEOPRENE	С
CPE	C
HYPALON	С
NAT+NEOPR+NITRILE	C
NATURAL RUBBER	C
NATURAL+NEOPRENE	С
NEOPRENE	C
NEOPRENE/NATURAL	C
NITRILE	C
NITRILE+PVC	С
PE	С
PE/EVAL/PE	С
PVA	C
PVC	С
PVDC/PE/PVDC	С
SARANEX-23	C
SARANEX-23 2-PLY	С
TEFLON	C
VITON	С
VITON/BUTYL	С
VITON/CHLOROBUTYL	С
VITON/NEOPRENE	С

* CPI - Chemwatch Performance Index

A: Best Selection

B: Satisfactory; may degrade after 4 hours continuous immersion

C: Poor to Dangerous Choice for other than short term immersion

NOTE: As a series of factors will influence the actual performance of the glove, a final selection must be based on detailed observation.

* Where the glove is to be used on a short term, casual or infrequent basis, factors such as "feel" or convenience (e.g. disposability), may dictate a choice of gloves which might otherwise be unsuitable following long-term or frequent use. A qualified practitioner should be consulted.

SECTION 9 Physical and chemical properties

Information on basic physical and chemical properties

Appearance	Black highly flammable liquid; does not mix with water.		
Physical state	Liquid	Relative density (Agua= 1)	0.93 @20C
Odour	Not Available	Partition coefficient n-octanol / water	Not Available
Odour threshold	Not Available	Auto-ignition temperature (°C)	Not Available
pH (as supplied)	Not Applicable	Decomposition temperature	Not Available
Melting point / freezing point (°C)	Not Available	Viscosity (cSt)	Not Available
Initial boiling point and boiling range (°C)	80	Molecular weight (g/mol)	Not Applicable
Flash point (°C)	<23	Taste	Not Available
Evaporation rate	Not Available	Explosive properties	Not Available
Flammability	HIGHLY FLAMMABLE.	Oxidising properties	Not Available

Respiratory protection

Type AX Filter of sufficient capacity. (AS/NZS 1716 & 1715, EN 143:2000 & 149:2001, ANSI Z88 or national equivalent)

Where the concentration of gas/particulates in the breathing zone, approaches or exceeds the "Exposure Standard" (or ES), respiratory protection is required. Degree of protection varies with both face-piece and Class of filter; the nature of protection varies with Type of filter.

Required Minimum Protection Factor	Half-Face Respirator	Full-Face Respirator	Powered Air Respirator
up to 10 x ES	AX-AUS	-	AX-PAPR-AUS / Class 1
up to 50 x ES	-	AX-AUS / Class 1	-
up to 100 x ES	-	AX-2	AX-PAPR-2 ^

^ - Full-face

A(All classes) = Organic vapours, B AUS or B1 = Acid gasses, B2 = Acid gas or hydrogen cyanide(HCN), B3 = Acid gas or hydrogen cyanide(HCN), E = Sulfur dioxide(SO2), G = Agricultural chemicals, K = Ammonia(NH3), Hg = Mercury, NO = Oxides of nitrogen, MB = Methyl bromide, AX = Low boiling point organic compounds(below 65 degC)

- Cartridge respirators should never be used for emergency ingress or in areas of unknown vapour concentrations or oxygen content.
- The wearer must be warned to leave the contaminated area immediately on detecting any odours through the respirator. The odour may indicate that the mask is not functioning properly, that the vapour concentration is too high, or that the mask is not properly fitted. Because of these limitations, only restricted use of cartridge respirators is considered appropriate.
- Cartridge performance is affected by humidity. Cartridges should be changed after 2 hr of continuous use unless it is determined that the humidity is less than 75%, in which case, cartridges can be used for 4 hr. Used cartridges should be discarded daily, regardless of the length of time used

Upper Explosive Limit (%)	Not Available	Surface Tension (dyn/cm or mN/m)	Not Available
Lower Explosive Limit (%)	Not Available	Volatile Component (%vol)	60.94 (VOC)
Vapour pressure (kPa)	Not Available	Gas group	Not Available
Solubility in water	Immiscible	pH as a solution (1%)	Not Applicable
Vapour density (Air = 1)	Not Available	VOC g/L	566.74

SECTION 10 Stability and reactivity

Reactivity	See section 7
Chemical stability	 Unstable in the presence of incompatible materials. Product is considered stable. Hazardous polymerisation will not occur.
Possibility of hazardous reactions	See section 7
Conditions to avoid	See section 7
Incompatible materials	See section 7
Hazardous decomposition products	See section 5

SECTION 11 Toxicological information

Information on toxicological effects

Chronic	drowsiness, reduced colour perception, blindness, nystagmus (rapid, involuntary eye movements), hearing loss leading to deafness and mild dementia. Exposure to styrene may aggravate central nervous system disorders, chronic respiratory disease, skin disease, kidney disease and liver disease. Exposure to styrene at work causes effects on the nervous system.			
	This material can cause serious damage if one is exposed to it for lon produce severe defects. Prolonged or repeated skin contact may cause drying with cracking, in Substance accumulation, in the human body, may occur and may cau Intentional abuse (glue sniffing) or occupational exposure to toluene of tremors of the extremeties (due to widespread cerebrum withering), h	ng periods. It can be assumed that it contains a substance which can rritation and possible dermatitis following. use some concern following repeated or long-term occupational exposure. can result in chronic habituation. Chronic abuse has caused inco-ordination, needache, abnormal speech, temporary memory loss, convulsions, coma,		
Eye	There is evidence that material may produce eye irritation in some persons and produce eye damage 24 hours or more after instillation. Severe inflammation may be expected with pain. The liquid produces a high level of eye discomfort and is capable of causing pain and severe conjunctivitis. Corneal injury may develop, with possible permanent impairment of vision, if not promptly and adequately treated. There has been concern that this material can cause cancer or mutations, but there is not enough data to make an assessment.			
Skin Contact	The material may cause moderate inflammation of the skin either following direct contact or after a delay of some time. Repeated exposure can cause contact dermatitis which is characterised by redness, swelling and blistering. Repeated exposure may cause skin cracking, flaking or drying following normal handling and use. Open cuts, abraded or irritated skin should not be exposed to this material Entry into the blood-stream, through, for example, cuts, abrasions or lesions, may produce systemic injury with harmful effects. Examine the skin prior to the use of the material and ensure that any external damage is suitably protected. Skin contact with the material may damage the health of the individual; systemic effects may result following absorption.			
Ingestion	Swallowing of the liquid may cause aspiration into the lungs with the risk of chemical pneumonitis; serious consequences may result. (ICSC13733) Accidental ingestion of the material may be damaging to the health of the individual.			
Inhaled	cause further lung damage. The main effects of simple esters are irritation, stupor and insensibility. Headache, drowsiness, dizziness, coma and behavioural changes may occur. Exposure to 400ppm ethyl acetate may cause mild eye, nose and throat irritation in an unacclimated persons. However, production workers with regular exposure have better tolerance. Inhalation hazard is increased at higher temperatures. Central nervous system (CNS) depression may include general discomfort, symptoms of giddiness, headache, dizziness, nausea, anaesthetic effects, slowed reaction time, slurred speech and may progress to unconsciousness. Serious poisonings may result in respiratory depression and may be fatal. Inhalation of high concentrations of gas/vapour causes lung irritation with coughing and nausea, central nervous depression with headache and dizziness, slowing of reflexes, fatigue and inco-ordination. Inhalation of aerosols (mists, fumes), generated by the material during the course of normal handling, may be damaging to the health of the individual.			

	TOXICITY	IRRITATION
	Dermal (rabbit) LD50: >1700 mg/kg ^[2]	Eye (human): 200 ppm irritant
	Inhalation(Rat) LC50; 5922 ppm4 ^[1]	Eye (rabbit): 5 mg/24h SEVERE
xylene	Oral(Rat) LD50; 11.494 mg/kg ^[1]	Eye (rabbit): 87 mg mild
-		Eye: adverse effect observed (irritating) ^[1]
		Skin (rabbit):500 mg/24h moderate
		Skin: adverse effect observed (irritating) ^[1]
	ΤΟΧΙΟΙΤΥ	IRRITATION
	Dermal (rabbit) LD50: >5000 mg/kg ^[1]	Eye (rabbit): 2mg/24h - SEVERE
	Inhalation(Rat) LC50; 12.528.8 mg/l4 ^[2]	Eye (rabbit):0.87 mg - mild
	Oral(Rat) LD50; 636 mg/kg ^[2]	Eye (rabbit):100 mg/30sec - mild
toluene		Eye: adverse effect observed (irritating) ^[1]
		Skin (rabbit):20 mg/24h-moderate
		Skin (rabbit):500 mg - moderate
		Skin: adverse effect observed (irritating) ^[1]
		Skin: no adverse effect observed (not irritating) ^[1]
	ΤΟΧΙΟΙΤΥ	IRRITATION
nonhtha natrolaum light	Dermal (rabbit) LD50: >1900 mg/kg ^[1]	Eye: no adverse effect observed (not irritating) ^[1]
naphtha petroleum, light aromatic solvent	Inhalation(Rat) LC50; >4.42 mg/L4 ^[1]	Skin: adverse effect observed (irritating) ^[1]
	Oral(Rat) LD50; >4500 mg/kg ^[1]	
	ΤΟΧΙϹΙΤΥ	IRRITATION
	Dermal (rabbit) LD50: >14100 mg/kg ^[2]	Eye (human): 300 mg
	Inhalation(Rat) LC50; 0.74 mg/l4 ^[2]	Eye (rabbit): 20 mg (open)-SEVERE
n-butyl acetate	Oral(Rat) LD50; 13.864 mg/kg ^[1]	Eye (rabbit): 20 mg/24h - moderate
		Eye: no adverse effect observed (not irritating) ^[1]
		Skin (rabbit): 500 mg/24h-moderate
		Skin (rabbit): 500 mg/24h-moderate Skin: no adverse effect observed (not irritating) ^[1]
	ΤΟΧΙΟΙΤΥ	
		Skin: no adverse effect observed (not irritating) ^[1]
ethyl acetate	Dermal (rabbit) LD50: >22.222 mg/kg ^[2]	Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 400 ppm
ethyl acetate		Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 400 ppm Eye: no adverse effect observed (not irritating) ^[1]
ethyl acetate	Dermal (rabbit) LD50: >22.222 mg/kg ^[2] Inhalation(Mouse) LC50; >18 mg/l4 ^[1] Oral(Rat) LD50; 12.556 mg/kg ^[1]	Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 400 ppm Eye: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1]
ethyl acetate	Dermal (rabbit) LD50: >22.222 mg/kg ^[2] Inhalation(Mouse) LC50; >18 mg/l4 ^[1] Oral(Rat) LD50; 12.556 mg/kg ^[1] TOXICITY	Skin: no adverse effect observed (not irritating)[1] IRRITATION Eye (human): 400 ppm Eye: no adverse effect observed (not irritating)[1] Skin: no adverse effect observed (not irritating)[1] IRRITATION
ethyl acetate	Dermal (rabbit) LD50: >22.222 mg/kg ^[2] Inhalation(Mouse) LC50; >18 mg/l4 ^[1] Oral(Rat) LD50; 12.556 mg/kg ^[1] TOXICITY Dermal (rabbit) LD50: >11.899 mg/kg ^[1]	Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 400 ppm Eye: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 500 ppm - irritant
ethyl acetate	Dermal (rabbit) LD50: >22.222 mg/kg ^[2] Inhalation(Mouse) LC50; >18 mg/l4 ^[1] Oral(Rat) LD50; 12.556 mg/kg ^[1] TOXICITY Dermal (rabbit) LD50: >11.899 mg/kg ^[1] Inhalation(Mouse) LC50; 44 mg/L4 ^[2]	Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 400 ppm Eye: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 500 ppm - irritant Eye (rabbit): 20mg/24hr -moderate
ethyl acetate	Dermal (rabbit) LD50: >22.222 mg/kg ^[2] Inhalation(Mouse) LC50; >18 mg/l4 ^[1] Oral(Rat) LD50; 12.556 mg/kg ^[1] TOXICITY Dermal (rabbit) LD50: >11.899 mg/kg ^[1]	Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 400 ppm Eye: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 500 ppm - irritant Eye (rabbit): 20mg/24hr -moderate Eye (rabbit): 3.95 mg - SEVERE
	Dermal (rabbit) LD50: >22.222 mg/kg ^[2] Inhalation(Mouse) LC50; >18 mg/l4 ^[1] Oral(Rat) LD50; 12.556 mg/kg ^[1] TOXICITY Dermal (rabbit) LD50: >11.899 mg/kg ^[1] Inhalation(Mouse) LC50; 44 mg/L4 ^[2]	Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 400 ppm Eye: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 500 ppm - irritant Eye (rabbit): 20mg/24hr -moderate Eye (rabbit): 3.95 mg - SEVERE Eye: adverse effect observed (irritating) ^[1]
	Dermal (rabbit) LD50: >22.222 mg/kg ^[2] Inhalation(Mouse) LC50; >18 mg/l4 ^[1] Oral(Rat) LD50; 12.556 mg/kg ^[1] TOXICITY Dermal (rabbit) LD50: >11.899 mg/kg ^[1] Inhalation(Mouse) LC50; 44 mg/L4 ^[2]	Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 400 ppm Eye: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 500 ppm - irritant Eye (rabbit): 20mg/24hr -moderate Eye (rabbit): 3.95 mg - SEVERE Eye: adverse effect observed (irritating) ^[1] Skin (rabbit): 500 mg/24hr - mild
	Dermal (rabbit) LD50: >22.222 mg/kg ^[2] Inhalation(Mouse) LC50; >18 mg/l4 ^[1] Oral(Rat) LD50; 12.556 mg/kg ^[1] TOXICITY Dermal (rabbit) LD50: >11.899 mg/kg ^[1] Inhalation(Mouse) LC50; 44 mg/L4 ^[2]	Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 400 ppm Eye: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 500 ppm - irritant Eye (rabbit): 20mg/24hr -moderate Eye (rabbit): 3.95 mg - SEVERE Eye: adverse effect observed (irritating) ^[1] Skin (rabbit): 500 mg/24hr - mild Skin (rabbit): 395mg (open) - mild
	Dermal (rabbit) LD50: >22.222 mg/kg ^[2] Inhalation(Mouse) LC50; >18 mg/l4 ^[1] Oral(Rat) LD50; 12.556 mg/kg ^[1] TOXICITY Dermal (rabbit) LD50: >11.899 mg/kg ^[1] Inhalation(Mouse) LC50; 44 mg/L4 ^[2]	Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 400 ppm Eye: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 500 ppm - irritant Eye (rabbit): 20mg/24hr -moderate Eye (rabbit): 3.95 mg - SEVERE Eye: adverse effect observed (irritating) ^[1] Skin (rabbit): 500 mg/24hr - mild
	Dermal (rabbit) LD50: >22.222 mg/kg ^[2] Inhalation(Mouse) LC50; >18 mg/l4 ^[1] Oral(Rat) LD50; 12.556 mg/kg ^[1] TOXICITY Dermal (rabbit) LD50: >11.899 mg/kg ^[1] Inhalation(Mouse) LC50; 44 mg/L4 ^[2]	Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 400 ppm Eye: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 500 ppm - irritant Eye (rabbit): 20mg/24hr -moderate Eye (rabbit): 3.95 mg - SEVERE Eye: adverse effect observed (irritating) ^[1] Skin (rabbit): 500 mg/24hr - mild Skin (rabbit): 395mg (open) - mild
	Dermal (rabbit) LD50: >22.222 mg/kg ^[2] Inhalation(Mouse) LC50; >18 mg/l4 ^[1] Oral(Rat) LD50; 12.556 mg/kg ^[1] TOXICITY Dermal (rabbit) LD50: >11.899 mg/kg ^[1] Inhalation(Mouse) LC50; 44 mg/L4 ^[2] Oral(Rat) LD50; 2.785 mg/kg ^[1]	Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 400 ppm Eye: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 500 ppm - irritant Eye (rabbit): 20mg/24hr -moderate Eye (rabbit): 3.95 mg - SEVERE Eye: adverse effect observed (irritating) ^[1] Skin (rabbit): 500 mg/24hr - mild Skin (rabbit): 395mg (open) - mild Skin: no adverse effect observed (not irritating) ^[1]
	Dermal (rabbit) LD50: >22.222 mg/kg ^[2] Inhalation(Mouse) LC50; >18 mg/l4 ^[1] Oral(Rat) LD50; 12.556 mg/kg ^[1] TOXICITY Dermal (rabbit) LD50: >11.899 mg/kg ^[1] Inhalation(Mouse) LC50; 44 mg/L4 ^[2] Oral(Rat) LD50; 2.785 mg/kg ^[1] Inhalation(Mouse) LC50; 44 mg/L4 ^[2] Oral(Rat) LD50; 2.785 mg/kg ^[1]	Skin: no adverse effect observed (not irritating)[1] IRRITATION Eye (human): 400 ppm Eye: no adverse effect observed (not irritating)[1] Skin: no adverse effect observed (not irritating)[1] Skin: no adverse effect observed (not irritating)[1] IRRITATION Eye (human): 500 ppm - irritant Eye (rabbit): 20mg/24hr -moderate Eye (rabbit): 3.95 mg - SEVERE Eye: adverse effect observed (irritating)[1] Skin (rabbit): 500 mg/24hr - mild Skin (rabbit): 395mg (open) - mild Skin: no adverse effect observed (not irritating)[1] IRRITATION
acetone	Dermal (rabbit) LD50: >22.222 mg/kg ^[2] Inhalation(Mouse) LC50; >18 mg/l4 ^[1] Oral(Rat) LD50; 12.556 mg/kg ^[1] TOXICITY Dermal (rabbit) LD50: >11.899 mg/kg ^[1] Inhalation(Mouse) LC50; 44 mg/L4 ^[2] Oral(Rat) LD50; 2.785 mg/kg ^[1] Inhalation(Mouse) LC50; 44 mg/L4 ^[2] Oral(Rat) LD50; 2.785 mg/kg ^[1] TOXICITY Dermal (rabbit) LD50: >12.346 mg/kg ^[1]	Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 400 ppm Eye: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 500 ppm - irritant Eye (rabbit): 20mg/24hr -moderate Eye (rabbit): 3.95 mg - SEVERE Eye: adverse effect observed (irritating) ^[1] Skin (rabbit): 500 mg/24hr - mild Skin (rabbit): 395mg (open) - mild Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 500 mg/24hr - mild Skin (rabbit): 395mg (open) - mild Skin: no adverse effect observed (not irritating) ^[1] Eye (human): 350 ppm -irritant
acetone	Dermal (rabbit) LD50: >22.222 mg/kg ^[2] Inhalation(Mouse) LC50; >18 mg/l4 ^[1] Oral(Rat) LD50; 12.556 mg/kg ^[1] TOXICITY Dermal (rabbit) LD50: >11.899 mg/kg ^[1] Inhalation(Mouse) LC50; 44 mg/L4 ^[2] Oral(Rat) LD50; 2.785 mg/kg ^[1] Inhalation(Mouse) LC50; 44 mg/L4 ^[2] Oral(Rat) LD50; 2.785 mg/kg ^[1] Inhalation(Mouse) LC50; 32 mg/kg ^[1]	Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 400 ppm Eye: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 500 ppm - irritant Eye (rabbit): 20mg/24hr -moderate Eye (rabbit): 3.95 mg - SEVERE Eye: adverse effect observed (irritating) ^[1] Skin (rabbit): 500 mg/24hr - mild Skin (rabbit): 395mg (open) - mild Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 350 ppm -irritant Eye (rabbit): 395mg (open) - mild Skin (rabbit): 395mg (open) - mild Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 350 ppm -irritant Eye (rabbit): 80 mg - irritant
acetone	Dermal (rabbit) LD50: >22.222 mg/kg ^[2] Inhalation(Mouse) LC50; >18 mg/l4 ^[1] Oral(Rat) LD50; 12.556 mg/kg ^[1] TOXICITY Dermal (rabbit) LD50: >11.899 mg/kg ^[1] Inhalation(Mouse) LC50; 44 mg/L4 ^[2] Oral(Rat) LD50; 2.785 mg/kg ^[1] Inhalation(Mouse) LC50; 44 mg/L4 ^[2] Oral(Rat) LD50; 2.785 mg/kg ^[1] Inhalation(Mouse) LC50; 32 mg/kg ^[1] Inhalation(Mouse) LC50; 32 mg/L4 ^[2] Oral(Rat) LD50; 2054 mg/kg ^[1]	Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 400 ppm Eye: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 500 ppm - irritant Eye (rabbit): 20mg/24hr -moderate Eye (rabbit): 3.95 mg - SEVERE Eye (rabbit): 500 mg/24hr - mild Skin (rabbit): 500 mg/24hr - mild Skin (rabbit): 395mg (open) - mild Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 350 ppm -irritant Eye (rabbit): 30 mg - irritant Eye (numan): 350 ppm -irritant Eye (rabbit): 80 mg - irritant Skin (rabbit): 402 mg/24 hr - mild Skin (rabbit): 402 mg/24 hr - mild Skin (rabbit): 13.78mg/24 hr open
acetone	Dermal (rabbit) LD50: >22.222 mg/kg ^[2] Inhalation(Mouse) LC50; >18 mg/l4 ^[1] Oral(Rat) LD50; 12.556 mg/kg ^[1] TOXICITY Dermal (rabbit) LD50: >11.899 mg/kg ^[1] Inhalation(Mouse) LC50; 44 mg/L4 ^[2] Oral(Rat) LD50; 2.785 mg/kg ^[1] Inhalation(Mouse) LC50; 44 mg/L4 ^[2] Oral(Rat) LD50; 2.785 mg/kg ^[1] Dermal (rabbit) LD50: >12.346 mg/kg ^[1] Inhalation(Mouse) LC50; 32 mg/L4 ^[2] Oral(Rat) LD50; 2054 mg/kg ^[1] Inhalation(Mouse) LC50; 32 mg/L4 ^[2] Oral(Rat) LD50; 2054 mg/kg ^[1]	Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 400 ppm Eye: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 500 ppm - irritant Eye (rabbit): 20mg/24hr -moderate Eye (rabbit): 3.95 mg - SEVERE Eye (rabbit): 500 mg/24hr - mild Skin (rabbit): 500 mg/24hr - mild Skin (rabbit): 395mg (open) - mild Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 350 ppm -irritant Eye (rabbit): 30 mg - irritant Eye (rabbit): 30 mg - irritant Eye (rabbit): 80 mg - irritant Eye (rabbit): 80 mg - irritant Skin (rabbit): 402 mg/24 hr - mild
acetone methyl ethyl ketone	Dermal (rabbit) LD50: >22.222 mg/kg ^[2] Inhalation(Mouse) LC50; >18 mg/kg ^[1] Oral(Rat) LD50; 12.556 mg/kg ^[1] TOXICITY Dermal (rabbit) LD50: >11.899 mg/kg ^[1] Inhalation(Mouse) LC50; 44 mg/L4 ^[2] Oral(Rat) LD50; 2.785 mg/kg ^[1] Inhalation(Mouse) LC50; 44 mg/L4 ^[2] Oral(Rat) LD50; 2.785 mg/kg ^[1] Inhalation(Mouse) LC50; 32 mg/kg ^[1] Oral(Rat) LD50: >12.346 mg/kg ^[1] Inhalation(Mouse) LC50; 32 mg/L4 ^[2] Oral(Rat) LD50; 2054 mg/kg ^[1] Inhalation(Mouse) LC50; 32 mg/L4 ^[2] Oral(Rat) LD50; 2054 mg/kg ^[1] Inhalation(Mouse) LC50; 32 mg/L4 ^[2] Oral(Rat) LD50; 2054 mg/kg ^[1]	Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 400 ppm Eye: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 500 ppm - irritant Eye (rabbit): 20mg/24hr -moderate Eye (rabbit): 3.95 mg - SEVERE Eye: adverse effect observed (irritating) ^[1] Skin (rabbit): 500 mg/24hr - mild Skin (rabbit): 395mg (open) - mild Skin: no adverse effect observed (not irritating) ^[1] Skin (rabbit): 395mg (open) - mild Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 350 ppm -irritant Eye (rabbit): 80 mg - irritant Eye (rabbit): 402 mg/24 hr - mild Skin (rabbit): 402 mg/24 hr open IRRITATION Eye (rabbit): 13.78mg/24 hr open IRRITATION Eye (rabbit): 100 mg/24h - moderate
acetone	Dermal (rabbit) LD50: >22.222 mg/kg ^[2] Inhalation(Mouse) LC50; >18 mg/l4 ^[1] Oral(Rat) LD50; 12.556 mg/kg ^[1] TOXICITY Dermal (rabbit) LD50: >11.899 mg/kg ^[1] Inhalation(Mouse) LC50; 44 mg/L4 ^[2] Oral(Rat) LD50; 2.785 mg/kg ^[1] Inhalation(Mouse) LC50; 44 mg/L4 ^[2] Oral(Rat) LD50; 2.785 mg/kg ^[1] Dermal (rabbit) LD50: >12.346 mg/kg ^[1] Inhalation(Mouse) LC50; 32 mg/L4 ^[2] Oral(Rat) LD50; 2054 mg/kg ^[1] Inhalation(Mouse) LC50; 32 mg/L4 ^[2] Oral(Rat) LD50; 2054 mg/kg ^[1]	Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 400 ppm Eye: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1] Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 500 ppm - irritant Eye (rabbit): 20mg/24hr -moderate Eye (rabbit): 3.95 mg - SEVERE Eye: adverse effect observed (irritating) ^[1] Skin (rabbit): 500 mg/24hr - mild Skin (rabbit): 395mg (open) - mild Skin: no adverse effect observed (not irritating) ^[1] Skin (rabbit): 395mg (open) - mild Skin: no adverse effect observed (not irritating) ^[1] IRRITATION Eye (human): 350 ppm -irritant Eye (rabbit): 80 mg - irritant Eye (rabbit): 102 mg/24 hr - mild Skin (rabbit): 402 mg/24 hr - mild Skin (rabbit): 13.78mg/24 hr open IRRITATION IRRITATION

	TOXICITY	IRRITATION
	Dermal (rabbit) LD50: ~20.531 mg/kg ^[1]	Eye (rabbit): 500 mg - SEVERE
ethylbenzene	Inhalation(Rat) LC50; 17.2 mg/l4 ^[2]	Eye: no adverse effect observed (not irritating) ^[1]
	Oral(Rat) LD50; ~3523 mg/kg ^[2]	Skin (rabbit): 15 mg/24h mild
		Skin: no adverse effect observed (not irritating) ^[1]
	τοχιςιτγ	IRRITATION
	Dermal (rabbit) LD50: >25 mg/kg ^[1]	Eye (human): 200 ppm/15m
methyl isobutyl ketone	Inhalation(Rat) LC50; ~8.216.4 mg/l4 ^[2]	Eye (rabbit): 40 mg - SEVERE
	Oral(Rat) LD50; 4.663 mg/kg ^[1]	Eye (rabbit): 500 mg/24h - mild
		Skin (rabbit): 500 mg/24h - mild
	τοχιςιτγ	IRRITATION
	dermal (rat) LD50: >4.624 mg/kg ^[1]	Eye (rabbit): 500 mg/24h mild
1,3,5-trimethyl benzene	Inhalation(Rat) LC50; 10.2 mg/L4 ^[1]	Eye: adverse effect observed (irritating) ^[1]
	Oral(Rat) LD50; 6000 mg/kg ^[1]	Skin (rabbit): 20 mg/24h moderate
		Skin: adverse effect observed (irritating) ^[1]
	ΤΟΧΙΟΙΤΥ	IRRITATION
	Dermal (rabbit) LD50: >2000 mg/kg ^[1]	Eye: no adverse effect observed (not irritating) ^[1]
heptane	Inhalation(Rat) LC50; >29.29 mg/l4 ^[1]	Skin: no adverse effect observed (not irritating) ^[1]
	Oral(Rat) LD50; >5000 mg/kg ^[1]	
	ΤΟΧΙΟΙΤΥ	IRRITATION
	Dermal (rabbit) LD50: >7.572 mg/kg ^[1]	Eye(rabbit): 10 mg - mild
n-hexane	Inhalation(Rat) LC50; 48000 ppm4 ^[2]	
	Oral(Rat) LD50; 36.347 mg/kg ^[1]	
Legend:	1. Value obtained from Europe ECHA Registered Substances - , specified data extracted from RTECS - Register of Toxic Effect of	Acute toxicity 2.* Value obtained from manufacturer's SDS. Unless otherwise f chemical Substances
XYLENE	Reproductive effector in rats The substance is classified by IARC as Group 3: NOT classifiable as to its carcinogenicity to humans. Evidence of carcinogenicity may be inadequate or limited in anim	nal testing.
TOLUENE	from headaches to intoxication, convulsions, narcosis (sleepines nervous system depression, and in large doses has a narcotic e congestion and bleeding of the lungs and kidney injury were all f	rt periods of time experience adverse central nervous system effects ranging s) and death. When inhaled or swallowed, toluene can cause severe central ffect. 60mL has caused death. Death of heart muscle fibres, liver swelling, ound on autopsy. I for 8 hours resulted in the same and more serious symptoms including eupho
NAPHTHA PETROLEUM, LIGHT AROMATIC SOLVENT	and dermal (LD50 in rabbits > 2000 mg/kg-bw) routes of exposu Most LBPNs are mild to moderate eye and skin irritants in rabbit naphthas, which have higher primary skin irritation indices. Sensitisation: LBPNs do not appear to be skin sensitizers, but a poor response Repeat dose toxicity: The lowest-observed-adverse-effect concentration (LOAEC) and short-term (2-89 days) and subchronic (greater than 90 days) ex endpoints after considering the toxicity data for all LBPNs in the Renal effects, including increased kidney weight, renal lesions (r rats exposed orally or by inhalation to most LBPNs, were consid	s, with the exception of heavy catalytic cracked and heavy catalytic reformed e in the positive control was also noted in these studies I lowest-observed-adverse-effect level (LOAEL) values identified following posure to the LBPN substances. These values were determined for a variety group. Most of the studies were carried out by the inhalation route of exposure enal tubule dilation, necrosis) and hyaline droplet formation, observed in male ered species- and sex-specific These effects were determined to be due to a eraction between hydrocarbon metabolites and alpha-2-microglobulin, an

Animal studies indicate that normal, branched and cyclic paraffins are absorbed from the gastrointestinal tract and that the absorption of n-paraffins is inversely proportional to the carbon chain length, with little absorption above C30. With respect to the carbon chain lengths likely to be present in mineral oil, n-paraffins may be absorbed to a greater extent than iso- or cyclo-paraffins.

The major classes of hydrocarbons are well absorbed into the gastrointestinal tract in various species. In many cases, the hydrophobic hydrocarbons are ingested in association with fats in the diet. Some hydrocarbons may appear unchanged as in the lipoprotein particles in the gut lymph, but most hydrocarbons partly separate from fats and undergo metabolism in the gut cell. For C9 aromatics (typically trimethylbenzenes – TMBs)

Acute toxicity: Animal testing shows that semi-lethal concentrations and doses vary amongst this group. The semilethal concentrations for inhalation range from 6000 to 10000 mg/cubic metre for C9 aromatic naphtha and 18000-24000 mg/cubic metre for 1,2,4- and 1,3,5-TMB, respectively.

Irritation and sensitization: Results from animal testing indicate that C9 aromatic hydrocarbon solvents are mildly to moderately irritating to the skin, minimally irritating to the eye, and have the potential to irritate the airway and cause depression of breathing rate. There is no evidence that it sensitizes skin.

	be relevant in humans. Mutation-causing potential: Most studies involving gasoline have returned negative results r all recent studies in living human subjects (such as in petrol service station attendants).	egarding the potential to cause mutations, including			
N-BUTYL ACETATE	Generally,linear and branched-chain alkyl esters are hydrolysed to their component alcohols and carboxylic acids in the intestinal tract, blood and most tissues throughout the body. Following hydrolysis the component alcohols and carboxylic acids are metabolized Oral acute toxicity studies have been reported for 51 of the 67 esters of aliphatic acyclic primary alcohols and aliphatic linear saturated carboxylic acids. The very low oral acute toxicity of this group of esters is demonstrated by oral LD50 values greater than 1850 mg/kg bw Genotoxicity studies have been performed in vitro using the following esters of aliphatic acyclic primary alcohols and aliphatic linear saturated carboxylic acids: methyl acetate, butyl acetate, butyl stearate and the structurally related isoamyl formate and demonstrates that these substances are not genotoxic. The JEFCA Committee concluded that the substances in this group would not present safety concerns at the current levels of intake the esters of aliphatic acyclic primary alcohols and aliphatic linear saturated carboxylic acids are generally used as flavouring substances up to average maximum levels of 200 mg/kg. Higher levels of use (up to 3000 mg/kg) are permitted in food categories such as chewing gum and hard candy.				
ACETONE	For acetone: The acute toxicity of acetone is low. Acetone is not a skin irritant or sensitizer, but it removes fat from the skin, and it also irritates the eye. Animal testing shows acetone may cause macrocytic anaemia. Studies in humans have shown that exposure to acetone at a level of 2375 mg/cubic metre has not caused neurobehavioural deficits.				
METHYL ETHYL KETONE	Methyl ethyl ketone is considered to have a low order of toxicity; however, methyl ethyl ketone is often used in combination with other solvents and the mixture may have greater toxicity than either solvent alone. Combinations of n-hexane with methyl ethyl ketone, and also methyl n-butyl ketone with methyl ethyl ketone may result in an increased in peripheral neuropathy, a progressive disorder of the nerves of the extremities. Combinations with chloroform also show an increase in toxicity.				
ETHYLBENZENE	Liver changes, utheral tract, effects on fertility, foetotoxicity, specific developmental abnormalities (musculoskeletal system) recorded. Ethylbenzene is readily absorbed when inhaled, swallowed or in contact with the skin. It is distributed throughout the body, and passed out through urine. It may irritate the skin, eyes and may cause hearing loss if exposed to high doses. Long Term exposure may cause damage to the kidney, liver and lungs, including a tendency to cancer formation, according to animal testing. NOTE: Substance has been shown to be mutagenic in at least one assay, or belongs to a family of chemicals producing damage or change to cellular DNA.				
METHYL ISOBUTYL KETONE	MIBK is primarily absorbed by the lungs in animals and humans but can be absorbed by the skin, stomach and gut. If inhaled, it may be found in the brain, liver, lung, vitreous fluid, kidney and blood. Oral and respiratory routes of exposure are of minimal effect with changes seen only in the liver and kidney. MIBK does not cause genetic damage or harm the foetus or offspring, and has low toxicity to aquatic organisms.				
1,3,5-TRIMETHYL BENZENE	Other Toxicity data is available for CHEMWATCH 12171 1,2,4-trimethylbenzene CHEMWAT	CH 12172 1,2,3-trimethylbenzene			
XYLENE & N-BUTYL ACETATE & ETHYLBENZENE	The material may produce severe irritation to the eye causing pronounced inflammation. Re	peated or prolonged exposure to irritants may			
XYLENE & TOLUENE & N-BUTYL ACETATE & ACETONE & METHYL ETHYL KETONE & STYRENE & ETHYLBENZENE & METHYL ISOBUTYL KETONE & 1,3,5- TRIMETHYL BENZENE	produce conjunctivitis. The material may cause skin irritation after prolonged or repeated exposure and may produ vesicles, scaling and thickening of the skin.	ce on contact skin redness, swelling, the production of			
NAPHTHA PETROLEUM, LIGHT AROMATIC SOLVENT & 1,3,5-TRIMETHYL BENZENE	For trimethylbenzenes: Absorption of 1,2,4-trimethylbenzene occurs after exposure by swallowing, inhalation, or skin contact. In the workplace, inhalation and skin contact are the most important routes of absorption; whole-body toxic effects from skin absorption are unlikely to occur as the skin irritation caused by the chemical generally leads to quick removal. The substance is fat-soluble and may accumulate in fatty tissues. It is also bound to red blood cells in the bloodstream.				
ETHYL ACETATE & METHYL ETHYL KETONE & METHYL ISOBUTYL KETONE & 1,3,5- TRIMETHYL BENZENE	Asthma-like symptoms may continue for months or even years after exposure to the material ends. This may be due to a non-allergic condition known as reactive airways dysfunction syndrome (RADS) which can occur after exposure to high levels of highly irritating compound. Main criteria for diagnosing RADS include the absence of previous airways disease in a non-atopic individual, with sudden onset of persistent asthma-like symptoms within minutes to hours of a documented exposure to the irritant. Other criteria for diagnosis of RADS include a reversible airflow pattern on lung function tests, moderate to severe bronchial hyperreactivity on methacholine challenge testing, and the lack of minimal lymphocytic inflammation, without eosinophilia.				
STYRENE & ETHYLBENZENE & METHYL ISOBUTYL KETONE	WARNING: This substance has been classified by the IARC as Group 2B: Possibly Carcino	genic to Humans.			
1,3,5-TRIMETHYL BENZENE & N-HEXANE	The material may be irritating to the eye, with prolonged contact causing inflammation. Rep conjunctivitis.	eated or prolonged exposure to irritants may produce			
Acute Toxicity	× Carcinogenicity	✓			
Skin Irritation/Corrosion	✓ Reproductivity	×			
Serious Eye Damage/Irritation	✓ STOT - Single Exposure	×			
Respiratory or Skin sensitisation	X STOT - Repeated Exposure	×			

SECTION 12 Ecological information

Toxicity

Not Available	Not AvailableTest Duration (hr)489672Not ReportedTest Duration (hr)964896969672Test Duration (hr)7296	F C	Not Available Species Crustacea Fish Algae or other aquatic plants Fish Species Fish Crustacea Crustacea Algae or other aquatic plants	3.78mg 0.104n	-	Not Available 2 2 2 4 Source 4 5
EC50 EC50 EC50(ECx) Endpoint EC50 EC5	48 96 72 Not Reported Test Duration (hr) 96 48 96 96 96 Test Duration (hr) 72	F C	Crustacea Fish Algae or other aquatic plants Fish Species Fish Crustacea Crustacea	>1.055 3.78mg 0.104n	1.8mg/l 2.6mg/l 4.6mg/l 0.017mg/L i<1.809mg/L g/L	2 2 2 4 Source 4 5
EC50 EC50 EC50(ECx) Endpoint EC50 EC5	48 96 72 Not Reported Test Duration (hr) 96 48 96 96 96 Test Duration (hr) 72	F C	Crustacea Fish Algae or other aquatic plants Fish Species Fish Crustacea Crustacea	>1.055 3.78mg 0.104n	2.6mg/l 4.6mg/l 0.017mg/L s<1.809mg/L g/L	2 2 4 Source 4 5
EC50 EC50(ECx) Endpoint C50 EC50 EC50 EC50 EC50 EC50 EC50 EC50	72 Not Reported Test Duration (hr) 96 48 96 96 96 Test Duration (hr) 72	F C	Algae or other aquatic plants Fish Species Fish Crustacea Crustacea	>1.055 3.78mg 0.104n	4.6mg/l 0.017mg/L 6<1.809mg/L g/L	2 4 Sourc 4 5
EC50(ECx) Endpoint C50 EC50 EC50 EC50 Endpoint EC50 EC50 EC50 EC50 EC50	Not Reported Test Duration (hr) 96 48 96 96 96 96 96 72	F C	Fish Species Crustacea Crustacea	>1.055 3.78mg 0.104n	0.017mg/L 6<1.809mg/L g/L	4 Source 4 5
Endpoint .C50 EC50 NOEC(ECx) EC50 EC50 EC50 EC50 NOEC(ECx)	Test Duration (hr) 96 48 96 96 96 96 96 72	F C	Species Fish Crustacea Crustacea	>1.055 3.78mg 0.104n	i<1.809mg/L g/L	Source 4 5
C50 EC50 NOEC(ECx) EC50 EC50 EC50 EC50 NOEC(ECx)	96 48 96 96 96 96 Test Duration (hr) 72 72	F C	rustacea Crustacea	>1.055 3.78mg 0.104n	g/L	4 5
EC50 NOEC(ECx) EC50 Endpoint EC50 EC50 EC50 NOEC(ECx)	48 96 96 Test Duration (hr) 72	((Crustacea Crustacea	3.78mg 0.104n	g/L	5
NOEC(ECx) EC50 Endpoint EC50 EC50 EC50 NOEC(ECx)	96 96 Test Duration (hr) 72	C	Crustacea	0.104n	-	
EC50 Endpoint EC50 EC50 NOEC(ECx)	96 Test Duration (hr) 72				na/L	
Endpoint EC50 EC50 NOEC(ECx)	Test Duration (hr) 72	Α	lgae or other aquatic plants	1 620	5	4
EC50 EC50 NOEC(ECx)	72			~1.032	2mg/L	4
EC50 EC50 NOEC(ECx)	72		Species		Value	Sourc
NOEC(ECx)	96		Algae or other aquatic plants		19mg/l	1
			Algae or other aquatic plants		64mg/l	2
	72		Algae or other aquatic plants		1mg/l	1
	48		Crustacea		6.14mg/l	1
Endpoint	Test Duration (hr)		Species		Value	Sourc
EC50	48		Crustacea		32mg/l	1
_C50	96		Fish		18mg/l	2
EC50	72		Algae or other aquatic plants		246mg/l	2
EC50(ECx)	96		Fish		18mg/l	2
Endpoint	Test Duration (hr)		Species		Value	Sourc
EC50	48		Crustacea		164mg/l	1
_C50	96		Fish		>75.6mg/l	2
NOEC(ECx)	72		Algae or other aquatic plants		>100mg/l	1
Endpoint			Value)	Sourc	
_C50	96		Fish	13.30)3mg/L	4
NOEC(ECx)	12		Fish	0.001	mg/L	4
EC50	48		Crustacea	6098.4mg/L		5
EC50	96		Algae or other aquatic plants	9.873	3-27.684mg/l	4
Endpoint	Test Duration (hr)		Species Va		Value	Sourc
NOEC(ECx)	96		Fish		1.18mg/L	4
_C50	96		Fish		>1.18mg/L	4
EC50	48		Crustacea		308mg/l	2
EC50	72		Algae or other aquatic plants		1972mg/l	2
EC50	96		Algae or other aquatic plants		>500mg/l	4
Endpoint	Test Duration (hr)		-			Sourc
EC50	48				-	1
_C50	96					4
NOEC(ECx)	96				-	1
EC50	72				-	1
_000	JU		Augae of other aquatic plants	0.72	ing/i	1
Endpoint	Test Duration (hr)		-			Sourc
EC50					-	4
_C50						4
NOEC(ECx)	720				-	4
EC50	72				-	4
	C50 C50(ECx) adpoint C50 C50 C50 C50 C50 C50 C50 C50 C50 C50	C50 72 C50(ECx) 96 Indpoint Test Duration (hr) C50 48 C50 96 C50 48 C50 96 DEC(ECx) 72 Indpoint Test Duration (hr) DEC(ECx) 72 Indpoint Test Duration (hr) DEC(ECx) 12 C50 48 C50 96 DEC(ECx) 96 DEC(ECx) 96 C50 96 DEC(ECx) 96	C50 72 C50(ECx) 96 Indpoint Test Duration (hr) C50 48 C50 96 C50 48 C50 96 CEC(ECx) 72 Indpoint Test Duration (hr) 1 C50 96 1 C50 48 1 C50 96 1	25072Algae or other aquatic plantsC50(ECx)96FishC50(ECx)96CrustaceaC5048CrustaceaC5096FishDEC(ECx)72Algae or other aquatic plantsDEC(ECx)72SpeciesDEC(ECx)72FishDEC(ECx)96FishDEC(ECx)96FishDEC(ECx)12FishC5096Algae or other aquatic plantsDEC(ECx)12FishC5096Algae or other aquatic plantsDEC(ECx)96FishDEC(ECx)96FishC5096FishC5096FishC5096Algae or other aquatic plantsC5096CrustaceaC5096CrustaceaC5096Algae or other aquatic plantsC5096FishC5048CrustaceaC5096FishDEC(ECx)96Algae or other aquatic plantsC5048CrustaceaC5096Algae or other aquatic plantsC5096Algae or other aquatic plants	250 72 Algae or other aquatic plants C50 (ECx) 96 Fish ddpoint Test Duration (hr) Species C50 48 Crustacea C50 96 Fish DEC(ECx) 72 Algae or other aquatic plants DEC(ECx) 72 Algae or other aquatic plants DEC(ECx) 72 Species Value D50 96 Fish 13.30 DEC(ECx) 72 Fish 0.001 C50 96 Fish 0.001 D50 96 Fish 0.001 C50 48 Crustacea 6098 C50 96 Fish 0.001 D50 96 Fish 0.001 C50 96 Fish 0.001 C50 96 Fish 0.001 D50 96 Fish 0.001 C50 96 Fish 0.001 C50 96 Fish 0.002 C50 96 Algae or other aquatic plants 0.0	250 72 Algae or other aquatic plants 246mg/l C50 96 Fish 18mg/l Adpoint Test Duration (hr) Species Value C50 96 Fish >75.6mg/l D50 96 Fish >75.6mg/l D50 96 Fish >75.6mg/l D50 72 Algae or other aquatic plants >100mg/l D50 96 Fish 13.303mg/L D50 96 Fish 0.001mg/L D50 96 Fish 0.001mg/L D50 96 Crustacea 6098.4mg/L D50 96 Algae or other aquatic plants 987-27.684mg/l D50 96 Fish 1.18mg/L D50 96 Fish 1.18mg/L D50 96 Fish >1.18mg/L D50 96 Fish 0.027mg/L D50 96 Algae or other aquatic plants 1972mg/L D50 96 Algae or other aquatic plants 0.027mg/L D50 96 Algae or ot

	Endpoint	Test Duration (hr)	Species		Value	Source
	EC50	48	Crustacea		170mg/l	1
methyl isobutyl ketone	LC50	96	Fish		>179mg/l	2
	EC50(ECx)	48	Crustacea		170mg/l	1
	EC50	96	Algae or other aquatic plants		400mg/l	1
	Endpoint	Test Duration (hr)	Species		Value	Source
	LC50	96	Fish		5.216mg/l	2
	EC50	48	Crustacea	Crustacea		5
1,3,5-trimethyl benzene	BCF	1680	Fish		23-342	7
	NOEC(ECx)	384	Crustacea		0.257mg/l	2
	EC50	96	Algae or other aquatic plants		3.084mg/l	2
	Endpoint	Test Duration (hr)	Species		Value	Source
	EC50	48	Crustacea		0.64mg/l	2
heptane	LC50	96	Fish		20.179mg/L	4
	NOEC(ECx)	504	Crustacea		0.17mg/l	2
	Endpoint	Test Duration (hr)	Species	Value	9	Source
n-hexane	EC50(ECx)	240	Algae or other aquatic plants	25.02	23-137.802mg/L	4
Legend:	V3.12 (QSAR) -	Aquatic Toxicity Data (Estimated) 4. US	Registered Substances - Ecotoxicological EPA, Ecotox database - Aquatic Toxicity D apan) - Bioconcentration Data 8. Vendor D	ata 5. ECETOC		

DO NOT discharge into sewer or waterways. Harmful to aquatic organisms, may cause long-term adverse effects in the aquatic environment.

Persistence and degradability

Ingredient	Persistence: Water/Soil	Persistence: Air
xylene	HIGH (Half-life = 360 days)	LOW (Half-life = 1.83 days)
toluene	LOW (Half-life = 28 days)	LOW (Half-life = 4.33 days)
n-butyl acetate	LOW	LOW
ethyl acetate	LOW (Half-life = 14 days)	LOW (Half-life = 14.71 days)
acetone	LOW (Half-life = 14 days)	MEDIUM (Half-life = 116.25 days)
methyl ethyl ketone	LOW (Half-life = 14 days)	LOW (Half-life = 26.75 days)
styrene	HIGH (Half-life = 210 days)	LOW (Half-life = 0.3 days)
ethylbenzene	HIGH (Half-life = 228 days)	LOW (Half-life = 3.57 days)
methyl isobutyl ketone	HIGH (Half-life = 7001 days)	LOW (Half-life = 1.9 days)
1,3,5-trimethyl benzene	HIGH	HIGH
heptane	LOW	LOW
n-hexane	LOW	LOW

Bioaccumulative potential

Ingredient	Bioaccumulation
xylene	MEDIUM (BCF = 740)
toluene	LOW (BCF = 90)
n-butyl acetate	LOW (BCF = 14)
ethyl acetate	HIGH (BCF = 3300)
acetone	LOW (BCF = 0.69)
methyl ethyl ketone	LOW (LogKOW = 0.29)
styrene	LOW (BCF = 77)
ethylbenzene	LOW (BCF = 79.43)
methyl isobutyl ketone	LOW (LogKOW = 1.31)
1,3,5-trimethyl benzene	LOW (BCF = 342)
heptane	HIGH (LogKOW = 4.66)
n-hexane	MEDIUM (LogKOW = 3.9)

Mobility in soil

Ingredient	Mobility
toluene	LOW (KOC = 268)
n-butyl acetate	LOW (KOC = 20.86)
ethyl acetate	LOW (KOC = 6.131)

Continued...

Ardex WPM 240 Primer (Aus)

GH (KOC = 1.981)
EDIUM (KOC = 3.827)
W (KOC = 517.8)
W (KOC = 517.8)
W (KOC = 10.91)
W (KOC = 703)
W (KOC = 274.7)
W (KOC = 149)
~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~

# **SECTION 13 Disposal considerations**

Product / Packaging disposal	<ul> <li>Containers may still present a chemical hazard/ danger when empty.</li> <li>Return to supplier for reuse/ recycling if possible.</li> <li>Otherwise:</li> <li>If container can not be cleaned sufficiently well to ensure that residuals do not remain or if the container cannot be used to store the same product, then puncture containers, to prevent re-use, and bury at an authorised landfill.</li> <li>Where possible retain label warnings and SDS and observe all notices pertaining to the product.</li> <li>DO NOT allow wash water from cleaning or process equipment to enter drains.</li> <li>It may be necessary to collect all wash water for treatment before disposal.</li> <li>In all cases disposal to sewer may be subject to local laws and regulations and these should be considered first.</li> <li>Where in doubt contact the responsible authority.</li> <li>Recycle wherever possible.</li> <li>Consult manufacturer for recycling options or consult local or regional waste management authority for disposal if no suitable treatment or disposal facility can be identified.</li> <li>Dispose of by: burial in a land-fill specifically licensed to accept chemical and / or pharmaceutical wastes or Incineration in a licensed apparatus (after admixture with suitable combustible material).</li> <li>Decontaminate empty containers.</li> </ul>
------------------------------	------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------

# **SECTION 14 Transport information**

# Labels Required



# Land transport (ADG)

UN number	1263		
UN proper shipping name	PAINT (including paint, lacquer, enamel, stain, shellac, varnish, polish, liquid filler and liquid lacquer base) or PAINT RELATED MATERIAL (including paint thinning or reducing compound)		
Transport hazard class(es)	Class     3       Subrisk     Not Applicable		
Packing group	II		
Environmental hazard	Not Applicable		
Special precautions for user	Special provisions163 367Limited quantity5 L		

# Air transport (ICAO-IATA / DGR)

UN number	1263		
UN proper shipping name	Paint (including paint, lacquer, enamel, stain, shellac, varnish, polish, liquid filler and liquid lacquer base)		
Transport hazard class(es)	ICAO/IATA Class ICAO / IATA Subrisk ERG Code	3 Not Applicable 3L	
Packing group	11		
Environmental hazard	Not Applicable		
Special precautions for user	Special provisions Cargo Only Packing Instructions Cargo Only Maximum Qty / Pack		A3 A72 A192 364 60 L

Passenger and Cargo Packing Instructions	353
Passenger and Cargo Maximum Qty / Pack	5 L
Passenger and Cargo Limited Quantity Packing Instructions	Y341
Passenger and Cargo Limited Maximum Qty / Pack	1 L

# Sea transport (IMDG-Code / GGVSee)

UN number	1263		
UN proper shipping name	PAINT (including paint, lacquer, enamel, stain, shellac, varnish, polish, liquid filler and liquid lacquer base) or PAINT RELATED MATERIAL (including paint thinning or reducing compound)		
Transport hazard class(es)	IMDG Class     3       IMDG Subrisk     Not Applicable		
Packing group	11		
Environmental hazard	Not Applicable		
Special precautions for user	EMS NumberF-E , S-ESpecial provisions163 367Limited Quantities5 L		

# Transport in bulk according to Annex II of MARPOL and the IBC code Not Applicable

### Transport in bulk in accordance with MARPOL Annex V and the IMSBC Code

Product name	Group
xylene	Not Available
toluene	Not Available
naphtha petroleum, light aromatic solvent	Not Available
n-butyl acetate	Not Available
ethyl acetate	Not Available
acetone	Not Available
methyl ethyl ketone	Not Available
styrene	Not Available
ethylbenzene	Not Available
methyl isobutyl ketone	Not Available
1,3,5-trimethyl benzene	Not Available
heptane	Not Available
n-hexane	Not Available

# Transport in bulk in accordance with the ICG Code

Product name	Ship Type
xylene	Not Available
toluene	Not Available
naphtha petroleum, light aromatic solvent	Not Available
n-butyl acetate	Not Available
ethyl acetate	Not Available
acetone	Not Available
methyl ethyl ketone	Not Available
styrene	Not Available
ethylbenzene	Not Available
methyl isobutyl ketone	Not Available
1,3,5-trimethyl benzene	Not Available
heptane	Not Available
n-hexane	Not Available

# **SECTION 15 Regulatory information**

Safety, health and environmental regulations / legislation specific for the substance or mixture

xylene is found on the following regulatory lists

Australia Hazardous Chemical Information System (HCIS) - Hazardous Chemicals Australian Inventory of Industrial Chemicals (AIIC) International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) -Schedule 5 Monographs Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) -Schedule 6 toluene is found on the following regulatory lists Australia Hazardous Chemical Information System (HCIS) - Hazardous Chemicals Australian Inventory of Industrial Chemicals (AIIC) Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) -Chemical Footprint Project - Chemicals of High Concern List Schedule 5 International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) -Monographs Schedule 6 naphtha petroleum, light aromatic solvent is found on the following regulatory lists Australia Hazardous Chemical Information System (HCIS) - Hazardous Chemicals Chemical Footprint Project - Chemicals of High Concern List International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Australian Inventory of Industrial Chemicals (AIIC) Monographs n-butyl acetate is found on the following regulatory lists Australia Hazardous Chemical Information System (HCIS) - Hazardous Chemicals Australian Inventory of Industrial Chemicals (AIIC) ethyl acetate is found on the following regulatory lists Australia Hazardous Chemical Information System (HCIS) - Hazardous Chemicals Australian Inventory of Industrial Chemicals (AIIC) acetone is found on the following regulatory lists Australia Hazardous Chemical Information System (HCIS) - Hazardous Chemicals Australian Inventory of Industrial Chemicals (AIIC) Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) -Schedule 5 methyl ethyl ketone is found on the following regulatory lists Australia Hazardous Chemical Information System (HCIS) - Hazardous Chemicals Australian Inventory of Industrial Chemicals (AIIC) Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) -Schedule 5 styrene is found on the following regulatory lists Australia Hazardous Chemical Information System (HCIS) - Hazardous Chemicals Chemical Footprint Project - Chemicals of High Concern List Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) -International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Schedule 5 Monographs Australian Inventory of Industrial Chemicals (AIIC) International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Monographs - Group 2A: Probably carcinogenic to humans ethylbenzene is found on the following regulatory lists Australia Hazardous Chemical Information System (HCIS) - Hazardous Chemicals Chemical Footprint Project - Chemicals of High Concern List International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) -Schedule 5 Monographs Australian Inventory of Industrial Chemicals (AIIC) International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Monographs - Group 2B: Possibly carcinogenic to humans methyl isobutyl ketone is found on the following regulatory lists Australia Hazardous Chemical Information System (HCIS) - Hazardous Chemicals Chemical Footprint Project - Chemicals of High Concern List Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) -International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Schedule 5 Monographs Australian Inventory of Industrial Chemicals (AIIC) International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Monographs - Group 2B: Possibly carcinogenic to humans 1,3,5-trimethyl benzene is found on the following regulatory lists Australia Hazardous Chemical Information System (HCIS) - Hazardous Chemicals Australian Inventory of Industrial Chemicals (AIIC) Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) -Schedule 5 heptane is found on the following regulatory lists Australia Hazardous Chemical Information System (HCIS) - Hazardous Chemicals Australian Inventory of Industrial Chemicals (AIIC) n-hexane is found on the following regulatory lists Australia Hazardous Chemical Information System (HCIS) - Hazardous Chemicals Chemical Footprint Project - Chemicals of High Concern List Australian Inventory of Industrial Chemicals (AIIC) National Inventory Status

National Inventory	Status
Australia - AIIC / Australia Non-Industrial Use	Yes
Canada - DSL	Yes
Canada - NDSL	No (xylene; toluene; naphtha petroleum, light aromatic solvent; n-butyl acetate; ethyl acetate; acetone; methyl ethyl ketone; styrene; ethylbenzene; methyl isobutyl ketone; 1,3,5-trimethyl benzene; heptane; n-hexane)
China - IECSC	Yes
Europe - EINEC / ELINCS / NLP	Yes
Japan - ENCS	Yes
Korea - KECI	Yes

National Inventory	Status
New Zealand - NZIoC	Yes
Philippines - PICCS	Yes
USA - TSCA	Yes
Taiwan - TCSI	Yes
Mexico - INSQ	Yes
Vietnam - NCI	Yes
Russia - ARIPS	Yes
Legend:	Yes = All CAS declared ingredients are on the inventory No = One or more of the CAS listed ingredients are not on the inventory and are not exempt from listing(see specific ingredients in brackets)

## **SECTION 16 Other information**

Revision Date	23/03/2021
Initial Date	23/03/2021

#### SDS Version Summary

Version	Issue Date	Sections Updated
2.1.1.1	23/03/2021	Acute Health (eye), Acute Health (inhaled), Acute Health (skin), Acute Health (swallowed), Advice to Doctor, Appearance, Chronic Health, Classification, Disposal, Engineering Control, Environmental, Fire Fighter (extinguishing media), Fire Fighter (fire/explosion hazard), Fire Fighter (fire fighting), Fire Fighter (fire incompatibility), First Aid (eye), First Aid (inhaled), First Aid (skin), First Aid (swallowed), Handling Procedure, Instability Condition, Personal Protection (other), Personal Protection (Respirator), Personal Protection (eye), Personal Protection (hands/feet), Physical Properties, Spills (major), Spills (minor), Storage incompatibility), Storage (storage requirement), Storage (suitable container), Transport, Transport Information, Name

#### Other information

Classification of the preparation and its individual components has drawn on official and authoritative sources as well as independent review by the Chemwatch Classification committee using available literature references.

The SDS is a Hazard Communication tool and should be used to assist in the Risk Assessment. Many factors determine whether the reported Hazards are Risks in the workplace or other settings. Risks may be determined by reference to Exposures Scenarios. Scale of use, frequency of use and current or available engineering controls must be considered.

#### **Definitions and abbreviations**

PC-TWA: Permissible Concentration-Time Weighted Average

- PC-STEL: Permissible Concentration-Short Term Exposure Limit
- IARC: International Agency for Research on Cancer
- ACGIH: American Conference of Governmental Industrial Hygienists
- STEL: Short Term Exposure Limit
- TEEL: Temporary Emergency Exposure Limit。
- IDLH: Immediately Dangerous to Life or Health Concentrations
- OSF: Odour Safety Factor
- NOAEL :No Observed Adverse Effect Level LOAEL: Lowest Observed Adverse Effect Level
- TLV: Threshold Limit Value
- LOD: Limit Of Detection
- OTV: Odour Threshold Value
- BCF: BioConcentration Factors
- BEI: Biological Exposure Index
- ______.
- This document is copyright.

Apart from any fair dealing for the purposes of private study, research, review or criticism, as permitted under the Copyright Act, no part may be reproduced by any process without written permission from CHEMWATCH.

TEL (+61 3) 9572 4700.