

TC-1630 PART B BARNES PRODUCTS PTY LTD

Chemwatch: 6010-90 Version No: 8.1

Chemwatch Hazard Alert Code: 2

Issue Date: 16/03/2023 Print Date: 19/06/2023 S.GHS.NZL.EN.E

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

Safety Data Sheet according to the Health and Safety at Work (Hazardous Substances) Regulations 2017

Product Identifier	
Product name	TC-1630 PART B
Chemical Name	Not Applicable
Synonyms	ULTRACAST PART B
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	Casting resin.
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Details of the manufacturer or supplier of the safety data sheet

Registered company name	BARNES PRODUCTS PTY LTD	
Address	5 GREENHILLS AVE MOOREBANK NSW 2170 Australia	
Telephone	Barnes Australia +612 9793 7555 Mon-Fri 8am-4:30pm	
Fax	Barnes Australia +612 9793 7091	
Website	www.barnesnz.co.nz	
Email	sales@barnes.com.au	

Emergency telephone number

Association / Organisation	New Zealand Poisons Information Centre	
Emergency telephone numbers	rnes NZ +649 9731 816 - Monday-Thursday 9am-5pm Friday 9am-4.30pm	
Other emergency telephone numbers	New Zealand Poisons Information Centre 0800 764 766 After Hours	

SECTION 2 Hazards identification

Classification of the substance or mixture		
Classification ^[1]	Flammable Liquids Category 4, Skin Corrosion/Irritation Category 2, Sensitisation (Skin) Category 1, Serious Eye Damage/Eye Irritation Category 2, Acute Toxicity (Inhalation) Category 2, Sensitisation (Respiratory) Category 1, Reproductive Toxicity Category 2, Specific Target Organ Toxicity - Repeated Exposure Category 1, Hazardous to the Aquatic Environment Long-Term Hazard Category 3	
Legend:	1. Classified by Chemwatch; 2. Classification drawn from CCID EPA NZ; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI	
Determined by Chemwatch using GHS/HSNO criteria	3 11) 6 1B (inhalation) 6 3A 6 4A 6 5A (respiratory) 6 5B (contact) 6 8B 6 9A 9 1C	

Label elements



Signal word

Danger

Hazard statement(s)

H227	Combustible liquid.
H315	Causes skin irritation.
H317	May cause an allergic skin reaction.
H319	Causes serious eye irritation.
H330	Fatal if inhaled.

H334	May cause allergy or asthma symptoms or breathing difficulties if inhaled.	
H361	S1 Suspected of damaging fertility or the unborn child.	
H372	Causes 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	Keep away from heat, hot surfaces, sparks, open flames and other ignition sources. No smoking.
P260	Do not breathe mist/vapours/spray.
P271	Use only outdoors or in a well-ventilated area.
P280	Wear protective gloves, protective clothing, eye protection and face protection.
P284	[In case of inadequate ventilation] wear respiratory protection.
P270	Do not eat, drink or smoke when using this product.

Precautionary statement(s) Response

P304+P340	IF INHALED: Remove person to fresh air and keep comfortable for breathing.	
P308+P313	IF exposed or concerned: Get medical advice/ attention.	
P310	Immediately call a POISON CENTER/doctor/physician/first aider.	
P342+P311	If experiencing respiratory symptoms: Call a POISON CENTER/doctor/physician/first aider.	
P370+P378	In case of fire: Use alcohol resistant foam or normal protein foam to extinguish.	
P302+P352	IF ON SKIN: Wash with plenty of water.	
P305+P351+P338	IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.	

Precautionary statement(s) Storage

P403+P233	Store in a well-ventilated place. Keep container tightly closed.	
P405	P405 Store locked up.	

Precautionary statement(s) Disposal

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

SECTION 3 Composition / information on ingredients

P501

Substances

See section below for composition of Mixtures

Mixtures

CAS No	%[weight]	Name
Not Available	60-70	filler mixture, proprietary
101-68-8	10-15	4.4'-diphenylmethane diisocyanate (MDI)
9016-87-9	10-15	polymeric diphenylmethane diisocyanate
6846-50-0	5-10	2.2.4-trimethyl-1.3-pentanediol diisobutyrate
64742-88-7	<2	solvent naphtha petroleum, medium aliphatic
64742-94-5	<2	solvent naphtha petroleum, heavy aromatic
91-20-3	<0.2	naphthalene
Not Available	balance	Ingredients determined not to be hazardous
Legend:	 Classified by Chernwatch; 2. Classification drawn from CCID EPA NZ; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; Classification drawn from C&L * EU IOELVs available 	

SECTION 4 First aid measures

Description of first aid measures		
Eye Contact	 If this product comes in contact with the eyes: 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 contact occurs: Immediately remove all contaminated clothing, including footwear. Flush skin and hair with running water (and soap if available). Seek medical attention in event of irritation. 	
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. Following uptake by inhalation, move person to an area free from risk of further exposure. Oxygen or artificial respiration should be administered as needed. Asthmatic-type symptoms may develop and may be immediate or delayed up to several hours. Treatment is essentially symptomatic. A physician should be consulted.
Ingestion	 IF SWALLOWED, REFER FOR MEDICAL ATTENTION, WHERE POSSIBLE, WITHOUT DELAY. For advice, contact a Poisons Information Centre or a doctor. Urgent hospital treatment is likely to be needed. In the mean time, qualified first-aid personnel should treat the patient following observation and employing supportive measures as indicated by the patient's condition. If the services of a medical officer or medical doctor are readily available, the patient should be placed in his/her care and a copy of the SDS should be provided. Further action will be the responsibility of the medical specialist. If medical attention is not available on the worksite or surroundings send the patient to a hospital together with a copy of the SDS. Where medical attention is not immediately available or where the patient is more than 15 minutes from a hospital or unless instructed otherwise: INDUCE vomiting with fingers down the back of the throat, ONLY IF CONSCIOUS. Lean patient forward or place on left side (head-down position, if possible) to maintain open airway and prevent aspiration. NOTE: Wear a protective glove when inducing vomiting by mechanical means.

Indication of any immediate medical attention and special treatment needed

for naphthalene intoxication: Naphthalene requires hepatic and microsomal activation prior to the production of toxic effects. Liver microsomes catalyse the initial synthesis of the reactive 1,2-epoxide intermediate which is subsequently oxidised to naphthalene dihydrodiol and alpha-naphthol. The 2-naphthoquinones are thought to produce haemolysis, the 1,2-naphthoquinones are thought to be responsible for producing cataracts in rabbits, and the glutathione-adducts of naphthalene-1,2-oxide are probably responsible for pulmonary toxicity. Suggested treatment regime:

- Induce emesis and/or perform gastric lavage with large amounts of warm water where oral poisoning is suspected.
- Instill a saline cathartic such as magnesium or sodium sulfate in water (15 to 30g).
- Demulcents such as milk, egg white, gelatin, or other protein solutions may be useful after the stomach is emptied but oils should be avoided because they promote absorption.
 If eyes/skin contaminated, flush with warm water followed by the application of a bland ointment.
- Severe anaemia, due to haemolysis, may require small repeated blood transfusions, preferably with red cells from a non-sensitive individual.
- Where intravascular haemolysis, with haemoglobinuria occurs, protect the kidneys by promoting a brisk flow of dilute urine with, for example, an osmotic diuretic such as mannitol.
- It may be useful to alkalinise the urine with small amounts of sodium bicarbonate but many researchers doubt whether this prevents blockage of the renal tubules.
- Use supportive measures in the case of acute renal failure. GOSSELIN, SMITH HODGE: Clinical Toxicology of Commercial Products, 5th Ed.

For sub-chronic and chronic exposures to isocyanates:

- This material may be a potent pulmonary sensitiser which causes bronchospasm even in patients without prior airway hyperreactivity.
- Clinical symptoms of exposure involve mucosal irritation of respiratory and gastrointestinal tracts.
- + Conjunctival irritation, skin inflammation (erythema, pain vesiculation) and gastrointestinal disturbances occur soon after exposure.
- Pulmonary symptoms include cough, burning, substernal pain and dyspnoea.
- Some cross-sensitivity occurs between different isocyanates.
- Noncardiogenic pulmonary oedema and bronchospasm are the most serious consequences of exposure. Markedly symptomatic patients should receive oxygen, ventilatory support and an intravenous line.
- Treatment for asthma includes inhaled sympathomimetics (epinephrine [adrenalin], terbutaline) and steroids.
- Activated charcoal (1 g/kg) and a cathartic (sorbitol, magnesium citrate) may be useful for ingestion.
- Mydriatics, systemic analgesics and topical antibiotics (Sulamyd) may be used for corneal abrasions.
- There is no effective therapy for sensitised workers.

[Ellenhorn and Barceloux; Medical Toxicology]

NOTE: Isocyanates cause airway restriction in naive individuals with the degree of response dependant on the concentration and duration of exposure. They induce smooth muscle contraction which leads to bronchoconstrictive episodes. Acute changes in lung function, such as decreased FEV1, may not represent sensitivity.

[Karol & Jin, Frontiers in Molecular Toxicology, pp 56-61, 1992]

Personnel who work with isocyanates, isocyanate prepolymers or polyisocyanates should have a pre-placement medical examination and periodic examinations thereafter, including a pulmonary function test. Anyone with a medical history of chronic respiratory disease, asthmatic or bronchial attacks, indications of allergic responses, recurrent eczema or sensitisation conditions of the skin should not handle or work with isocyanates. Anyone who develops chronic respiratory distress when working with isocyanates should be removed from exposure and examined by a physician. Further exposure must be avoided if a sensitivity to isocyanates or polyisocyanates has developed.

SECTION 5 Firefighting measures

Extinguishing media

Flooding quantities of water only.

- Foam.
- Dry chemical powder.
- BCF (where regulations permit).
- Carbon dioxide.
- Water spray or fog Large fires only.

Fire Incompatibility

Special hazards arising from the substrate or mixture

Advice for firefighters	
Fire Fighting	 Alert Fire Brigade and tell them location and nature of hazard. Wear full body protective clothing with breathing apparatus. Prevent, by any means available, spillage from entering drains or water course. Use water delivered as a fine spray to control fire and cool adjacent area. Avoid spraying water onto liquid pools. DO NOT approach containers suspected to be hot. Cool fire exposed containers with water spray from a protected location.
Fire/Explosion Hazard	 Combustible. Moderate fire hazard when exposed to heat or flame. When heated to high temperatures decomposes rapidly generating vapour which pressures and may then rupture containers with release of flammable and highly toxic isocyanate vapour.

Avoid contamination with oxidising agents i.e. nitrates, oxidising acids, chlorine bleaches, pool chlorine etc, as ignition may result

 Burns with acrid black smoke and poisonous fumes. Due to reaction with water producing CO2-gas, a hazardous build-up of pressure could result if contaminated containers are re-sealed. Combustion yields traces of highly toxic hydrogen cyanide HCN, plus toxic nitrogen oxides NOx and carbon monoxide.
Combustion products include:
carbon dioxide (CO2)
isocyanates
and minor amounts of
hydrogen cyanide
nitrogen oxides (NOx)
other pyrolysis products typical of burning organic material.
May emit corrosive fumes.
When heated at high temperatures many isocyanates decompose rapidly generating a vapour which pressurises containers, possibly to the point
of rupture. Release of toxic and/or flammable isocyanate vapours may then occur

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. Contain and absorb spill with sand, earth, inert material or vermiculite. Wipe up. Place in a suitable, labelled container for waste disposal.
Major Spills	 Liquid Isocyanates and high isocyanate vapour concentrations will penetrate seals on self contained breathing apparatus - SCBA should be used inside encapsulating suit where this exposure may occur. For isocyanate spills of less than 40 litres (2 m2): Evacuate area from everybody not dealing with the emergency, keep them upwind and prevent further access, remove ignition sources and, if inside building, ventilate area as well as possible. Notify supervision and others as necessary. Put on personal protective equipment (suitable respiratory protection, face and eye protection, protective suit, gloves and impermeable boots). Control source of leakage (where applicable). Dike the spill to prevent spreading and to contain additions of decontaminating solution. Prevent the material from entering drains. Estimate spill pool volume or area. Avoid contamination with water, alkalies and detergent solutions. Material reacts with water and generates gas, pressurises containers with even drum rupture resulting. Do NOT reseal container if contamination is suspected. Open all containers with care. Moderate hazard. Clear area of personnel and move upwind. Alert Fire Brigade and tell them location and nature of hazard. Wear breathing apparatus plus protective gloves. Prevent, by any means available, spillage from entering drains or water course. No smoking, naked lights or ignition sources. Increase ventilation.

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

SECTION 7 Handling and storage

ecautions for safe handling	
Safe handling	 DO NOT allow clothing wet with material to stay in contact with skin 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. DO NOT enter confined spaces until atmosphere has been checked. Avoid smoking, naked lights or ignition sources. Avoid contact with incompatible materials.
Other information	Consider storage under inert gas. for commercial quantities of isocyanates: • Isocyanates should be stored in adequately bunded areas. Nothing else should be kept within the same bunding. Pre-polymers need not be segregated. Drums of isocyanates should be stored under cover, out of direct sunlight, protected from rain, protected from physical damage and well away from moisture, acids and alkalis. • Where isocyanates are stored at elevated temperatures to prevent solidifying, adequate controls should be installed to prevent the high temperatures and precautions against fire should be taken. • Where stored in tanks, the more reactive isocyanates should be blanketed with a non-reactive gas such as nitrogen and equipped with absorptive type breather valve (to prevent vapour emissions) • Transfer systems for isocyanates in bulk storage should be fully enclosed and use pump or vacuum systems. • Store in original containers. • Keep containers securely sealed. • No smoking, naked lights or ignition sources. • Store in a cool, dry, well-ventilated area. • Store in a cool, dry, well-ventilated area.
	Continue

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Conditions for safe storage, in	 Protect containers against physical damage and check regularly for leaks. Observe manufacturer's storage and handling recommendations contained within this SDS.
Suitable container	 Metal can or drum Packaging as recommended by manufacturer. Check all containers are clearly labelled and free from leaks.
Storage incompatibility	 Avoid reaction with water, alcohols and detergent solutions. Isocyanates are electrophiles, and as such they are reactive toward a variety of nucleophiles including alcohols, amines, and even water. Upon treatment with an alcohol, an isocyanate forms a urethane linkage. If a di-isocyanate is treated with a compound containing two or more hydroxyl groups, such as a diol or a polyol, polymer chains are formed, which are known as polyurethanes. Reaction between a di-isocyanate and a compound containing two or more amine groups, produces long polymer chains known as polyurethanes. Reaction between a di-isocyanate and a compound containing two or more amine groups, produces long polymer chains known as polyurethanes. Reaction between a di-isocyanate and a compound containing two or more amine groups, produces long polymer chains known as polyurethanes. Reaction between a di-isocyanate and a compound containing two or more amine groups, produces long polymer chains known as polyurethanes. Reaction between a di-isocyanate and a compound containing two or more amine groups, produces long polymer chains known as polyurethanes. Reaction between a di-isocyanates and thioisocyanates are incompatible with many classes of compounds, reacting exothermically to release toxic gases. Reactions with amines, strong bases, aldehydes, alcohols, alkali metals, ketones, mercaptans, strong oxidisers, hydrides, phenols, and peroxides can cause vigorous releases of heat. A range of exothermic decomposition energies for isocyanates is given as 20-30 kJ/mol. The relationship between energy of decomposition and processing hazards has been the subject of discussion; it is suggested that values of energy released per unit of mass, rather than on a molar basis (J/g) be used in the assessment. For example, in "open vessel processes" (with man-hole size openings, in an industrial setting), substances with exothermic decomposition energy exceeds 150 J/g. BRETHERICK: Handbook of Reac

NOTE: May develop pressure in containers; open carefully. Vent periodically.

SECTION 8 Exposure controls / personal protection

Control parameters

Occupational Exposure Limits (OEL)

INGREDIENT DATA

Source	Ingredient	Material name	TWA	STEL	Peak	Notes
New Zealand Workplace Exposure Standards (WES)	4,4'-diphenylmethane diisocyanate (MDI)	Diphenylmethane diisocyanate	0.02 mg/m3	0.07 mg/m3	Not Available	(dsen) - Dermal sensitiser (rsen) - Respiratory sensitiser (ifv) - The Inhalable Fraction and Vapour (ifv) notation is used when a material exerts sufficient vapour pressure such that it may be present in both particle and vapour phases, with each contributing to a significant portion of exposure
New Zealand Workplace Exposure Standards (WES)	4,4'-diphenylmethane diisocyanate (MDI)	MDI	0.02 mg/m3	0.07 mg/m3	Not Available	(dsen) - Dermal sensitiser (rsen) - Respiratory sensitiser (ifv) - The Inhalable Fraction and Vapour (ifv) notation is used when a material exerts sufficient vapour pressure such that it may be present in both particle and vapour phases, with each contributing to a significant portion of exposure
New Zealand Workplace Exposure Standards (WES)	4,4'-diphenylmethane diisocyanate (MDI)	Methylene bisphenyl isocyanate	0.02 mg/m3	0.07 mg/m3	Not Available	(dsen) - Dermal sensitiser (rsen) - Respiratory sensitiser (ifv) - The Inhalable Fraction and Vapour (ifv) notation is used when a material exerts sufficient vapour pressure such that it may be present in both particle and vapour phases, with each contributing to a significant portion of exposure
New Zealand Workplace Exposure Standards (WES)	polymeric diphenylmethane diisocyanate	lsocyanates, all, (as -NCO)	0.02 mg/m3	0.07 mg/m3	Not Available	(dsen) - Dermal sensitiser (rsen) - Respiratory sensitiser (ifv) - The Inhalable Fraction and Vapour (ifv) notation is used when a material exerts sufficient vapour pressure such that it may be present in both particle and vapour phases, with each contributing to a significant portion of exposure
New Zealand Workplace Exposure Standards (WES)	solvent naphtha petroleum, medium aliphatic	Oil mist, mineral	5 mg/m3	10 mg/m3	Not Available	(om) - Sampled by a method that does not collect vapour
New Zealand Workplace Exposure Standards (WES)	naphthalene	Naphthalene	0.5 ppm / 2.6 mg/m3	10 mg/m3 / 2 ppm	Not Available	carcinogen category 2 - Suspected human carcinogen (skin) - Skin absorption

Emergency Limits

Ingredient	TEEL-1	TEEL-2	TEEL-3
4,4'-diphenylmethane diisocyanate (MDI)	0.45 mg/m3	Not Available	Not Available
4,4'-diphenylmethane diisocyanate (MDI)	29 mg/m3	40 mg/m3	240 mg/m3
polymeric diphenylmethane diisocyanate	0.15 mg/m3	3.6 mg/m3	22 mg/m3
solvent naphtha petroleum, medium aliphatic	1,200 mg/m3	6,700 mg/m3	40,000 mg/m3
naphthalene	15 ppm	83 ppm	500 ppm

Ingredient	Original IDLH	Revised IDLH	
4,4'-diphenylmethane diisocyanate (MDI)	75 mg/m3	Not Available	
polymeric diphenylmethane diisocyanate	Not Available	Not Available	
2,2,4-trimethyl-1,3-pentanediol diisobutyrate	Not Available	Not Available	
solvent naphtha petroleum, medium aliphatic	2,500 mg/m3	Not Available	
solvent naphtha petroleum, heavy aromatic	Not Available	Not Available	
naphthalene	250 ppm	Not Available	
Occupational Exposure Banding			
Ingredient	Occupational Exposure Band Rating	Occupational Exposure Band Limit	
2,2,4-trimethyl-1,3-pentanediol diisobutyrate	E	≤ 0.1 ppm	
Notes:	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.		

- All processes in which isocyanates are used should be enclosed wherever possible. + Total enclosure, accompanied by good general ventilation, should be used to keep atmospheric concentrations below the relevant exposure standards. F If total enclosure of the process is not feasible, local exhaust ventilation may be necessary. Local exhaust ventilation is essential where lower molecular weight isocyanates (such as TDI or HDI) is used or where isocyanate or polyurethane is sprayed. • Where other isocyanates or pre-polymers are used and aerosol formation cannot occur, local exhaust ventilation may not be necessary if the atmospheric concentration can be kept below the relevant exposure standards. Appropriate engineering Where local exhaust ventilation is installed, exhaust vapours should not be vented to the exterior in such a manner as to create a hazard. controls 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. Ventilation can remove or dilute an air contaminant if designed properly. The design of a ventilation system must match the particular process and chemical or contaminant in use. Employers may need to use multiple types of controls to prevent employee overexposure. Individual protection measures, such as personal protective equipment Safety glasses with side shields. Chemical goggles. [AS/NZS 1337.1, EN166 or national equivalent] Contact lenses may pose a special hazard; soft contact lenses may absorb and concentrate irritants. A written policy document, describing the wearing of lenses or restrictions on use, should be created for each workplace or task. This should include a review of lens absorption Eye and face protection and adsorption for the class of chemicals in use and an account of injury experience. Medical and first-aid personnel should be trained in their removal and suitable equipment should be readily available. In the event of chemical exposure, begin eye irrigation immediately and remove contact lens as soon as practicable. Skin protection See Hand protection below NOTE: The material may produce skin sensitisation in predisposed individuals. Care must be taken, when removing gloves and other protective equipment, to avoid all possible skin contact. Contaminated leather items, such as shoes, belts and watch-bands should be removed and destroyed. 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. Hands/feet protection 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. Gloves must only be worn on clean hands. After using gloves, hands should be washed and dried thoroughly. Application of a non-perfumed moisturiser is recommended. Isocyanate resistant materials include Teflon, Viton, nitrile rubber and some PVA gloves. Protective gloves and overalls should be worn as specified in the appropriate national standard. Contaminated garments should be removed promptly and should not be re-used until they have been decontaminated. NOTE: Natural rubber, neoprene, PVC can be affected by isocyanates Body protection See Other protection below All employees working with isocyanates must be informed of the hazards from exposure to the contaminant and the precautions necessary to prevent damage to their health. They should be made aware of the need to carry out their work so that as little contamination as possible is produced, and of the importance of the proper use of all safeguards against exposure to themselves and their fellow workers. Adequate training, both in the proper execution of the task and in the use of all associated engineering controls, as well as of any personal protective equipment, is essential. Employees exposed to contamination hazards should be educated in the need for, and proper use of, facilities, clothing and equipment and Other protection thereby maintain a high standard of personal cleanliness. Special attention should be given to ensuring that all personnel understand instructions, especially newly recruited employees and those with local-language difficulties, where they are known. Overalls P.V.C apron. Barrier cream.
 - Skin cleansing cream

Eye wash unit.

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:

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Material	СРІ
PE/EVAL/PE	С
TEFLON	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.

Respiratory protection

Type A-P 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	A-AUS P2	-	A-PAPR-AUS / Class 1 P2
up to 50 x ES	-	A-AUS / Class 1 P2	-
up to 100 x ES	-	A-2 P2	A-PAPR-2 P2 ^

^ - 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

For spraying or operations which might generate aerosols:

Full face respirator with supplied air.

- In certain circumstances, personal protection of the individual employee is necessary. Personal protective devices should be regarded as being supplementary to substitution and engineering control and should not be used in preference to them as they do nothing to eliminate the hazard.
- However, in some situations, minimising exposure to isocyanates by enclosure and ventilation is not possible, and occupational exposure standards may be exceeded, particularly during on-site mixing of paints, spray-painting, foaming and maintenance of machine and ventilation systems. In these situations, air-line respirators or self-contained breathing apparatus complying with the appropriate nationals standard must be used.
- Organic vapour respirators with particulate pre- filters and powered, air-purifying respirators are NOT suitable.
- Personal protective equipment must be appropriately selected, individually fitted and workers trained in their correct use and maintenance. Personal protective equipment must be regularly checked and maintained to ensure that the worker is being protected.
- Air- line respirators or self-contained breathing apparatus complying with the appropriate national standard should be used during the clean-up of spills and the repair or clean-up of contaminated equipment and similar situations which cause emergency exposures to hazardous atmospheric concentrations of isocyanate.

SECTION 9 Physical and chemical properties

Information on basic physical and chemical properties

Appearance	White viscous liquid with aromatic odour odour; partly mixes with water.		
Physical state	Liquid	Relative density (Water = 1)	1.730
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 (°C)	Not Available
Melting point / freezing point (°C)	Not Available	Viscosity (cSt)	Not Available
Initial boiling point and boiling range (°C)	Not Available	Molecular weight (g/mol)	Not Applicable
Flash point (°C)	73 CC	Taste	Not Available
Evaporation rate	Not Available	Explosive properties	Not Available
Flammability	Combustible.	Oxidising properties	Not Available

Upper Explosive Limit (%)	Not Available	Surface Tension (dyn/cm or mN/m)	Not Available
Lower Explosive Limit (%)	Not Available	Volatile Component (%vol)	3.9
Vapour pressure (kPa)	Not Available	Gas group	Not Available
Solubility in water	Reacts	pH as a solution (1%)	Not Available
Vapour density (Air = 1)	Not Available	VOC g/L	67.500

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

Inhaled	Inhalation of vapours or aerosols (mists, fumes), generated by the materi There is some evidence to suggest that the material can cause respirator cause further lung damage. The vapour/mist may be highly irritating to the upper respiratory tract and pulmonary oedema. Possible neurological symptoms arising from isocyan neurosis, depression and paranoia. Gastrointestinal disturbances are cha produce asthmatic reactions ranging from minor breathing difficulties to s or may develop without warning for several hours after exposure. Sensiti work in situations allowing exposure to this material. Continued exposure impairment. Inhalation hazard is increased at higher temperatures. Inhalation of naphthalene vapour is linked with headache, loss of appetite long term exposure may cause excessive weakness and increased saliva- to the skin, liver and lungs.	y irritation in some persons. The body's response to such irritation can lungs; the response may be severe enough to produce bronchitis and nate exposure include headache, insomnia, euphoria, ataxia, anxiety aracterised by nausea and vomiting. Pulmonary sensitisation may evere allergic attacks; this may occur following a single acute exposure zed people can react to very low doses, and should not be allowed to of sensitised persons may lead to possible long term respiratory e, nausea, damage to the eyes and kidneys. According to animal testing,
Ingestion	Accidental ingestion of the material may be seriously damaging to the he than 40 gram may be fatal. Ingestion of naphthalene and related compounds may produce abdomina listlessness, confusion, and in severe poisonings, coma with or without co painful urination, and the passage of brown or black urine with or without	al cramps with nausea, vomiting, diarrhoea, headache, profuse sweating, provulsions. Irritation of the bladder may also occur, producing urgency,
Skin Contact	Skin contact with the material may damage the health of the individual; systemic effects may result following absorption. There is some evidence to suggest that this material can cause inflammation of the skin on contact in some persons. Workers sensitised to naphthalene and related compounds show an inflammation of the skin with scaling and reddening. Some individuals show an allergic reaction. 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.	
Eye	There is some evidence to suggest that this material can cause eye irritation and damage in some persons. Long term exposure to naphthalene has produced clouding of the lens (cataracts) in workers.	
Chronic	There has been concern that this material can cause cancer or mutations Inhaling this product is more likely to cause a sensitisation reaction in sor Skin contact with the material is more likely to cause a sensitisation react Harmful: danger of serious damage to health by prolonged exposure thro This material can cause serious damage if one is exposed to it for long p produce severe defects. Substance accumulation, in the human body, may occur and may cause Persons with a history of asthma or other respiratory problems or are kno handling of isocyanates. The chemistry of reaction of isocyanates, as evidenced by MDI, in biologi doses to the mouth, reactions will commence at once with biological mac tract prior to reaching the stomach. Reaction products will be a variety of proteins and cell components. This is corroborated by the results from an MDI inhalation study. Followin was excreted in faeces. The faecal excretion in these animals was consisi ingestion of deposited material from the nasopharangeal region via the m radioactivity was tentatively identified as mixed molecular weight polyures Animal testing indicates that inhalation of naphthalene may increase the inflammation. Isocyanate vapours are irritating to the airways and can cause their inflam consciousness and fluid in the lungs. Nervous system symptoms that ma anxiety, depression and paranoia.	ne persons compared to the general population. ion in some persons compared to the general population. ugh inhalation. eriods. It can be assumed that it contains a substance which can some concern following repeated or long-term occupational exposure. where the sensitised, should not be engaged in any work involving the ical milieu is such that in the event of a true exposure of small MDI romolecules in the buccal region and will continue along the digestive polyureas and macromolecular conjugates with for example mucus, g an inhalation exposure of rats to radiolabelled MDI, 79% of the dose dered entirely due to ingestion of radioactivity from grooming and nuccoliary escalator, i.e. not following systemic absorption. The faecal as derived from MDI. incidence of respiratory tumours and may aggravate chronic nmation, with wheezing, gasping, severe distress, even loss of
	τοχιατγ	
TC-1630 PART B		IRRITATION
	Not Available	Not Available

	ΤΟΧΙΟΙΤΥ	IRRITATION
	Dermal (rabbit) LD50: >6200 mg/kg ^[2]	Eye: no adverse effect observed (not irritating) ^[1]
4,4'-diphenylmethane diisocyanate (MDI)	Inhalation(Rat) LC50: 0.368 mg/L4h ^[1]	Skin (rabbit): 500 mg /24 hours Dermal Sensitiser *Respiratory Sensitiser (g.pig) *[* = Bayer CCINFO 2133615]
	Oral (Mouse) LD50; 2200 mg/kg ^[2]	Skin: adverse effect observed (irritating) ^[1]
	ΤΟΧΙCITY	IRRITATION
	Dermal (rabbit) LD50: >9400 mg/kg ^[2]	Eye (rabbit): 100 mg - mild
polymeric diphenylmethane diisocyanate	Inhalation(Rat) LC50: 0.49 mg/L4h ^[2]	
	Oral (Rat) LD50: 43000 mg/kg ^[2]	
	ΤΟΧΙΟΙΤΥ	IRRITATION
	Dermal (rabbit) LD50: >2000 mg/kg ^[1]	Eye (rabbit): very slight** **[Eastman] *[Patty]
2,2,4-trimethyl-1,3-pentanediol diisobutyrate	Oral (Rat) LD50: >2000 mg/kg ^[1]	Eye: no adverse effect observed (not irritating) ^[1]
unsobutyrate		Skin (guinea pig): 5000mg/kg-mild
		Skin: no adverse effect observed (not irritating) ^[1]
	ΤΟΧΙΟΙΤΥ	IRRITATION
solvent naphtha petroleum,	Dermal (rabbit) LD50: >2000 mg/kg ^[2]	Eye: no adverse effect observed (not irritating) ^[1]
medium aliphatic	Inhalation(Rat) LC50: >4.3 mg/l4h ^[1]	Skin: no adverse effect observed (not irritating) ^[1]
	Oral (Rat) LD50: >5000 mg/kg ^[2]	
	ΤΟΧΙΟΙΤΥ	IRRITATION
colvent nanhtha natroloum	Dermal (rabbit) LD50: >2000 mg/kg ^[2]	Eye (rabbit): Irritating [PETROFIN]
solvent naphtha petroleum, heavy aromatic	Inhalation(Rat) LC50: >0.003 mg/L4h ^[1]	Eye: no adverse effect observed (not irritating) ^[1]
	Oral (Rat) LD50: >2000 mg/kg ^[1]	Skin: adverse effect observed (irritating) ^[1]
	ΤΟΧΙΟΙΤΥ	IRRITATION
	dermal (rat) LD50: >2500 mg/kg ^[2]	Eye (rabbit): 100 mg - mild
naphthalene	Inhalation(Rat) LC50: >0.4 mg/l4h ^[1]	Skin (rabbit):495 mg (open) - mild
	Oral (Rat) LD50: 490 mg/kg ^[2]	
Legend:	1. Value obtained from Europe ECHA Registered Substar specified data extracted from RTECS - Register of Toxic I	nces - Acute toxicity 2. Value obtained from manufacturer's SDS. Unless otherwise Effect of chemical Substances
4,4'-DIPHENYLMETHANE	Inhalation (human) TCLo: 0.13 ppm/30 mins Eye (rabbit):	0 10 mg moderate
DIISOCYANATE (MDI) POLYMERIC		
DIPHENYLMETHANE DIISOCYANATE	product	
2,2,4-TRIMETHYL- 1,3-PENTANEDIOL DIISOBUTYRATE	NOAEL oral (rat), 103 days = 1% in diet *** NOEL oral (dog), 90 days = 1% in diet *** Mutagenicity/Genotoxicity Data: *** Chromosomal aberration assay: Negative (+/- activation) CHO/HGPRT assay: Negative (+/- activation) Salmonella-E.coli reverse mutation assay (Ames test): Negative (+/- activation) *,**,*** Various suppliers MSDS Sensitization Species:Guinea pig: Result: sensitizing Effects on foetal development: Species: Rabbit Application Route: Oral Developmental Toxicity: NOAEL: 300 mg/kg body weight Reproductive toxicity;Assessment: Some evidence of adverse effects on development, based on animal experiments. * Eastman Benzoflex 6000 Plasticiser For 2,2,4-trimethyl-1,3-pentanediol diisobutyrate (TXIB) Laboratory testing showed that TXIB does not cause genetic toxicity. It may damage the kidneys of developing animals but only at levels that als affect the adult.	
SOLVENT NAPHTHA PETROLEUM, MEDIUM ALIPHATIC	produce conjunctivitis. For toluene: Acute toxicity: Humans exposed to high levels of toluene I from headaches to intoxication, convulsions, narcosis (sle nervous system depression, and in large doses has a nar congestion and bleeding of the lungs and kidney injury we Exposure to inhalation at a concentration of 600 parts per (a feeling of well-being), dilated pupils, convulsions and na narcosis and death. Toluene can also strip the skin of lipic	million for 8 hours resulted in the same and more serious symptoms including euphor ausea. Exposure to 10000-30000 parts per million (1-3%) has been reported to cause ls, causing skin inflammation. the adverse central nervous system effects and can damage the upper airway, the liver
NAPHTHALENE	conjunctivitis.	contact causing inflammation. Repeated or prolonged exposure to irritants may produce
	WARNING: This substance has been classified by the IAR	
4,4'-DIPHENYLMETHANE DIISOCYANATE (MDI) & POLYMERIC DIPHENYLMETHANE	eczema involves a cell-mediated (T lymphocytes) immune	group and may not be specific to this product. eczema, more rarely as urticaria or Quincke's oedema. The pathogenesis of contact e reaction of the delayed type. Other allergic skin reactions, e.g. contact urticaria, ance of the contact allergen is not simply determined by its sensitisation potential: the

TRIMETHYL- 1,3-PENTANEDIOL DIISOBUTYRATE	distributed can be a more important allergen than one with stronger sensitising potential wit clinical point of view, substances are noteworthy if they produce an allergic test reaction in	
4,4'-DIPHENYLMETHANE DIISOCYANATE (MDI) & POLYMERIC DIPHENYLMETHANE DIISOCYANATE	Exogenous allergic alveolitis is induced essentially by allergen specific immune-complexes of the IgG type; cell-mediated reactions (1 lymphocytes) may be involved. Such allergy is of the delayed type with onset up to four hours following exposure. 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 animal testing. Isocyanate vapours are irritating to the airways and can cause their inflammation, with wheezing, gasping, severe distress, even loss of consciousness and fluid in the lungs. Nervous system symptoms that may occur include headache, sleep disturbance, euphoria, inco-ordination anxiety, depression and paranoia. The material may produce moderate eye irritation leading to inflammation. Repeated or prolonged exposure to irritants may produce conjunctivitis. Aromatic and aliphatic diisocyanates may cause airway toxicity and skin sensitization. Monomers and prepolymers exhibit similar respiratory effect. Of the several members of diisocyanates tested on experimental animals by inhalation and oral exposure, some caused cancer while others produced a harmless outcome. This group of compounds has therefore been classified as cancer-causing.	
2,2,4-TRIMETHYL- 1,3-PENTANEDIOL		
DIISOBUTYRATE & SOLVENT NAPHTHA PETROLEUM, MEDIUM ALIPHATIC & NAPHTHALENE	The material may cause skin irritation after prolonged or repeated exposure and may produvesicles, scaling and thickening of the skin.	ce on contact skin redness, swelling, the production of
NAPHTHA PETROLEUM, MEDIUM ALIPHATIC &		astrointestinal tract and that the absorption of C30. With respect to the carbon chain lengths likely to paraffins. us species. In many cases, the hydrophobic ieear unchanged as in the lipoprotein particles in the gut cell. The gut cell may play a major role in red in peripheral tissues such as in the body fat stores and n-hexane, which can be metabolized to hal studies suggest high concentrations of toluene lead esting shows evidence of tumour formation. ver and kidney; these are however not considered to regarding the potential to cause mutations, including cause developmental effects such as lower birth no adverse effects on the foetus. ad to skin inflammation and may make the skin more
NAPHTHA PETROLEUM, MEDIUM ALIPHATIC & NAPHTHALENE SOLVENT NAPHTHA PETROLEUM, MEDIUM ALIPHATIC & SOLVENT NAPHTHA PETROLEUM,	 vesicles, scaling and thickening of the skin. Animal studies indicate that normal, branched and cyclic paraffins are absorbed from the gan-paraffins is inversely proportional to the carbon chain length, with little absorption above (be present in mineral oil, n-paraffins may be absorbed to a greater extent than iso- or cyclo The major classes of hydrocarbons are well absorbed into the gastrointestinal tract in variou hydrocarbons are ingested in association with fats in the diet. Some hydrocarbons may app gut lymph, but most hydrocarbons partly separate from fats and undergo metabolism in the determining the proportion of hydrocarbon that becomes available to be deposited unchang or the liver. For petroleum: This product contains benzene, which can cause acute myeloid leukaemia, compounds which are toxic to the nervous system. This product contains toluene, and anim to hearing loss. This product contains ethyl benzene and naphthalene, from which animal te Cancer-causing potential: Animal testing shows inhaling petroleum causes tumours of the live relevant in humans. Mutation-causing potential: Most studies involving gasoline have returned negative results all recent studies in living human subjects (such as in petrol service station attendants). Reproductive toxicity: Animal studies show that high concentrations of toluene (>0.1%) can weight and developmental toxicity to the nervous system of the foetus. Other studies show thum an effects: Prolonged or repeated contact may cause defatting of the skin which can le susceptible to irritation and penetration by other materials. 	astrointestinal tract and that the absorption of C30. With respect to the carbon chain lengths likely to paraffins. us species. In many cases, the hydrophobic ieear unchanged as in the lipoprotein particles in the gut cell. The gut cell may play a major role in red in peripheral tissues such as in the body fat stores and n-hexane, which can be metabolized to hal studies suggest high concentrations of toluene lead esting shows evidence of tumour formation. ver and kidney; these are however not considered to regarding the potential to cause mutations, including cause developmental effects such as lower birth no adverse effects on the foetus. ad to skin inflammation and may make the skin more
NAPHTHA PETROLEUM, MEDIUM ALIPHATIC & NAPHTHALENE SOLVENT NAPHTHA PETROLEUM, MEDIUM ALIPHATIC & SOLVENT NAPHTHA PETROLEUM, HEAVY AROMATIC	vesicles, scaling and thickening of the skin. Animal studies indicate that normal, branched and cyclic paraffins are absorbed from the gr n-paraffins is inversely proportional to the carbon chain length, with little absorption above 0 be present in mineral oil, n-paraffins may be absorbed to a greater extent than iso- or cyclo The major classes of hydrocarbons are well absorbed into the gastrointestinal tract in varion hydrocarbons are ingested in association with fats in the diet. Some hydrocarbons may app gut lymph, but most hydrocarbons partly separate from fats and undergo metabolism in the determining the proportion of hydrocarbon that becomes available to be deposited unchang or the liver. For petroleum: This product contains benzene, which can cause acute myeloid leukaemia, compounds which are toxic to the nervous system. This product contains toluene, and anim to hearing loss. This product contains ethyl benzene and naphthalene, from which animal to Cancer-causing potential: Animal testing shows inhaling petroleum causes tumours of the I be relevant in humans. Mutation-causing potential: Most studies involving gasoline have returned negative results i all recent studies in living human subjects (such as in petrol service station attendants). Reproductive toxicity: Animal studies show that high concentrations of toluene (>0.1%) can weight and developmental toxicity to the nervous system of the foetus. Other studies show Human effects: Prolonged or repeated contact may cause defatting of the skin which can le susceptible to irritation and penetration by other materials. Animal testing shows that exposure to gasoline over a lifetime can cause kidney cancer, bu	astrointestinal tract and that the absorption of C30. With respect to the carbon chain lengths likely to paraffins. us species. In many cases, the hydrophobic tear unchanged as in the lipoprotein particles in the gut cell. The gut cell may play a major role in led in peripheral tissues such as in the body fat stores and n-hexane, which can be metabolized to hal studies suggest high concentrations of toluene lead esting shows evidence of tumour formation. ver and kidney; these are however not considered to regarding the potential to cause mutations, including cause developmental effects such as lower birth no adverse effects on the foetus. ad to skin inflammation and may make the skin more t the relevance in humans is questionable.
NAPHTHA PETROLEUM, MEDIUM ALIPHATIC & NAPHTHALENE SOLVENT NAPHTHA PETROLEUM, MEDIUM ALIPHATIC & SOLVENT NAPHTHA PETROLEUM, HEAVY AROMATIC	vesicles, scaling and thickening of the skin. Animal studies indicate that normal, branched and cyclic paraffins are absorbed from the gan-paraffins is inversely proportional to the carbon chain length, with little absorption above (be present in mineral oil, n-paraffins may be absorbed to a greater extent than iso- or cyclo. The major classes of hydrocarbons are well absorbed into the gastrointestinal tract in variou hydrocarbons are ingested in association with fats in the diet. Some hydrocarbons may app gut lymph, but most hydrocarbons partly separate from fats and undergo metabolism in the determining the proportion of hydrocarbon that becomes available to be deposited unchang or the liver. For petroleum: This product contains benzene, which can cause acute myeloid leukaemia, compounds which are toxic to the nervous system. This product contains toluene, and anim to hearing loss. This product contains ethyl benzene and naphthalene, from which animal the cancer-causing potential: Animal testing shows inhaling petroleum causes tumours of the libe relevant in humans. Mutation-causing potential: Most studies involving gasoline have returned negative results in all recent studies in living human subjects (such as in petrol service station attendants). Reproductive toxicity: Animal studies show that high concentrations of toluene (>0.1%) can weight and developmental toxicity to the nervous system of the foetus. Other studies show Human effects: Prolonged or repeated contact may cause defatting of the skin which can le susceptible to irritation and penetration by other materials. Animal testing shows that exposure to gasoline over a lifetime can cause kidney cancer, but and testing shows that exposure to gasoline over a lifetime can cause kidney cancer, but and testing shows that exposure to gasoline over a lifetime can cause kidney cancer, but and testing shows that exposure to gasoline over a lifetime can cause kidney cancer, but and testing shows that exposure to gasoline over a lifetime can cause kidney cance	astrointestinal tract and that the absorption of C30. With respect to the carbon chain lengths likely to paraffins. us species. In many cases, the hydrophobic tear unchanged as in the lipoprotein particles in the gut cell. The gut cell may play a major role in yed in peripheral tissues such as in the body fat stores and n-hexane, which can be metabolized to hal studies suggest high concentrations of toluene lead esting shows evidence of tumour formation. ver and kidney; these are however not considered to regarding the potential to cause mutations, including cause developmental effects such as lower birth no adverse effects on the foetus. ad to skin inflammation and may make the skin more t the relevance in humans is questionable.
NAPHTHA PETROLEUM, MEDIUM ALIPHATIC & NAPHTHALENE SOLVENT NAPHTHA PETROLEUM, MEDIUM ALIPHATIC & SOLVENT NAPHTHA PETROLEUM, HEAVY AROMATIC Acute Toxicity Skin Irritation/Corrosion	vesicles, scaling and thickening of the skin. Animal studies indicate that normal, branched and cyclic paraffins are absorbed from the gas n-paraffins is inversely proportional to the carbon chain length, with little absorption above (be present in mineral oil, n-paraffins may be absorbed to a greater extent than iso- or cyclo The major classes of hydrocarbons are well absorbed into the gastrointestinal tract in variou hydrocarbons are ingested in association with fats in the diet. Some hydrocarbons may apg gut lymph, but most hydrocarbons partly separate from fats and undergo metabolism in the determining the proportion of hydrocarbon that becomes available to be deposited unchang or the liver. For petroleum: This product contains benzene, which can cause acute myeloid leukaemia, compounds which are toxic to the nervous system. This product contains toluene, and anim to hearing loss. This product contains thy benzene and naphthalene, from which animal the cancer-causing potential: Animal testing shows inhaling petroleum causes tumours of the libe relevant in humans. Mutation-causing potential: Most studies involving gasoline have returned negative results all recent studies in living human subjects (such as in petrol service station attendants). Reproductive toxicity: Animal studies show that high concentrations of toluene (>0.1%) can weight and developmental toxicity to the nervous system of the foetus. Other studies show Human effects: Prolonged or repeated contact may cause defatting of the skin which can lesusceptible to irritation and penetration by other materials. Animal testing shows that exposure to gasoline over a lifetime can cause kidney cancer, but the astroine of the skin which can lesusceptible to irritation and penetration by other materials. Animal testing shows that exposure to gasoline over a lifetime can cause kidney cancer, but the shime and the show that high concent can cause kidney cancer, but the shime and the show that exposure to gasoline over a lifetime can cause kidney cancer.	astrointestinal tract and that the absorption of 230. With respect to the carbon chain lengths likely to paraffins. us species. In many cases, the hydrophobic ear unchanged as in the lipoprotein particles in the gut cell. The gut cell may play a major role in ted in peripheral tissues such as in the body fat stores and n-hexane, which can be metabolized to hal studies suggest high concentrations of toluene lead esting shows evidence of tumour formation. ver and kidney; these are however not considered to regarding the potential to cause mutations, including cause developmental effects such as lower birth no adverse effects on the foetus. ad to skin inflammation and may make the skin more t the relevance in humans is questionable.

Legend:

X − Data either not available or does not fill the criteria for classification
→ Data available to make classification

SECTION 12 Ecological information

	Endpoint	Test Duration (hr)	Species	Valu	le Source
TC-1630 PART B Not Available		Not Available	Not Available	Not Ava	Not ilable Available
	Endpoint	Test Duration (hr)	Species	Value	Source
4,4'-diphenylmethane diisocyanate (MDI)	LC50	96h	Fish	95.24-134.3	7mg/l Not Availabl
	BCF	672h	Fish	61-150	7
	EC50	48h	Crustacea	>100mg/l	2
	NOEC(ECx)	504h	Crustacea	>=10mg/l	2

a share sin dia baardaa athaa a	Endpoint	Test Duration (hr)	Species	Value	Source
polymeric diphenylmethane diisocyanate	Not Available	Not Available	Not Available	Not Available	Not Availabl
	Endpoint	Test Duration (hr)	Species	Value	Source
2,2,4-trimethyl-1,3-pentanediol	BCF	1008h	Fish	0.6-0.8	7
	NOEC(ECx)	504h	Crustacea	0.7mg/l	2
diisobutyrate	LC50	96h	Fish	>1.55mg/l	2
	EC50	72h	Algae or other aquatic plants	>7.49mg/l	2
	EC50	48h	Crustacea	>1.46mg/l	1
	Endpoint	Test Duration (hr)	Species	Value	Sourc
solvent naphtha petroleum,	EC50(ECx)	48h	Crustacea	>100mg/l	1
medium aliphatic	EC50	96h	Algae or other aquatic plants	Algae or other aquatic plants 450mg/l	
	EC50	48h	Crustacea	>100mg/l	1
	Endpoint	Test Duration (hr)	Species	Value	Source
	EC50(ECx)	48h	Crustacea	0.95mg/l	1
	EC50	72h	Algae or other aquatic plants	<1mg/l	1
solvent naphtha petroleum, heavy aromatic	LC50	96h	Fish	2-5mg/l	Not Availab
	EC50	96h	Algae or other aquatic plants	11.7mg/l	2
	EC50	48h	Crustacea	0.95mg/l	1
	Endpoint	Test Duration (hr)	Species	Value	Sourc
	BCF	1344h	Fish	23-146	7
	EC50(ECx)	0.05h	Crustacea	<0.00001mg/l	4
naphthalene	EC50	72h	Algae or other aquatic plants	ca.0.4mg/l	1
	EC50	48h	Crustacea	1.09-3.4mg/l	4
	LC50	96h	Fish	0.213mg/l	4
Legend:	Ecotox databa		CHA Registered Substances - Ecotoxicological Inform C Aquatic Hazard Assessment Data 6. NITE (Japan) -		

Harmful to aquatic organisms, may cause long-term adverse effects in the aquatic environment.

Do NOT allow product to come in contact with surface waters or to intertidal areas below the mean high water mark. Do not contaminate water when cleaning equipment or disposing of equipment wash-waters.

Wastes resulting from use of the product must be disposed of on site or at approved waste sites.

For Isocyanate Monomers:

Environmental Fate: Isocyanates, (di- and polyfunctional isocyanates), are commonly used to make various polymers, such as polyurethanes. Polyurethanes find significant application in the manufacture of rigid and flexible foams. They are also used in the production of adhesives, elastomers, and coatings.

Atmospheric Fate: These substances are not expected to be removed from the air via precipitation washout or dry deposition.

Terrestrial Fate: These substances are expected to sorb strongly to soil. Migration to groundwater and surface waters is not expected to occur.

Aquatic Fate: Breakdown by water, (hydrolysis), is the primary fate mechanism for the majority of commercial isocyanate monomers, however; the low solubility of these substances will generally lessen the effectiveness of hydrolysis as a fate pathway.

For naphthalene:

Environmental Fate: Naphthalene may be reach surface water and soil through transportation in water or being carried by air. Most airborne naphthalene is in a vapour form and hence deposition is expected to be slow. A minimal amount of naphthalene emitted to the air is transported to other environmental components mostly by dry deposition. Naphthalene in surface water may volatililize into the atmosphere, depending on environmental condiditons. It remains in solution in water, with only small amounts associated with suspended material and benthic sediments. While naphthalene is readily volatilized from aerated soils, it adheres to soils with a high organic content. Adsorption to aquifer material reduces transportation of naphthalene through groundwater, and the presence of nonionic organic compounds such as tetrachloroethene may enhance sorption to materials that contain low carbon content.

Drinking Water Standards: hydrocarbon total: 10 ug/l (UK max.).

For 2,2,4-trimethyl-1,3-pentanediol diisobutyrate (TMPD)

This chemical is used as additives to plastic (plasticiser). This chemical is stable in neutral and acidic solutions, and is considered as inherently biodegradable .

Photodegradation half-life: 90.7 years

Stability in water: stable at pH 4 and pH 7 (OECD TG 111); half-life 178 days at pH9

Transport and Distribution (calculated Fugacity Level III):

In air 3.4E-10 mg/l

In water1.2E-05 mg/l

In soil 7.4E-06 mg/kg

In sediment 3.2E-03 mg/kg

Oxygen Demand Data:

ThOD: 2.40 g oxygen/g

BOD-5 and BOD-20 were not determined because the aqueous solubility of the test article was below that which is required for these tests.

For 2,2,4-trimethyl-1,3-pentanediol diisobutyrate (TXIB)

Environmental Fate:

Biodegradability: If released into water, this substance is inherently biodegraded.

Hydrolysis as a function to pH: The chemical is stable in water at pH 4 and 7. The half-life at pH 9 is 178 days.

Photodegradability (estimation): A half-life time of 90.7 years is estimated for the direct photodegradation of the chemical in water.

DO NOT discharge into sewer or waterways.

Persistence and degradability

Ingredient

Persistence: Water/Soil

Persistence: Air

Ingredient	Persistence: Water/Soil	Persistence: Air
4,4'-diphenylmethane diisocyanate (MDI)	LOW (Half-life = 1 days)	LOW (Half-life = 0.24 days)
2,2,4-trimethyl-1,3-pentanediol diisobutyrate	HIGH	HIGH
naphthalene	HIGH (Half-life = 258 days)	LOW (Half-life = 1.23 days)

Bioaccumulative potential

Bioaccumulation
LOW (BCF = 15)
LOW (BCF = 1)
LOW (BCF = 159)
HIGH (BCF = 18000)

Mobility in soil

Ingredient	Mobility
4,4'-diphenylmethane diisocyanate (MDI)	LOW (KOC = 376200)
2,2,4-trimethyl-1,3-pentanediol diisobutyrate	LOW (KOC = 607.5)
naphthalene	LOW (KOC = 1837)

SECTION 13 Disposal considerations

	Containers may still present a chemical hazard/ danger when empty.
	Return to supplier for reuse/ recycling if possible.
	Otherwise:
	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 sam product, then puncture containers, to prevent re-use, and bury at an authorised landfill.
	Where possible retain label warnings and SDS and observe all notices pertaining to the product.
	Legislation addressing waste disposal requirements may differ by country, state and/ or territory. Each user must refer to laws operating in the area. In some areas, certain wastes must be tracked.
	A Hierarchy of Controls seems to be common - the user should investigate:
	▶ Reduction
	▶ Reuse
	▶ Recycling
	 Disposal (if all else fails)
Product / Packaging disposal	This material may be recycled if unused, or if it has not been contaminated so as to make it unsuitable for its intended use. If it has been contaminated, it may be possible to reclaim the product by filtration, distillation or some other means. Shelf life considerations should also be
	applied in making decisions of this type. Note that properties of a material may change in use, and recycling or reuse may not always be appropriate.
	 DO NOT allow wash water from cleaning or process equipment to enter drains.
	 It may be necessary to collect all wash water for treatment before disposal.
	 In all cases disposal to sever may be subject to local laws and regulations and these should be considered first.
	• Where in doubt contact the responsible authority.
	DO NOT recycle spilled material.
	 Consult State Land Waste Management Authority for disposal.
	Neutralise spill material carefully and decontaminate empty containers and spill residues with 10% ammonia solution plus detergent or a
	proprietary decontaminant prior to disposal.
	DO NOT seal or stopper drums being decontaminated as CO2 gas is generated and may pressurise containers.
	Puncture containers to prevent re-use.
	Bury or incinerate residues at an approved site.

Ensure that the hazardous substance is disposed in accordance with the Hazardous Substances (Disposal) Notice 2017

Disposal Requirements

Packages that have been in direct contact with the hazardous substance must be only disposed if the hazardous substance was appropriately removed and cleaned out from the package. The package must be disposed according to the manufacturer's directions taking into account the material it is made of. Packages which hazardous content have been appropriately treated and removed may be recycled.

The hazardous substance must only be disposed if it has been treated by a method that changed the characteristics or composition of the substance and it is no longer hazardous. DO NOT deposit the hazardous substance into or onto a landfill or a sewage facility.

Burning the hazardous substance must happen under controlled conditions with no person or place exposed to

(1) a blast overpressure of more than 9 kPa; or

(2) an unsafe level of heat radiation.

The disposed hazardous substance must not come into contact with class 1 or 5 substances.

SECTION 14 Transport information

Labels Required

Marine Pollutant NO

HAZCHEM Not Applicable

Land transport (UN): NOT REGULATED FOR TRANSPORT OF DANGEROUS GOODS

Air transport (ICAO-IATA / DGR): NOT REGULATED FOR TRANSPORT OF DANGEROUS GOODS

Sea transport (IMDG-Code / GGVSee): NOT REGULATED FOR TRANSPORT OF DANGEROUS GOODS

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
4,4'-diphenylmethane diisocyanate (MDI)	Not Available
polymeric diphenylmethane diisocyanate	Not Available
2,2,4-trimethyl-1,3-pentanediol diisobutyrate	Not Available
solvent naphtha petroleum, medium aliphatic	Not Available
solvent naphtha petroleum, heavy aromatic	Not Available
naphthalene	Not Available

Transport in bulk in accordance with the IGC Code

Product name	Ship Type
4,4'-diphenylmethane diisocyanate (MDI)	Not Available
polymeric diphenylmethane diisocyanate	Not Available
2,2,4-trimethyl-1,3-pentanediol diisobutyrate	Not Available
solvent naphtha petroleum, medium aliphatic	Not Available
solvent naphtha petroleum, heavy aromatic	Not Available
naphthalene	Not Available

SECTION 15 Regulatory information

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

This substance is to be managed using the conditions specified in an applicable Group Standard

HSR Number	Group Standard	
HSR002509	Additives Process Chemicals and Raw Materials Combustible Acutely Toxic Group Standard 2020	
HSR002676	Surface Coatings and Colourants Combustible Acutely Toxic Group Standard 2020	
HSR100425	Pharmaceutical Active Ingredients Group Standard 2020	
HSR002580	Food Additives and Fragrance Materials Combustible Acutely Toxic Group Standard 2020	
HSR002594	Industrial and Institutional Cleaning Products Combustible Acutely Toxic Group Standard 2020	
HSR100757	Veterinary Medicines Limited Pack Size Finished Dose Group Standard 2020	
HSR100758	Veterinary Medicines Non dispersive Closed System Application Group Standard 2020	
HSR100756	Active Ingredients for Use in the Manufacture of Agricultural Compounds Group Standard 2020	

Please refer to Section 8 of the SDS for any applicable tolerable exposure limit or Section 12 for environmental exposure limit.

4,4'-diphenylmethane diisocyanate (MDI) is found on the following regulatory lists

 International Agency for Research on Cancer (IARC) - Agents Classified by the IARC
 New Zealand Hazardous Substances and New Organisms (HSNO) Act - Classification of Chemicals - Classification Data

 New Zealand Approved Hazardous Substances with controls
 New Zealand Hazardous Substances and New Organisms (HSNO) Act - Classification

 New Zealand Hazardous Substances and New Organisms (HSNO) Act - Classification
 New Zealand Inventory of Chemicals (NZIoC)

 New Zealand Hazardous Substances and New Organisms (HSNO) Act - Classification
 New Zealand Inventory of Chemicals (WES)

polymeric diphenylmethane diisocyanate is found on the following regulatory lists International Agency for Research on Cancer (IARC) - Agents Classified by the IARC

Monographs - Not Classified as Carcinogenic

New Zealand Approved Hazardous Substances with controls

New Zealand Hazardous Substances and New Organisms (HSNO) \mbox{Act} - Classification of Chemicals

New Zealand Hazardous Substances and New Organisms (HSNO) Act - Classification of Chemicals - Classification Data

New Zealand Inventory of Chemicals (NZIoC)

New Zealand Workplace Exposure Standards (WES)

2,2,4-trimethyl-1,3-pentanediol diisobutyrate is found on the following regulatory lists

New Zealand Hazardous Substances and New Organisms (HSNO) Act - Classification of Chemicals	New Zealand Inventory of Chemicals (NZIoC)
New Zealand Hazardous Substances and New Organisms (HSNO) Act - Classification of Chemicals - Classification Data	
solvent naphtha petroleum, medium aliphatic is found on the following regulatory lists	S
Chemical Footprint Project - Chemicals of High Concern List	New Zealand Approved Hazardous Substances with controls
International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Monographs	New Zealand Hazardous Substances and New Organisms (HSNO) Act - Classification of Chemicals
International Agency for Research on Cancer (IARC) - Agents Classified by the IARC	New Zealand Inventory of Chemicals (NZIoC)
Monographs - Group 1: Carcinogenic to humans	New Zealand Workplace Exposure Standards (WES)
International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Monographs - Not Classified as Carcinogenic	
solvent naphtha petroleum, heavy aromatic is found on the following regulatory lists	
International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Monographs - Not Classified as Carcinogenic	New Zealand Inventory of Chemicals (NZIoC)
naphthalene is found on the following regulatory lists	
Chemical Footprint Project - Chemicals of High Concern List	New Zealand Approved Hazardous Substances with controls
International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Monographs	New Zealand Hazardous Substances and New Organisms (HSNO) Act - Classification of Chemicals
International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Monographs - Group 2B: Possibly carcinogenic to humans	New Zealand Hazardous Substances and New Organisms (HSNO) Act - Classification of Chemicals - Classification Data
International WHO List of Proposed Occupational Exposure Limit (OEL) Values for Manufactured Nanomaterials (MNMS)	New Zealand Workplace Exposure Standards (WES)

Hazardous Substance Location

Subject to the Health and Safety at Work (Hazardous Substances) Regulations 2017.

Hazard Class	Quantity (Compliance Certificate)	Quantity (Compliance Certificate - Farms >4 ha)	
6.1B	250 kg or 250 L	500 kg or 500 L	

Certified Handler

Subject to Part 4 of the Health and Safety at Work (Hazardous Substances) Regulations 2017.

Class of substance	Quantities
6.1B	Any quantity

Refer Group Standards for further information

Maximum quantities of certain hazardous substances permitted on passenger service vehicles

Subject to Regulation 13.14 of the Health and Safety at Work (Hazardous Substances) Regulations 2017.

Hazard Class	Gas (aggregate water capacity in mL)	Liquid (L)	Solid (kg)	Maximum quantity per package for each classification
6.1B	120	0,1	0,5	
6.5A or 6.5B	120	1	3	
3.1C or 3.1D				10 L

Tracking Requirements

Subject to tracking according to the Health and Safety at Work (Hazardous Substances) Regulations 2017 - Refer to the regulation for more information

National Inventory Status

National Inventory	Status	
Australia - AIIC / Australia Non-Industrial Use	Yes	
Canada - DSL	Yes	
Canada - NDSL	No (4,4'-diphenylmethane diisocyanate (MDI); polymeric diphenylmethane diisocyanate; 2,2,4-trimethyl-1,3-pentanediol diisobutyrate; solvent naphtha petroleum, heavy aromatic; naphthalene)	
China - IECSC	Yes	
Europe - EINEC / ELINCS / NLP	No (polymeric diphenylmethane diisocyanate)	
Japan - ENCS	Yes	
Korea - KECI	Yes	
New Zealand - NZIoC	No (naphthalene)	
Philippines - PICCS	Yes	
USA - TSCA	Yes	
Taiwan - TCSI	Yes	
Mexico - INSQ	Yes	
Vietnam - NCI	Yes	
Russia - FBEPH	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. These ingredients may be exempt or will require registration.	

SECTION 16 Other information

Revision Date	16/03/2023
Initial Date	01/11/2006

SDS Version Summary

Version	Date of Update	Sections Updated
7.1	23/12/2022	Classification review due to GHS Revision change.
8.1	16/03/2023	Hazards identification - Classification, Identification of the substance / mixture and of the company / undertaking - Supplier 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 ES: Exposure Standard 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 AIIC: Australian Inventory of Industrial Chemicals DSL: Domestic Substances List NDSL: Non-Domestic Substances List IECSC: Inventory of Existing Chemical Substance in China EINECS: European INventory of Existing Commercial chemical Substances ELINCS: European List of Notified Chemical Substances NLP: No-Longer Polymers ENCS: Existing and New Chemical Substances Inventory KECI: Korea Existing Chemicals Inventory NZIoC: New Zealand Inventory of Chemicals PICCS: Philippine Inventory of Chemicals and Chemical Substances TSCA: Toxic Substances Control Act TCSI: Taiwan Chemical Substance Inventory INSQ: Inventario Nacional de Sustancias Químicas NCI: National Chemical Inventory FBEPH: Russian Register of Potentially Hazardous Chemical and Biological Substances This document is copyright.

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