Altex Coatings Ltd

Version No: 6.11

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

Chemwatch Hazard Alert Code: 3

Issue Date: **11/04/2024** Print Date: **11/04/2024** S.GHS.NZL.EN

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

Product Identifier	
Product name	Flowcrete Flowfast 319 Flexible Sealer Base A
Synonyms	Not Available
Proper shipping name	RESIN SOLUTION, flammable
Other means of identification	Not Available

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

Relevant identified uses	Part A of a two pack industrial flooring coating	

Details of the manufacturer or supplier of the safety data sheet

Registered company name	Itex Coatings Ltd	
Address	1 Oropi Road, Tauranga 3112 Tauranga New Zealand	
Telephone	+64 7 541 1221	
Fax	lot Available	
Website	www.altexcoatings.com	
Email	neil.debenham@altexcoatings.co.nz	

Emergency telephone number

Association / Organisation	NZ POISONS (24hr 7days)	CHEMWATCH EMERGENCY RESPONSE (24/7)
Emergency telephone numbers	0800 764766	+64 800 700 112
Other emergency telephone numbers	0800 700112	+61 3 9573 3188

Once connected and if the message is not in your preferred language then please dial 01

SECTION 2 Hazards identification

Classification of the substance or mixture

Considered a Hazardous Substance according to the criteria of the New Zealand Hazardous Substances New Organisms legislation. Classified as Dangerous Goods for transport	
purposes.	

Classification ^[1]	Flammable Liquids Category 2, Skin Corrosion/Irritation Category 2, Sensitisation (Skin) Category 1, Serious Eye Damage/Eye Irritation Category 2, Acute Toxicity (Inhalation) Category 4, 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	

Label elements

Hazard pictogram(s)	
Signal word	Danger

Hazard statement(s)

H225	Highly flammable liquid and vapour.
H315	Causes skin irritation.
H317	May cause an allergic skin reaction.
H319	Causes serious eye irritation.
H332	Harmful if inhaled.
H372	Causes damage to organs through prolonged or repeated exposure.

Harmful to aquatic life with long lasting effects.		
Precautionary statement(s) Prevention		
Keep away from heat, hot surfaces, sparks, open flames and other ignition sources. No smoking.		
Keep container tightly closed.		
Do not breathe mist/vapours/spray.		
Use only a well-ventilated area.		
Wear protective gloves, protective clothing, eye protection and face protection.		
Ground and bond container and receiving equipment.		
Use explosion-proof electrical/ventilating/lighting/intrinsically safe equipment.		
Use non-sparking tools.		
Take action to prevent static discharges.		
Do not eat, drink or smoke when using this product.		
Avoid release to the environment.		
Wash all exposed external body areas thoroughly after handling.		
Contaminated work clothing should not be allowed out of the workplace.		

Precautionary statement(s) Response

• • • • •	
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 and soap.
P305+P351+P338	IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.
P312	Call a POISON CENTER/doctor/physician/first aider/if you feel unwell.
P333+P313	If skin irritation or rash occurs: Get medical advice/attention.
P337+P313	If eye irritation persists: Get medical advice/attention.
P362+P364	Take off contaminated clothing and wash it before reuse.
P303+P361+P353	IF ON SKIN (or hair): Take off immediately all contaminated clothing. Rinse skin with water [or shower].
P304+P340	IF INHALED: Remove person to fresh air and keep comfortable for breathing.

Precautionary statement(s) Storage

P403+P235

Store in a well-ventilated place. Keep cool.

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
80-62-6	50-75	methyl methacrylate
38668-48-3	1-2.5	dipropoxy-p-toluidine
103-11-7	10-25	2-ethylhexyl acrylate
2082-81-7	1-2.5	1,4-butanediol dimethacrylate
2440-22-4	<1	2-(2'-hydroxy-5'-methylphenyl)benzotriazole
Legend:	1. Classified by Chemwatch; 2. VI; 4. Classification drawn from	Classification drawn from CCID EPA NZ; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex 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: Wash out immediately with fresh 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. Seek medical attention without delay; if pain persists or recurs seek medical attention. 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.
Ingestion	 Immediately give a glass of water. First aid is not generally required. If in doubt, contact a Poisons Information Centre or a doctor.

Indication of any immediate medical attention and special treatment needed

As in all cases of suspected poisoning, follow the ABCDEs of emergency medicine (airway, breathing, circulation, disability, exposure), then the ABCDEs of toxicology (antidotes, basics, change absorption, change distribution, change elimination). For poisons (where specific treatment regime is absent):

BASIC TREATMENT

- Establish a patent airway with suction where necessary.
- Watch for signs of respiratory insufficiency and assist ventilation as necessary.
- Administer oxygen by non-rebreather mask at 10 to 15 L/min.
- Monitor and treat, where necessary, for pulmonary oedema.
- Monitor and treat, where necessary, for shock.
- Anticipate seizures.
- DO NOT use emetics. Where ingestion is suspected rinse mouth and give up to 200 ml water (5 ml/kg recommended) for dilution where patient is able to swallow, has a strong gag reflex and does not drool.

ADVANCED TREATMENT

- Consider orotracheal or nasotracheal intubation for airway control in unconscious patient or where respiratory arrest has occurred.
- Positive-pressure ventilation using a bag-valve mask might be of use.
- Monitor and treat, where necessary, for arrhythmias.
- Start an IV D5W TKO. If signs of hypovolaemia are present use lactated Ringers solution. Fluid overload might create complications.
- Drug therapy should be considered for pulmonary oedema.
- Hypotension with signs of hypovolaemia requires the cautious administration of fluids. Fluid overload might create complications.
- Treat seizures with diazepam.
- Proparacaine hydrochloride should be used to assist eye irrigation.

BRONSTEIN, A.C. and CURRANCE, P.L.

EMERGENCY CARE FOR HAZARDOUS MATERIALS EXPOSURE: 2nd Ed. 1994 Treat symptomatically.

SECTION 5 Firefighting measures

Extinguishing media

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

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

Advice for fillengilters	
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. Consider evacuation (or protect in place). Fight fire from a safe distance, with adequate cover. If safe, switch off electrical equipment until vapour fire hazard removed. Use water delivered as a fine spray to control the 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. If safe to do so, remove containers from path of fire.
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. On combustion, may emit toxic fumes of carbon monoxide (CO). Combustion products include: carbon dioxide (CO2) nitrogen oxides (NOx) other pyrolysis products typical of burning organic material. May emit clouds of acrid smoke

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	 Environmental hazard - contain spillage. 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 small quantities with vermiculite or other absorbent material. Wipe up. Collect residues in a flammable waste container.
Major Spills	 Environmental hazard - contain spillage. 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. Prevent, by any means available, spillage from entering drains or water course. Consider evacuation (or protect in place). No smoking, naked lights or ignition sources. Increase ventilation. Stop leak if safe to do so. Water spray or fog may be used to disperse /absorb vapour. Contain spill with sand, earth or vermiculite. Use only spark-free shovels and explosion proof equipment. Collect recoverable product into labelled containers for recycling. Absorb remaining product with sand, earth or vermiculite. Collect solid residues and seal in labelled drums for disposal. Wash area and prevent runoff into drains. If contamination of drains or waterways occurs, advise emergency services.

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

SECTION 7 Handling and storage

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. 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, heat or ignition sources. When handling, DO NOT eat, drink or smoke. Vapour may ignite on pumping or pouring due to static electricity. DO NOT use plastic buckets. Earth and secure metal containers when dispensing or pouring product. Use spark-free tools when handling. Avoid contact with incompatible materials. Keep containers securely sealed. Avoid physical damage to containers. Always wash hands with soap and water after handling. Work clothes should be laundered separately. Use good occupational work practice. Observe manufacturer's storage and handling recommendations contained within this SDS. Atmosphere should be regularly checked against established exposure standards to ensure safe working conditions.
Other information	 Store below 38 deg. C. Store in original containers in approved flame-proof area. No smoking, naked lights, heat or ignition sources. DO NOT store in pits, depression, basement or areas where vapours may be trapped. Keep containers securely sealed. Store away from incompatible materials in a cool, dry well ventilated area. Protect containers against physical damage and check regularly for leaks. Observe manufacturer's storage and handling recommendations contained within this MSDS.

Conditions for safe storage, including any incompatibilities

Suitable container	 Manufactured product that requires stirring before use and having a viscosity of at least 20 cSt (25 deg. C): (i) Removable head packaging; (ii) Cans with friction closures and (iii) low pressure tubes and cartridges may be used. 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.
Storage incompatibility	 Stable under controlled storage conditions provided material contains adequate stabiliser / polymerisation inhibitor. Bulk storages may have special storage requirements WARNING: Gradual decomposition in strong, sealed containers may lead to a large pressure build-up and subsequent explosion. Rapid and violent polymerisation possible at temperatures above 32 deg c. Avoid reaction with oxidising agents



X — Must not be stored together

0 — May be stored together with specific preventions

+ — May be stored together

Note: Depending on other risk factors, compatibility assessment based on the table above may not be relevant to storage situations, particularly where large volumes of dangerous goods are stored and handled. Reference should be made to the Safety Data Sheets for each substance or article and risks assessed accordingly.

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)	methyl methacrylate	Methyl methacrylate	50 ppm / 208 mg/m3	416 mg/m3 / 100 ppm	Not Available	(skin) - Skin absorption (dsen) - Dermal sensitiser
New Zealand Workplace Exposure Standards (WES)	2-(2'-hydroxy-5'- methylphenyl)benzotriazole	Inhalable dust (not otherwise classified)	10 mg/m3	Not Available	Not Available	Not Available
New Zealand Workplace Exposure Standards (WES)	2-(2'-hydroxy-5'- methylphenyl)benzotriazole	Respirable dust (not otherwise classified)	3 mg/m3	Not Available	Not Available	Not Available

Exposure controls

Appropriate engineering controls	can be highly effective in protecting workers and will typical The basic types of engineering controls are: Process controls which involve changing the way a job acti Enclosure and/or isolation of emission source which keeps strategically 'adds' and 'removes' air in the work environme design of a ventilation system must match the particular pro Employers may need to use multiple types of controls to pr For flammable liquids and flammable gases, local exhaust equipment should be explosion-resistant.	a selected hazard 'physically' away from the worker and ventila nt. Ventilation can remove or dilute an air contaminant if design ocess and chemical or contaminant in use. event employee overexposure. ventilation or a process enclosure ventilation system may be re ng 'escape' velocities which, in turn, determine the 'capture vel	vel of protection. ation that hed properly. The equired. Ventilation
	Type of Contaminant:		Air Speed:
	solvent, vapours, degreasing etc., evaporating from tank (in still air).	0.25-0.5 m/s (50-100 f/min.)
	aerosols, fumes from pouring operations, intermittent cont plating acid fumes, pickling (released at low velocity into z	ainer filling, low speed conveyer transfers, welding, spray drift, one of active generation)	0.5-1 m/s (100-200 f/min.)
	direct spray, spray painting in shallow booths, drum filling, generation into zone of rapid air motion)	conveyer loading, crusher dusts, gas discharge (active	1-2.5 m/s (200-500 f/min.)
	Within each range the appropriate value depends on:		
	Lower end of the range	Upper end of the range	
	1: Room air currents minimal or favourable to capture	1: Disturbing room air currents	
	2: Contaminants of low toxicity or of nuisance value only.	2: Contaminants of high toxicity	
	3: Intermittent, low production.	3: High production, heavy use	
	4: Large hood or large air mass in motion	4: Small hood-local control only	
	decreases with the square of distance from the extraction p adjusted, accordingly, after reference to distance from the of a minimum of 1-2 m/s (200-400 f/min.) for extraction of solv mechanical considerations, producing performance deficits multiplied by factors of 10 or more when extraction systems · Adequate ventilation is typically taken to be that which lim room or enclosure containing the dangerous substance. · Ventilation for plant and machinery is normally considered might potentially be present to no more than 25% of the LE additional safeguards are provided to prevent the formation emergency shutdown of the process might be used togethe ovens and gas turbine enclosures. · Temporary exhaust ventilation systems may be provided f	nce away from the opening of a simple extraction pipe. Velocity ooint (in simple cases). Therefore the air speed at the extraction contaminating source. The air velocity at the extraction fan, for rents generated in a tank 2 meters distant from the extraction p within the extraction apparatus, make it essential that theoretic are installed or used. It is the average concentration to no more than 25% of the LEL adequate if it limits the average concentration of any dangero L. However, an increase up to a maximum 50% LEL can be ac of a hazardous explosive atmosphere. For example, gas dete or with maintaining or increasing the exhaust ventilation on solv or non-routine higher-risk activities, such as cleaning, repair or release. The work procedures for such activities should be care	point should be example, should be oint. Other cal air velocities are within the building, us substance that ceptable where ctors linked to ent evaporating maintenance in

tanks or other confined spaces or in an emergency after a release. The work procedures for such activities should be carefully considere The atmosphere should be continuously monitored to ensure that ventilation is adequate and the area remains safe. Where workers will

	enter the space, the ventilation should ensure (irrespective of the provision of suitable breat	e that the concentration of the dangerous substance does not exceed 10% of the LEL hing apparatus)	
Individual protection measures, such as personal protective equipment			
Eye and face protection	describing the wearing of lenses or restric lens absorption and adsorption for the cla should be trained in their removal and sui irrigation immediately and remove contac	66 or national equivalent] rd; soft contact lenses may absorb and concentrate irritants. A written policy document, ctions on use, should be created for each workplace or task. This should include a review of ass of chemicals in use and an account of injury experience. Medical and first-aid personnel itable equipment should be readily available. In the event of chemical exposure, begin eye as soon as practicable. Lens should be removed at the first signs of eye redness or ean environment only after workers have washed hands thoroughly. [CDC NIOSH Current	
Skin protection	See Hand protection below		
Hands/feet protection	equipment, to avoid all possible skin cont Contaminated leather items, such as show The selection of suitable gloves does not only manufacturer. Where the chemical is a prepara advance and has therefore to be checked pric The exact break through time for substances when making a final choice. Personal hygiene is a key element of effective washed and dried thoroughly. Application of a Suitability and durability of glove type is deper- frequency and duration of contact, chemical resistance of glove material, glove thickness and dexterity Select gloves tested to a relevant standard (e When prolonged or frequently repeated cont 240 minutes according to EN 374, AS/NZS 21 When only brief contact is expected, a glove EN 374, AS/NZS 2161.10.1 or national equiva- some glove polymer types are less affected use. Contaminated gloves should be replaced. As defined in ASTM F-739-96 in any applicati Excellent when breakthrough time > 20 min Fair when breakthrough time < 20 min Fair when breakthrough time < 20 min Soor when glove material degrades For general applications, gloves with a thickness permeation efficiency of the glove will be depriced be based on consideration of the task require Glove thickness may also vary depending on technical data should always be taken into ac Note: Depending on the activity being conduc Thinner gloves (down to 0.1 mm or less) ma only likely to give short duration protection an Thicker gloves (up to 3 mm or more) may be or puncture potential Gloves must only be worn on clean hands. Aff moisturiser is recommended.	 ion in predisposed individuals. Care must be taken, when removing gloves and other protective fact. ies, belts and watch-bands should be removed and destroyed. y depend on the material, but also on further marks of quality which vary from manufacturer to ration of several substances, the resistance of the glove material can not be calculated in or to the application. has to be obtained from the manufacturer of the protective gloves and has to be observed e hand care. Gloves must only be worn on clean hands. After using gloves, hands should be a non-perfumed moisturiser is recommended. indent on usage. Important factors in the selection of gloves include: a.g. Europe EN 374, US F739, AS/NZS 2161.1 or national equivalent). tact may occur, a glove with a protection class of 5 or higher (breakthrough time greater than 161.10.1 or national equivalent) is recommended. a with a protection class of 3 or higher (breakthrough time greater than 60 minutes according to alent) is recommended. b wovement and this should be taken into account when considering gloves for long-term ion, gloves are rated as: 	
	Exposure condition Short time use; (few minutes less than 0.5 hour) Little physical stress	Jse only recommended gloves - using the wrong gloves may increase the risk: Use of thin nitrile rubber gloves: Nitrile rubber (0.1 mm) Excellent tactibility ('feel'), powder-free Disposable Inexpensive Give adequate protection to low molecular weigh acrylic monomers	
	Exposure condition Medium time use; less than 4 hours Physical stress (opening drums, using tools, etc.)	Use of medium thick nitrile rubber gloves Nitrile rubber, NRL (latex) free; <0.45 mm Moderate tactibility ('feel'), powder-free Disposable Moderate price Gives adequate protection for most acrylates up to 4 hours Do NOT give adequate protection to low molecular weight monomers at exposures longer than 1 hour	
	Exposure condition Long time Cleaning operations	Nitrile rubber, NRL (latex) free; >0.56 mm low tactibility ('feel'), powder free High price Gives adequate protection for most acrylates in combination with commonly used solvents up to 8 hours Do NOT give adequate protection to low molecular weight monomers at exposures longer than 1 hour Avoid use of ketones and acetates in wash-up solutions.	

Avoid use of ketones and acetates in wash-up solutions.

	Where none of this gloves ensure safe handling (for example in long term handling of acrylates containing high levels of acetates and/ or ketones, use laminated multilayer gloves. Guide to the Classification and Labelling of UV/EB Acrylates Third edition, 231 October 2007 - Cefic
Body protection	See Other protection below
Other protection	 Overalls. PVC Apron. PVC protective suit may be required if exposure severe. Eyewash unit. Ensure there is ready access to a safety shower. 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 ground the foot an shall dissipate static electricity from the body to reduce the possibility of ignition of volatile compounds. Electrical resistance must range between 0 to 500,000 ohms. Conductive shoes should be stored in lockers close to the room in which they are worn. Personnel who have been issued conductive footwear should not wear them from their place of work to their homes and return.

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:

Flowcrete Flowfast 319 Flexible Sealer Base A

Material	СРІ
PE/EVAL/PE	A
PVA	A
TEFLON	A
BUTYL	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.

Ansell Glove Selection

Glove — In order of recommendation
AlphaTec® 15-554
AlphaTec® Solvex® 37-675
AlphaTec® Solvex® 37-185
AlphaTec® 58-008
AlphaTec® 58-735
AlphaTec® 79-700
AlphaTec® 38-612
AlphaTec® 53-001
AlphaTec® 58-005
AlphaTec® 58-530B

The suggested gloves for use should be confirmed with the glove supplier.

SECTION 9 Physical and chemical properties

Information on basic physical and chemical properties

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

Respiratory protection

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	AK-AUS P2	-	AK-PAPR-AUS / Class 1 P2
up to 50 x ES	-	AK-AUS / Class 1 P2	-
up to 100 x ES	-	AK-2 P2	AK-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

information on basic physical and chemical properties			
Appearance	Violet viscous liquid		
Physical state	Liquid	Relative density (Water = 1)	0.99
Odour	Not Available	Partition coefficient n-octanol / water	Not Available
Odour threshold	Not Available	Auto-ignition temperature (°C)	430
pH (as supplied)	Not Available	Decomposition temperature (°C)	Not Available

Melting point / freezing point (°C)	-48	Viscosity (cSt)	161.62
Initial boiling point and boiling range (°C)	101	Molecular weight (g/mol)	Not Available
Flash point (°C)	12	Taste	Not Available
Evaporation rate	Not Available BuAC = 1	Explosive properties	Not Available
Flammability	HIGHLY FLAMMABLE.	Oxidising properties	Not Available
Upper Explosive Limit (%)	12.5	Surface Tension (dyn/cm or mN/m)	Not Available
Lower Explosive Limit (%)	2.1	Volatile Component (%vol)	Not Available
Vapour pressure (kPa)	3.87	Gas group	Not Available
Solubility in water	Immiscible	pH as a solution (1%)	Not Available
Vapour density (Air = 1)	Not Available	VOC g/L	910.80

SECTION 10 Stability and reactivity

Reactivity	See section 7
Chemical stability	 Stable under controlled storage conditions provided material contains adequate stabiliser / polymerisation inhibitor. Bulk storages may have special storage requirements WARNING: Gradual decomposition in strong, sealed containers may lead to a large pressure build-up and subsequent explosion. Rapid and violent polymerisation possible at temperatures above 32 deg c.
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

Inhaled	The material is not thought to produce either adverse health effects or irritation of the respiratory tract following inhalation (as classified by EC Directives using animal models). Nevertheless, adverse systemic effects have been produced following exposure of animals by at least one other route and good hygiene practice requires that exposure be kept to a minimum and that suitable control measures be used in an occupational setting. Inhalation of vapours may cause drowsiness and dizziness. This may be accompanied by sleepiness, reduced alertness, loss of reflexes,
	lack of co-ordination, and vertigo.
	The material is not thought to produce adverse health effects following ingestion (as classified by EC Directives using animal models). Nevertheless, adverse systemic effects have been produced following exposure of animals by at least one other route and good hygiene practice requires that exposure be kept to a minimum. At sufficiently high doses the material may be hepatotoxic (i.e. poisonous to the liver).
Ingestion	Oral doses can produce low blood pressure, central nervous system depression and drowsiness, liver and kidney degeneration and death
ingestion	after cessation of breathing.
	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.
	Accidental ingestion of the material may be damaging to the health of the individual.
	The liquid may be able to be mixed with fats or oils and may degrease the skin, producing a skin reaction described as non-allergic contact
	dermatitis. The material is unlikely to produce an irritant dermatitis as described in EC Directives.
Skin Contact	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	This material can cause eye irritation and damage in some persons.
Chronic	There has been concern that this material can cause cancer or mutations, but there is not enough data to make an assessment.
	Inhaling this product is more likely to cause a sensitisation reaction in some persons compared to the general population.
	Skin contact with the material is more likely to cause a sensitisation reaction in some persons compared to the general population.
	Toxic: danger of serious damage to health by prolonged exposure through inhalation, in contact with skin and if swallowed. This material can cause serious damage if one is exposed to it for long periods. It can be assumed that it contains a substance which can
	produce severe defects.
	Prolonged and repeated exposures can cause liver and kidney damage, low blood pressure and heart attack. There may be increased deaths from colon or rectal cancer. Long term local injection may cause tumour of the local tissues. When inhaled, it may cause watery and sore nostrils and destruction of the organ of smell.
	Respiratory sensitisation may result in allergic/asthma like responses; from coughing and minor breathing difficulties to bronchitis with wheezing, gasping.

Flowcrete Flowfast 319	TOXICITY		IRRITATION	
Flexible Sealer Base A	Not Available		Not Available	
	TOXICITY		IRRITATION	
	Dermal (rabbit) LD50: >5000 mg/kg ^[2]		Eye (rabbit): 150 mg	
methyl methacrylate	Inhalation (Rat) LC50: 29.8 mg/l4h ^[1]		Skin (rabbit): 10000 mg/kg (open)	
	Oral (Rat) LD50: 7872 mg/kg ^[2]			
	ΤΟΧΙCΙΤΥ		IRRITATION	
dipropoxy-p-toluidine	dermal (rat) LD50: >2000 mg/kg ^[1]		Eye (rabbit): slight* * = BAYER	
	Oral (Rat) LD50: >25<200 mg/kg ^[1]		Skin (rabbit): 4h - Non irrit.*	
	ΤΟΧΙΟΙΤΥ		IRRITATION	
	Dermal (rabbit) LD50: >177 mg/kg ^[1]		Eyes (rabbit) 500mg/24h mild	
2-ethylhexyl acrylate	Oral (Mouse) LD50; >5000 mg/kg ^[1]		Skin (rabbit) 10mg/24h - SEVERE	
			Skin (rabbit) 20mg/24h mod.	
			Skin (rabbit) 500mg mild	
	ΤΟΧΙCΙΤΥ	IRRITATION	l .	
4-butanediol dimethacrylate	Oral (Rat) LD50: 10.066 mg/kg ^[1]	Eye: no adve	erse effect observed (not irritating) ^[1]	
		Skin: no adv	erse effect observed (not irritating) ^[1]	
	ΤΟΧΙCΙΤΥ	IRRITA	TION	
2-(2'-hydroxy-5'-	Dermal (rabbit) LD50: >1000 mg/kg ^[2] Eye (rabl		obit): 500 mg/24 h - mild	
methylphenyl)benzotriazole	Inhalation (Rat) LC50: >0.59 mg/L4h ^[1] Eye: no a		adverse effect observed (not irritating) ^[1]	
			o adverse effect observed (not irritating) ^[1]	

Acute Toxicity	×	Carcinogenicity	×
Skin Irritation/Corrosion	×	Reproductivity	×
Serious Eye Damage/Irritation	✓	STOT - Single Exposure	×
Respiratory or Skin sensitisation	×	STOT - Repeated Exposure	*
Mutagenicity	×	Aspiration Hazard	×
			t available or does not fill the criteria for classification

Data available to make classification

SECTION 12 Ecological information

Flowcrete Flowfast 319 Flexible Sealer Base A	Endpoint	Test Duration (hr)		Species	Value	Sou	urce
	Not Available	Not Available	Not Available Not A		Not Available	Available Not Ava	
	Endpoint	Test Duration (hr)	Spec	ies		Value	Source
methyl methacrylate	EC0(ECx)	48h	Crust	acea		48mg/l	1
	EC50	96h	Algae	or other aquatic plant	S	170mg/l	1
	EC50	72h	Algae	or other aquatic plant	s	>110mg/l	2
	EC50	48h	Crust	acea		69mg/l	1
	LC50	96h	Fish			>79mg/l	2
dipropoxy-p-toluidine							
aipiopoxy-p-totalaille	Endpoint	Test Duration (hr)	Spe	cies		Value	Source

	EC50	72h	Algae or other aquatic plants		245mg/l	2
	EC50(ECx)	48h	Crustacea		28.8mg/l	2
	EC50	48h	Crustacea		28.8mg/l	2
	LC50	96h	Fish		17mg/l	2
	Endpoint	Test Duration (hr)	Species		Value	Source
	EC50	72h	Algae or other aquatic plants		1.71mg/l	2
2-ethylhexyl acrylate	LC50	96h	Fish		1.1mg/l	2
2-ethymexyr acrylate	EC50	48h	Crustacea		1.3mg/l	2
	NOEC(ECx)	504h	Crustacea		0.136mg/l	2
	EC50	96h	Algae or other aquatic plants	Algae or other aquatic plants		2
	Endpoint	Test Duration (hr)	Species		Value	Source
1.4 hutenediel dimetheemdete	NOEC(ECx)	72h	Algae or other aquatic plants		2.11mg/l	2
1,4-butanediol dimethacrylate	EC50	72h	Algae or other aquatic plants		4.97mg/l	2
	LC50	96h	Fish		12.4mg/l	2
	Endpoint	Test Duration (hr)	Species	Value	5	Source
	BCF	1344h	Fish	123-49	94 7	7
2-(2'-hydroxy-5'- methylphenyl)benzotriazole	EC50	72h	Algae or other aquatic plants	0.072n	ng/L 2	2
methylphenylphenzoulazole	EC50(ECx)	24h	Crustacea	20mg/l	1 1	Not Available
	LC50	96h	Fish	7.9mg/	/I I	Not Available
	L					

Harmful to aquatic organisms. **DO NOT** discharge into sewer or waterways.

Persistence and degradability

Ingredient	Persistence: Water/Soil	Persistence: Air
methyl methacrylate	LOW	LOW
dipropoxy-p-toluidine	HIGH	HIGH
2-ethylhexyl acrylate	LOW	LOW
1,4-butanediol dimethacrylate	LOW	LOW
2-(2'-hydroxy-5'- methylphenyl)benzotriazole	HIGH	HIGH

Bioaccumulative potential

Ingredient	Bioaccumulation
methyl methacrylate	LOW (BCF = 6.6)
dipropoxy-p-toluidine	LOW (LogKOW = 2.0121)
2-ethylhexyl acrylate	LOW (BCF = 289.73)
1,4-butanediol dimethacrylate	LOW (LogKOW = 3.191)
2-(2'-hydroxy-5'- methylphenyl)benzotriazole	LOW (BCF = 494)

Mobility in soil

Ingredient	Mobility
methyl methacrylate	LOW (Log KOC = 10.14)
dipropoxy-p-toluidine	LOW (Log KOC = 10)
2-ethylhexyl acrylate	LOW (Log KOC = 429)
1,4-butanediol dimethacrylate	LOW (Log KOC = 92.37)
2-(2'-hydroxy-5'- methylphenyl)benzotriazole	LOW (Log KOC = 100800)

SECTION 13 Disposal considerations

Waste treatment methods Product / Packaging disposal Containers may still present a chemical bazard/ danger w

- 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 same 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 their 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)

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 sewer may be subject to local laws and regulations and these should be considered first.
- Where in doubt contact the responsible authority.
- Recycle wherever possible.
- 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.
- 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).
- Decontaminate empty containers. Observe all label safeguards until containers are cleaned and destroyed.

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	•3YE

Land transport (UN)

1866		
RESIN SOLUTION, fla	RESIN SOLUTION, flammable	
Class Subsidiary Hazard	3 Not Applicable	
II		
Not Applicable		
Special provisions Limited quantity	Not Applicable 5 L	
	RESIN SOLUTION, fla Class Subsidiary Hazard II Not Applicable Special provisions	

Air transport (ICAO-IATA / DGR)

14.1. UN number	1866		
14.2. UN proper shipping name	Resin solution flammable		
14.3. Transport hazard class(es)	ICAO/IATA Class ICAO / IATA Subsidiary Hazard ERG Code	3 Not Applicable 3L	-
14.4. Packing group	П		

14.5. Environmental hazard	Not Applicable			
	Special provisions	A3		
14.6. Special precautions for user	Cargo Only Packing Instructions	364		
	Cargo Only Maximum Qty / Pack	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)

14.1. UN number	1866		
14.2. UN proper shipping name	RESIN SOLUTION flammable		
14.3. Transport hazard class(es)	IMDG Class IMDG Subsidiary Haz	3 rard Not Applicable	
14.4. Packing group	П		
14.5 Environmental hazard	Not Applicable		
14.6. Special precautions for user	EMS Number Special provisions Limited Quantities	F-E , S-E Not Applicable 5 L	

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

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
HSR002662	Surface Coatings and Colourants Flammable Group Standard 2020

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

Additional Regulatory Information

Not Applicable

Hazardous Substance Location

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

Hazard Class	Quantity (Closed Containers)	Quantity (Open Containers)
Flammable Liquid Category 2	100 L in containers more than 5 L	50 L
Flammable Liquid Category 2	250 L in containers up to and including 5 L	50 L

Certified Handler

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

Class of substance	Quantities
Not Applicable	Not Applicable

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	Liquid (L)	Maximum quantity per package for each classification
Skin Sensitisation Category 1	1	
Flammable Liquid Category 2		1L

Tracking Requirements

Not Applicable

National Inventory Status

National Inventory	Status
Australia - AIIC / Australia Non- Industrial Use	Yes
Canada - DSL	Yes
Canada - NDSL	No (methyl methacrylate; dipropoxy-p-toluidine; 2-ethylhexyl acrylate; 1,4-butanediol dimethacrylate)
China - IECSC	Yes
Europe - EINEC / ELINCS / NLP	Yes
Japan - ENCS	Yes
Korea - KECI	Yes
New Zealand - NZIoC	Yes
Philippines - PICCS	Yes
USA - TSCA	Yes
Taiwan - TCSI	Yes
Mexico - INSQ	No (dipropoxy-p-toluidine; 1,4-butanediol dimethacrylate)
Vietnam - NCI	Yes
Russia - FBEPH	No (dipropoxy-p-toluidine)
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	11/04/2024
Initial Date	25/01/2018

SDS Version Summary

Version	Date of Update	Sections Updated
5.11	11/04/2024	Hazards identification - Classification, Ecological Information - Environmental

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
- DNEL: Derived No-Effect Level
- PNEC: Predicted no-effect concentration
- 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

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Altex Coatings Ltd

Version No: 5.9

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

Chemwatch Hazard Alert Code: 4

Issue Date: 23/10/2023 Print Date: 23/10/2023 S.GHS.NZL.EN

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

Product Identifier	
Product name	Flowcrete Flowfast Catalyst
Synonyms	Not Available
Proper shipping name	ORGANIC PEROXIDE TYPE D, SOLID
Other means of identification	Not Available

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

Relevant identified uses	Part B of a multi-pack industrial coatings
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Details of the manufacturer or supplier of the safety data sheet

Registered company name	Altex Coatings Ltd
Address	91-111 Oropi Road,Tauranga 3112 Tauranga New Zealand
Telephone	+64 7 541 1221
Fax	Not Available
Website	www.altexcoatings.com
Email	neil.debenham@altexcoatings.co.nz

Emergency telephone number

Association / Organisation	NZ POISONS (24hr 7days)	CHEMWATCH EMERGENCY RESPONSE (24/7)
Emergency telephone numbers	0800 764766	+64 800 700 112
Other emergency telephone numbers	0800 700112	+61 3 9573 3188

Once connected and if the message is not in your preferred language then please dial 01

SECTION 2 Hazards identification

Classification of the substance or mixture

Considered a Hazardous Substance according to the criteria of the New Zealand Hazardous Substances New Organisms legislation. Classified as Dangerous Goods for transport purposes.

pulpecce.		
Classification ^[1]	Organic Peroxides Type D, Sensitisation (Skin) Category 1, Hazardous to the Aquatic Environment Acute Hazard Category 1, Hazardous to the Aquatic Environment Long-Term Hazard Category 1	
Legend:	1. Classified by Chemwatch; 2. Classification drawn from CCID EPA NZ; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI	

Label elements

Hazard pictogram(s)	
Signal word	Danger

Hazard statement(s)

H242	Heating may cause a fire.
H317	May cause an allergic skin reaction.
H410	Very toxic to aquatic life with long lasting effects.

Precautionary statement(s) Prevention

P210	Keep away from heat, hot surfaces, sparks, open flames and other ignition sources. No smoking.	
P234	Keep only in original packaging.	
P235	Keep cool.	

P240	Ground and bond container and receiving equipment.
P280	Wear protective gloves, protective clothing, eye protection and face protection.
P261	Avoid breathing dust/fumes.
P273	Avoid release to the environment.
P264	Wash all exposed external body areas thoroughly after handling.
P272	Contaminated work clothing should not be allowed out of the workplace.

Precautionary statement(s) Response

P370+P378	In case of fire: Use water jets to extinguish.
P302+P352	IF ON SKIN: Wash with plenty of water and soap.
P333+P313	If skin irritation or rash occurs: Get medical advice/attention.
P362+P364	Take off contaminated clothing and wash it before reuse.
P391	Collect spillage.

Precautionary statement(s) Storage

P403	Store in a well-ventilated place.
P411	Store at temperatures not exceeding 25°C
P410	Protect from sunlight.

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	
94-36-0*	40-50	dibenzoyl peroxide	
Legend:	 Classified by Chernwatch; 2. Classification drawn from CCID EPA NZ; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex Classification drawn from C&L * EU IOELVs available 		

SECTION 4 First aid measures

Description of first aid measur	es
Eye Contact	 If this product comes in contact with the eyes: Immediately hold the eyelids apart and flush the eye with 2% sodium carbonate solution or 5% sodium ascorbate solution then wash continuously for at least 15 minutes with fresh 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. Transport to hospital (or doctor) without further delay. Removal of contact lenses should only be undertaken by trained 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, aerosols or combustion products are inhaled remove from contaminated area. Other measures are usually unnecessary.
Ingestion	 Immediately give a glass of water. First aid is not generally required. If in doubt, contact a Poisons Information Centre or a doctor.

Indication of any immediate medical attention and special treatment needed

Treat symptomatically.

SECTION 5 Firefighting measures

Extinguishing media

FOR SMALL FIRE:

Water spray, foam, CO2 or dry chemical.

• **DO NOT** use water jets. FOR <u>LARGE FIRE</u>:

Flood fire area with water from a distance.

Special hazards arising from the substrate or mixture

Fire Incompatibility

Avoid storage with reducing agents.

Continued...

	Avoid any contamination of this material as it is very reactive and any contamination is potentially hazardous
vice for firefighters	
Fire Fighting	 Alert Fire Brigade and tell them location and nature of hazard. May be violently or explosively reactive. Wear full body protective clothing with breathing apparatus. Prevent, by any means available, spillage from entering drains or water courses. Consider evacuation (or protect in place). Fight fire from a safe distance, with adequate cover. Extinguishers should be used only by trained personnel. 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. If safe to do so, remove containers from path of fire. If fire gets out of control withdraw personnel and warn against entry. Equipment should be thoroughly decontaminated after use.
Fire/Explosion Hazard	 Will not burn but increases intensity of fire. May explode from friction, shock, heat or containment. Heating may cause expansion or decomposition leading to violent rupture of containers. Heat affected containers remain hazardous. Contact with combustibles such as wood, paper, oil or finely divided metal may produce spontaneous combustion or violent decomposition May emit irritating, poisonous or corrosive fumes. Combustion/decomposition may produce acrid/toxic fumes of carbon monoxide (CO). Combustion products include: carbon monoxide (CO) carbon dioxide (CO2) other pyrolysis products typical of burning organic material. Organic peroxides provide internal oxygen for combustion, so burn intensely. Simple smothering actions are not effective against established fires. NOTE: A Type D Organic Peroxide: may partially detonate does not deflagrate rapidly and shows no violent effect when heated under confinement

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	 Clean up all spills immediately. No smoking, naked lights, ignition sources. Avoid all contact with any organic matter including fuel, solvents, sawdust, paper or cloth and other incompatible materials, as ignition may result. Avoid breathing dust or vapours and all contact with skin and eyes. Control personal contact with the substance, by using protective equipment. Contain and absorb spill with dry sand, earth, inert material or vermiculite. DO NOT use sawdust as fire may result. Scoop up solid residues and seal in labelled drums for disposal. Neutralise/decontaminate area.
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 full body protective clothing with breathing apparatus. Prevent, by any means available, spillage from entering drains or water course. Consider evacuation (or protect in place). No smoking, flames or ignition sources. Increase ventilation. Contain spill with sand, earth or other clean, inert materials. NEVER use organic absorbents such as sawdust, paper, cloth; as fire may result. Avoid any contamination by organic matter. Use spark-free and explosion-proof equipment. Collect any recoverable product into labelled containers for possible recycling. DO NOT mix fresh with recovered material. Collect residues and prevent runoff into drains. Decontaminate equipment and launder all protective clothing before storage and re-use. If contamination of drains or waterways occurs advise emergency services.

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

SECTION 7 Handling and storage

Safe handling	 Nor only as much as is required Do NOT return the mixed material to original containers For oxidisers, including peroxides. Avoid personal contact and inhalation of dust, mist or vapours. Provide adequate ventilation. Always wear protective equipment and wash off any spillage from clothing. Keep material away from incompatible materials. Avoid physical damage to containers. Do NOT repack or return unused portions to original containers. Withdraw only sufficient amounts for immediate use. Use only minimum quantity required. Avoid using solutions of peroxides in volatile solvents. Solvent evaporation should be controlled to avoid dangerous concentration of the peroxide. Do NOT use peaks or return unused portions to original containers. Withdraw only sufficient amounts for immediate use. Use only minimum quantity required. Avoid using solutions of peroxides in volatile solvents. Solvent evaporation should be controlled to avoid dangerous concentration of the peroxide. Do NOT use glass containers with screw cap lids or glass stoppers. Store peroxides at the lowest possible temperature, consistent with their solubility and freezing point. CAUTION: Do NOT store liquids or solutions of peroxides at a temperature bolow that at which the coxidiser freezes or precipitates. Peroxides, in particular, in this form are extremely shock and heat-sensitive. Refrigerated storage of peroxides must ONLY be in explosion-proof units. The hazard sand consequences of fires and explosions outing synthesis and use of oxidisers is widely recognised; spontaneous or induced decomposition may culminate in a variety of ways, ranging from moderate gassing to spontaneous ignition or explosion. The heat released from spontaneous divers of the burning of peroxides an the surrounding temperature; the temperature will
Other information	 Store in original containers in an isolated approved flammable materials storage area. Keep containers securely sealed as supplied. WARNING: Gradual decomposition during storage in sealed containers may lead to a large pressure build-up and subsequent explosion. No smoking, naked lights, heat or ignition sources. Store in a cool, dry, well ventilated area. Store under cover and away from sunlight. Store below safe storage (control) temperature. Always store below 35 deg.C. Store away from flammable or combustible materials, debris and waste. Contact may cause fire or violent reaction. Store away from foodstuff containers DO NOT stack on wooden floors or wooden pallets. Protect containers against physical damage. Check regularly for spills and leaks. Observe manufacturer's storage and handling recommendations contained within this SDS. Keep locked up. Restrictions may apply on quantities and to other materials permitted in the same location.

Conditions for safe storage, including any incompatibilities

Suitable container	 Metal packagings meeting the test criteria of Packing Group I, must NOT be used; this avoids unnecessary confinement. Packagings for organic peroxides must be constructed so that none of the materials, which are in contact with the contents, will catalyse or otherwise dangerously affect the properties of their contents. For combination packages, cushioning materials must not be readily combustible and must NOT cause decomposition of the organic peroxide if leakage occurs. Generally only stainless steel 316, polyethylene or glass lined equipment is suitable for use when working with organic peroxides. NOTE: Dangerous decomposition reactions may occur at or above the SADT (self-accelerating decomposition temperature).Under certain circumstances explosion or fire may result. Contact with incompatible substances may cause decomposition at or below the SADT. Some plastics may be incompatible with this material, check with manufacturer for storage suitability. DO NOT repack. Use containers supplied by manufacturer only. Check that containers are clearly labelled Type D Solid Organic Peroxides, UN 3106, UN 3116 are to be packed to the requirements of Packing method OP7B of the ADG Code, with maximum mass of 50 ka.
Storage incompatibility	 Steel, Aluminium, Plastic drum / container or plastic inner receptacle in fibre-board or metal outer container. Incidents involving interaction of active oxidants and reducing agents, either by design or accident, are usually very energetic and examples of so-called redox reactions. Organic peroxides as a class are highly reactive. They are thermally unstable and prone to undergoing exothermic self-accelerating decomposition. Organic peroxides may decompose explosively, burn rapidly, be impact and/or friction sensitive and react dangerously with many other substances. Avoid storage with reducing agents.
	 Peroxides decompose over time and give off oxygen. Peroxides require controlled storage for stability. DANGER: Explosion hazard, never mix peroxides with accelerators or promoters. Avoid any contamination of this material as it is very reactive and any contamination is potentially hazardous



Х - Must not be stored together

 May be stored together with specific preventions
 May be stored together 0

+

Note: Depending on other risk factors, compatibility assessment based on the table above may not be relevant to storage situations, particularly where large volumes of dangerous goods are stored and handled. Reference should be made to the Safety Data Sheets for each substance or article and risks assessed accordingly.

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)	dibenzoyl peroxide	Benzoyl peroxide	5 mg/m3	Not Available	Not Available	(dsen) - Dermal sensitiser

Emergency Limits

Ingredient	TEEL-1	TEEL-2		TEEL-3
dibenzoyl peroxide	15 mg/m3	1,200 mg/m3		7,000 mg/m3
Ingredient	Original IDLH		Revised IDLH	
dibenzoyl peroxide	1,500 mg/m3		Not Available	

Exposure 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. * Employees exposed to confirmed human carcinogens should be authorized to do so by the employer, and work in a regulated area. * Work should be undertaken in an isolated system such as a 'glove-box'. Employees should wash their hands and arms upon completion of the assigned task and before engaging in other activities not associated with the isolated system. * Within regulated areas, the carcinogen should be stored in sealed containers, or enclosed in a closed system, including piping systems, with any sample ports or openings closed while the carcinogens are contained within. * Open-vessel systems are prohibited. * Each operation should be provided with continuous local exhaust ventilation so that air movement is always from ordinary work areas to the operation. * Exhaust air should not be discharged to regulated areas, non-regulated areas or the external environment unless decontaminated. Clean make-up air should be introduced in sufficient volume to maintain correct operation of the local exhaust system. * For maintenance and decontamin
 Chemical goggles. [AS/NZS 1337.1, EN166 or national equivalent] Full face shield may be required for supplementary but never for primary protection of eyes. 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 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. Lens should be removed at the first signs of eye redness or irritation - lens should be removed in a clean environment only after workers have washed hands thoroughly. [CDC NIOSH Current Intelligence Bulletin 59].
See Hand protection below
 Wear chemical protective gloves, e.g. PVC. Wear safety footwear or safety gumboots, e.g. Rubber 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.

	 DO NOT wear cotton or cotton-backed gloves. DO NOT wear leather gloves. Promptly hose all spills off leather shoes or boots or ensure that such footwear is protected with PVC over-shoes.
Body protection	See Other protection below
Other protection	 Employees working with confirmed human carcinogens should be provided with, and be required to wear, clean, full body protective clothing (smocks, coveralls, or long-sleeved shirt and pants), shoe covers and gloves prior to entering the regulated area. [AS/NZS ISO 6529:2006 or national equivalent] Employees engaged in handling operations involving carcinogens should be provided with, and required to wear and use half-face filter-type respirators with filters for dusts, mists and fumes, or air purifying canisters or cartridges. A respirator affording higher levels of protection may be substituted. [AS/NZS 1715 or national equivalent] Emergency deluge showers and eyewash fountains, supplied with potable water, should be located near, within sight of, and on the same level with locations where direct exposure is likely. Prior to each exit from an area containing confirmed human carcinogens, employees should be required to remove and leave protective clothing and equipment at the point of exit and at the last exit of the day, to place used clothing and equipment in impervious containers at the point of exit for purposes of decontamination activities, authorized employees entering the area should be provided with and required to wear clean, impervious garments, including gloves, boots and continuous-air supplied hod. Prior to removing protective garments the employee should undergo decontamination and be required to shower upon removal of the garments and hood. Overalls. PVC protective suit may be required if exposure severe. Eyewash unit. Ensure there is ready access to a safety shower. 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. Conduct

Respiratory protection

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

Required Minimum Protection Factor	Half-Face Respirator	Full-Face Respirator	Powered Air Respirator
up to 10 x ES	P1 Air-line*	-	PAPR-P1 -
up to 50 x ES	Air-line**	P2	PAPR-P2
up to 100 x ES	-	P3	-
		Air-line*	-
100+ x ES	-	Air-line**	PAPR-P3

* - Negative pressure demand ** - Continuous flow

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)

SECTION 9 Physical and chemical properties

Appearance	white powder		
Physical state	Divided Solid Powder	Relative density (Water = 1)	1.23
Odour	Not Available	Partition coefficient n-octanol / water	Not Available
Odour threshold	Not Available	Auto-ignition temperature (°C)	Not Available
pH (as supplied)	Not Available	Decomposition temperature (°C)	55
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 Available
Flash point (°C)	Not Available	Taste	Not Available
Evaporation rate	Not Available	Explosive properties	Not Available
Flammability	Not Available	Oxidising properties	Not Available
Upper Explosive Limit (%)	Not Available	Surface Tension (dyn/cm or mN/m)	Not Applicable
Lower Explosive Limit (%)	Not Available	Volatile Component (%vol)	Not Available

Vapour pressure (kPa)	Not Available	Gas group	Not Available
Solubility in water	Immiscible	pH as a solution (1%)	Not Available
Vapour density (Air = 1)	Not Available	VOC g/L	0.00

SECTION 10 Stability and reactivity

Reactivity	See section 7
Chemical stability	 Unstable in the presence of incompatible materials. Product is considered stable under normal handling conditions. Prolonged exposure to heat. Hazardous polymerisation will not occur. NOTE: A range of exothermic decomposition energies for peroxides is given as 200-340 kJ/mol. The relationship between energy of decomposition and processing hazards has been the subject of discussion; it is suggested that values of energy releases per unit of mass, rather than on a molar mass 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 energies below 500 J/g are unlikely to present a danger, whilst those in closed vessel processes (opening is a safety valve or bursting disk) present some danger where the decomposition energy exceeds 150 J/g. BRETHERICK: Handbook of Reactive Chemical Hazards, 4th Edition
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	The material can cause respiratory irritation in some persons. The body's response to such irritation can cause further lung damage.			
Ingestion	The material has NOT been classified by EC Directives or other classification systems as 'harmful by ingestion'. This is because of the lack of corroborating animal or human evidence.			
Skin Contact	Skin contact is not thought to have harmful health effects (as classified under EC Directives); the material may still produce health damage following entry through wounds, lesions or abrasions. 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. There is some evidence to suggest that the material may cause mild but significant 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.			
Eye	Evidence exists, or practical experience predicts, that the material may cause eye irritation in a substantial number of individuals. Prolonged eye contact may cause inflammation characterised by a temporary redness of the conjunctiva (similar to windburn).			
	Long-term exposure to respiratory irritants may result in airways			
Chronic	Skin contact with the material is more likely to cause a sensitisati Ample evidence from experiments exists that there is a suspicior Based on experience with animal studies, exposure to the materi not cause significant toxic effects to the mother. Substance accumulation, in the human body, may occur and may There has been some concern that this material can cause cance	n this material dire ial may result in to y cause some cor	ectly reduces fertility. bxic effects to the development of the foetus, at levels which neern following repeated or long-term occupational exposure	
Chronic	Ample evidence from experiments exists that there is a suspicior Based on experience with animal studies, exposure to the materi not cause significant toxic effects to the mother. Substance accumulation, in the human body, may occur and may	n this material dire ial may result in to y cause some cor	ectly reduces fertility. boxic effects to the development of the foetus, at levels which neern following repeated or long-term occupational exposure ut there is not enough data to make an assessment.	
Chronic Flowcrete Flowfast Catalyst	Ample evidence from experiments exists that there is a suspicior Based on experience with animal studies, exposure to the materi not cause significant toxic effects to the mother. Substance accumulation, in the human body, may occur and may There has been some concern that this material can cause cance	n this material dire ial may result in to y cause some cor er or mutations bu	ectly reduces fertility. boxic effects to the development of the foetus, at levels which incern following repeated or long-term occupational exposure at there is not enough data to make an assessment.	
	Ample evidence from experiments exists that there is a suspicior Based on experience with animal studies, exposure to the materi not cause significant toxic effects to the mother. Substance accumulation, in the human body, may occur and may There has been some concern that this material can cause cance	n this material dire ial may result in to y cause some cor er or mutations bu IRRITAT	ectly reduces fertility. boxic effects to the development of the foetus, at levels which incern following repeated or long-term occupational exposure at there is not enough data to make an assessment.	
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Flowcrete Flowfast Catalyst	Ample evidence from experiments exists that there is a suspicior Based on experience with animal studies, exposure to the materi not cause significant toxic effects to the mother. Substance accumulation, in the human body, may occur and may There has been some concern that this material can cause cance TOXICITY Not Available TOXICITY	n this material dire ial may result in to y cause some cor er or mutations bu IRRITAT	ectly reduces fertility. Description of the foetus, at levels which accern following repeated or long-term occupational exposure at there is not enough data to make an assessment. FION illable IRRITATION	
	Ample evidence from experiments exists that there is a suspicion Based on experience with animal studies, exposure to the materi not cause significant toxic effects to the mother. Substance accumulation, in the human body, may occur and may There has been some concern that this material can cause cance TOXICITY Not Available TOXICITY Inhalation (Human) TCLo: 12 mg/m3 ^[2]	n this material dire ial may result in to y cause some cor er or mutations bu IRRITAT	ectly reduces fertility. oxic effects to the development of the foetus, at levels which incern following repeated or long-term occupational exposure ut there is not enough data to make an assessment. FION ilable Eye (rabbit): 500 mg/24h - mild	
Flowcrete Flowfast Catalyst	Ample evidence from experiments exists that there is a suspicior Based on experience with animal studies, exposure to the materi not cause significant toxic effects to the mother. Substance accumulation, in the human body, may occur and may There has been some concern that this material can cause cance TOXICITY Not Available TOXICITY Inhalation (Human) TCLo: 12 mg/m3 ^[2] Intraperitoneal (Mouse) LD50: 440 mg/kg ^[2]	n this material dire ial may result in to y cause some cor er or mutations bu IRRITAT	ectly reduces fertility. oxic effects to the development of the foetus, at levels which incern following repeated or long-term occupational exposure ut there is not enough data to make an assessment. FION ilable Eye (rabbit): 500 mg/24h - mild	
Flowcrete Flowfast Catalyst	Ample evidence from experiments exists that there is a suspicior Based on experience with animal studies, exposure to the materinot cause significant toxic effects to the mother. Substance accumulation, in the human body, may occur and may There has been some concern that this material can cause cance TOXICITY Not Available TOXICITY Inhalation (Human) TCLo: 12 mg/m3 ^[2] Intraperitoneal (Mouse) LD50: 440 mg/kg ^[2] Intravenous (Rabbit) LD: 16 mg/kg ^[2]	n this material dire ial may result in to y cause some cor er or mutations bu IRRITAT	ectly reduces fertility. oxic effects to the development of the foetus, at levels which incern following repeated or long-term occupational exposure ut there is not enough data to make an assessment. FION ilable Eye (rabbit): 500 mg/24h - mild	

×

Skin Irritation/Corrosion	×	Reproductivity	×
Serious Eye Damage/Irritation	×	STOT - Single Exposure	×
Respiratory or Skin sensitisation	*	STOT - Repeated Exposure	×
Mutagenicity	×	Aspiration Hazard	×
		legend: 🗙 – Data either not	available or does not fill the criteria for classification

Data available to make classification

SECTION 12 Ecological information

Flowcrete Flowfast Catalyst	Endpoint	Test Duration (hr)		Species	Value	Sou	rce
	Not Available	Not Available		Not Available	Not Available	Not	Available
	Endpoint	Test Duration (hr)	Spe	cies		Value	Source
dibenzoyl peroxide	EC50	72h	Alga	e or other aquatic plan	ts	0.042mg/l	2
	EC50	48h	Crus	tacea		0.11mg/l	2
	EC10(ECx)	504h	Crus	tacea		0.001mg/l	2
	LC50	96h	Fish			0.06mg/l	2
Legend:	E	IUCLID Toxicity Data 2. Europe				diana Annadia Tau	

Very toxic to aquatic organisms.

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.

DO NOT discharge into sewer or waterways.

Persistence and degradability

Ingredient	Persistence: Water/Soil	Persistence: Air
dibenzoyl peroxide	LOW (Half-life = 14 days)	LOW (Half-life = 21.25 days)

Bioaccumulative potential

Ingredient	Bioaccumulation	
dibenzoyl peroxide	LOW (LogKOW = 3.46)	

Mobility in soil

Ingredient	Mobility
dibenzoyl peroxide	LOW (KOC = 771)

SECTION 13 Disposal considerations

Waste treatment methods

Product / Packaging disposal	 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 same 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. For small quantities of oxidising agent: Cautiously acidify a 3% solution to pH 2 with sulfuric acid. Gradually add a 50% excess of sodium bisulfite solution with stirring. Add a further 10% sodium bisulfite. If no further reaction occurs (as indicated by a rise in temperature) cautiously add more acid.
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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.

Detonation, deflagration or controlled combustion of 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, 2, 3 or 4 substances.

Remove any ignition source from the disposal site.

SECTION 14 Transport information

Labels Required

	52
Marine Pollutant	
HAZCHEM	1WE

Land transport (UN)

14.1. UN number or ID number	3106		
14.2. UN proper shipping name	ORGANIC PEROXIDE TYPE D, SOLID		
14.3. Transport hazard class(es)	Class Subsidiary Hazard	5.2 Not Applicable	
14.4. Packing group	Not Applicable		
14.5. Environmental hazard	Environmentally hazardous		
14.6. Special precautions for user	Special provisions Limited quantity	Special provisions 122; 274	

Air transport (ICAO-IATA / DGR)

14.1. UN number	3106					
14.2. UN proper shipping name	Organic peroxide type D, solid *					
	ICAO/IATA Class	5.2				
14.3. Transport hazard class(es)	ICAO / IATA Subsidiary Hazard	ard Not Applicable				
01400(00)	ERG Code	ERG Code 5L				
14.4. Packing group	Not Applicable					
14.5. Environmental hazard	Environmentally hazardous					
	Special provisions		A20 A802			
14.6. Special precautions for user	Cargo Only Packing Instructions		570			
	Cargo Only Maximum Qty / Pack		10 kg			
	Passenger and Cargo Packing Instructions		570			
	Passenger and Cargo Maximum Qty / Pack		5 kg			
	Passenger and Cargo Limited Quantity Packing Instructions		Forbidden			
	Passenger and Cargo Limited Maximum Qty / Pack		Forbidden			
	1					

Sea transport (IMDG-Code / GGVSee)

14.1. UN number	3106		
14.2. UN proper shipping name	ORGANIC PEROXIDE TYPE D, SOLID		
14.3. Transport hazard class(es)	IMDG Class IMDG Subsidiary Hazard	5.2 Not Applicable	
14.4. Packing group	Not Applicable		

14.5 Environmental hazard	Marine Pollutant	
14.6. Special precautions for user	EMS Number	F-J, S-R
	Special provisions	122 274
	Limited Quantities	500 g

14.7.1. Transport in bulk according to Annex II of MARPOL and the IBC code

Not Applicable

SECTION 15 Regulatory information

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

This substance can be managed under the controls specified in the Transfer Notice or alternatively it may be managed using the conditions specified in an applicable Group Standard.

HSR Number	Group Standard
HSR002629(Proposed)	Organic Peroxides Group Standard 2020

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

Hazardous Substance Location

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

Hazard Class	Quantities
Organic Peroxide Type D	> 10 kg

Certified Handler

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

Class of substance	Quantities
Not Applicable	Not Applicable

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	Solid (kg)	Maximum quantity per package for each classification
Skin Sensitisation Category 1	3	
Organic Peroxide Type D		125 ml or 0.5 kg

Tracking Requirements

Not Applicable

National Inventory Status

National Inventory	Status
Australia - AIIC / Australia Non-Industrial Use	Yes
Canada - DSL	Yes
Canada - NDSL	No (dibenzoyl peroxide)
China - IECSC	Yes
Europe - EINEC / ELINCS / NLP	Yes
Japan - ENCS	Yes
Korea - KECI	Yes
New Zealand - NZIoC	Yes
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

202	Version	Summary	

Initial Date

08/07/2019

Version	Date of Update	Sections Updated
4.9 2	23/10/2023	Toxicological information - Acute Health (eye), First Aid measures - Advice to Doctor, Toxicological information - Chronic Health, Hazards identification - Classification, Disposal considerations - Disposal, Exposure controls / personal protection - Engineering Control, Exposure controls / personal protection - Exposure Standard, Composition / information on ingredients - Ingredients, Exposure controls / personal protection - Personal Protection (other), Identification of the substance / mixture and of the company / undertaking - Supplier Information

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
- DNEL: Derived No-Effect Level
- PNEC: Predicted no-effect concentration
- 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
 NI P: No-Longer Polymers
- NLP: No-Longer Polymers
- ENCS: Existing and New Chemical Substances Inventory
 KECI: Korea Existing Chemicals 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

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