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Infosafe No™ 3CHHF

Issue Date :March 2021

RE-ISSUED by ABS

Product Name RAPID DIFF FIXATIVE

Classified as hazardous

1. Identification						
GHS Product Identifier	RAPID DIFF FIXATIVE					
Company Name	AUSTRALIAN BIOSTAIN Pty Ltd					
Address	24 - 28 Stratton Drive, Traralgon, Victoria, Australia, 3844 www.australianbiostain.com.au					
Telephone/Fax Number	Tel: (03) 5176 2855					
Emergency phone number	CHEMCALL (24 hours): 1800 127 406 (Australia) / +64-4-917-9888 (International)					
E-mail Address	www.australianbiostain.com.au					
Recommended use of the chemical and restrictions on use Other Names	Laboratory reagent. Name Product Code					
Other Names	RAPID DIFF FIXATIVE ARDF					
Other Information	Australian Biostain Pty Ltd does not warrant that this product is suitable for any use or purpose. The user must ascertain the suitability of the product before use or application intended purpose. Preliminary testing of the product before use or application is recommended. Any reliance or purported reliance upon Australian Biostain Pty Ltd with respect to any skill or judgement or advice in relation to the suitability of this product of any purpose is disclaimed. Except to the extent prohibited at law, any condition implied by any statute as to the merchantable quality of this product or fitness for any purpose is hereby excluded. This product is not sold by description. Where the provisions of Part V, Division 2 of the Trade Practices Act apply, the liability of Australian Biostain Pty Ltd is limited to the replacement of supply of equivalent goods or payment of the cost of replacing the goods or acquiring equivalent goods.					
2. Hazard Identifi	cation					
the substance/mixture	Acute Toxicity - Dermal: Category 3 Flammable Liquids: Category 2 Acute Toxicity - Inhalation: Category 3 Acute Toxicity - Oral: Category 3 Specific target organ toxicity - Single Exposure Category 1, Eyes					
Signal Word (s)	DANGER					
Hazard Statement (s) Pictogram (s)	H225 Highly flammable liquid and vapour. H301 Toxic if swallowed. H311 Toxic in contact with skin. H331 Toxic if inhaled. H370 Causes damage to organs, eyes. Flame, Health hazard, Skull and crossbones					
g (<i>s</i>)						

Precautionary statement – Prevention
P210 Keep away from heat/sparks/open flames/hot surfaces. - No smoking. P233 Keep container tightly closed. P240 Ground/bond container and receiving equipment. P241 Use explosion-proof electrical/ventilating/lighting/.../equipment. P242 Use only non-sparking tools. P243 Take precautionary measures against static discharge. P261 Avoid breathing dust/fume/gas/mist/vapours/spray. P264 Wash thoroughly after handling. P270 Do not eat, drink or smoke when using this product.



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	P271 Use only outdoors or in a well-ventilated area. P280 Wear protective gloves/protective clothing/eye protection/face protection.
Precautionary statement – Response	Swallowed P301+P310 IF SWALLOWED: Immediately call a POISON CENTER or doctor/physician. P330 Rinse mouth. Skin
	<pre>P303+P361+P353 IF ON SKIN (or hair): Remove/Take off immediately all contaminated clothing. Rinse skin with water/shower. P312 Call a POISON CENTER or doctor/physician if you feel unwell. P361 Remove/Take off immediately all contaminated clothing. P363 Wash contaminated clothing before reuse. Inhaled P304+P340 IF INHALED: Remove victim to fresh air and keep at rest in a position comfortable for breathing. P311 Call a POISON CENTER or doctor/physician. Fire P370+P378 In case of fire: Use foam, dry chemical, carbon dioaxide or water spray for extinction.</pre>
Precautionary statement – Storage	P403+P233+P235 Store in a well-ventilated place. Keep container tightly closed. Keep cool. P405 Store locked up.
Precautionary statement – Disposal	P501 Dispose of contents/container to an approved waste disposal plant.

3. Composition/information on ingredients

Ingredients	Name	CAS	Proportion	
	Methyl Alcohol Thiazin dye	67-56-1 61-73-4	>99 % <0.001 %	

4. First-aid measures

Inhalation	If inhaled, remove from contaminated area to fresh air immediately, avoid
	becoming a casualty. Make patient comfortable, keep warm and at rest until fully recovered. If breathing is difficult (or develops a bluish skin
	discolouration), supply oxygen by a qualified person. Apply artificial respiration with a respiratory medical device if not breathing. Do not use mouth to mouth resuscitation. Immediately medical attention is required.
Ingestion	Rinse mouth thoroughly with water immediately. DO NOT INDUCE VOMITING. Seek immediate medical advice.
Skin	Wash affected areas with copious quantities of water and soap. Remove contaminated clothing and wash before re-use. If rapid recovery does not occur, obtain medical attention
Eye contact	If contact with the eye(s) occurs, wash with copious amounts of water for approximately 15 minutes holding eyelid(s) open. Take care not to rinse contaminated water into the non-affected eye. Seek medical attention.
First Aid Facilities	Maintain eyewash fountain and safety shower in work area.
Advice to Doctor	Effects may be delayed. Treat symptomatically based on judgement of doctor and individual reactions of the patient.
	The severity of outcome following methanol ingestion may be more related to the time between ingestion and treatment, rather than the amount ingested. Therefore, there is a need for rapid treatment of any ingestion exposure. Ethanol (contained in alcoholic beverages) can slow the metabolism of methanol, thus reducing the potential for harmful effects.
Other Information	For advice, contact a Poisons Information Centre (Phone eg Australia 13 1126; New Zealand 0800 764 766) or a doctor.
5. Fire-fighting n	neasures

Hazards from Combustion	Carbon dioxide, carbon monoxide, formaldehyde and other toxic, irritating chemicals.
Products Specific Methods	Caution: Use of water spray when fighting fire may be inefficient.

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		ed as hazardou	Classified as hazardous					
If safe to do so, m with flooding quant water inside contai HIGHLY FLAMMABLE: T by heat, sparks or Vapours may travel heavier than air ar tanks). Many liquid heated. Fire will p	m, fog op ove undar ities of ners. hese liqu flame. Va to source d will co s are liq roduce in	r water spray - maged containers water until we uids have a low apours will forr of ignition an ollect in low or ghter than water rritating, poise	Do not use water jets. s from fire area. Cool containers ll after fire is out. Avoid getting flashpoint - Will be easily ignited m explosive mixtures with air. nd flash back. Most vapours are r confined areas (drains, basements, r. Containers may explode when onous and/or corrosive gases.					
I V V H k V H k V H k V F K V V V V V V V V V V V V V V V V V	If safe to do so, movith flooding quant. Water inside contain HIGHLY FLAMMABLE: The Dy heat, sparks or Mapours may travel heavier than air and tanks). Many liquid heated. Fire will p Mapours from runoff 20WE Near SCBA and fully substances. Structure materials.	If safe to do so, move undar with flooding quantities of water inside containers. HIGHLY FLAMMABLE: These lique by heat, sparks or flame. Va Vapours may travel to source heavier than air and will con- tanks). Many liquids are lique heated. Fire will produce in Vapours from runoff may created v2WE Wear SCBA and fully-encapsus substances. Structural fires materials.	If safe to do so, move undamaged containers with flooding quantities of water until we water inside containers. HIGHLY FLAMMABLE: These liquids have a low by heat, sparks or flame. Vapours will for Vapours may travel to source of ignition an heavier than air and will collect in low of canks). Many liquids are lighter than water heated. Fire will produce irritating, poise Vapours from runoff may create explosion ha 2WE Wear SCBA and fully-encapsulating, gas-tight substances. Structural firefighter's unifor					

Spills & Disposal	ELIMINATE all ignition sources (no smoking, flares, sparks or flame) within at least 50m - All equipment used when handling the product must be earthed. Do not touch or walk through spilled material. Stop leak if safe to do so - Prevent entry into waterways, drains or confined areas. Vapour-suppressing foam may be used to control vapours - Water spray may be used to knock down or divert vapour clouds. Absorb with earth, sand or other non-combustible material. Use clean, non-sparking tools to collect absorbed material and place it into loosely-covered metal or plastic containers for later disposal. SEEK EXPERT ADVICE ON HANDLING AND DISPOSAL.
Personal Precautions	Evacuate the area of all non-essential personnel. Avoid inhalation, contact with skin, eyes and clothing.
Personal Protection	Wear protective clothing specified for normal operations (see Section 8)

7. Handling and storage

Precautions for Safe Handling	Avoid contact with eyes. Avoid contact with skin. Avoid breathing dust (or) vapour (or) spray mist. Keep locked up. Keep containers tightly sealed. Protect against physical damage. Avoid use in confined spaces. Ensure good ventilation/exhaustion at the workplace. Work under hood. In case of insufficient ventilation, wear suitable respiratory equipment. Avoid prolonged or repeated exposure. Do not ingest. If ingested, seek medical advice immediately and show the container or the label. Wear suitable protective clothing. Safety glasses. Wash thoroughly after handling. Remove contaminated clothing and wash before reuse. Keep away from heat and ignition sources - Do not smoke. Take precautions against static discharge. All electrical equipment must be flameproofed. Fumes can combine with air to form an explosive mixture. Avoid generation of vapours/aerosols. Containers should be bonded and grounded for transfers to avoid static sparks. Storage and use areas should be No Smoking areas. Use non-sparking type tools and equipment, including explosion proof ventilation. Containers of this material may be hazardous when empty since they retain product residues (vapours, liquid); observe all warnings and precautions listed for the product. Do Not attempt to clean empty containers since residue is difficult to remove. Do not pressurize, cut, weld, braze, solder, drill, grind or expose such containers to heat, sparks, flame, static electricity or other sources of ignition: they may explode and cause injury or death. Do not expose to temperatures above 60 °C.
Conditions for safe storage, including any incompatibilities	Store in a locked cabinet or with access restricted to technical experts or their assistants. Store small containers in suitable flammable liquid storage cabinets when not in use. Larger drums (200L) must be kept in purpose-built stores. Outside or detached storage is preferred. Store in well-sealed, dry containers, in a cool, well-ventilated location, away from any area where the fire hazard may be acute and protected from direct sunlight. Keep away from heat, sparks, open flames and all possible sources of ignition. Protect against physical damage. Separate from incompatibles. Do not store together with oxidizing and acidic materials. Aluminium, magnesium powder. Containers



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Corrosiveness	naval bronze, nic resistance (less types 304/347, 31	corrosive to most metals. Admiral ckel-resist and silicon copper hav than 2 mm (50.8 µm) penetration/ 6 and 400 stainless steels, copper talum, titanium and zirconium hav ear).	ve excellent corrosion year), while carbon steel, er, brass, bronze, aluminium,			
Storage Regulations	substances'. Ref	Standard AS/NZS 4452:1997 'The st fer Australian Standard AS 1940-20 combustible liquids'.				
Handling Temperatures	60°C maximum.					
Storage Temperatures	Store at room tem	perature (15 to 25 °C recommended	d). 60 °C Maximum.			
Unsuitable Materials	elastomers (such	ach as ABS and Isophthalic polyest as Viton A, hard and soft rubber, poxy general purpose coatings, all	, polyether-urethane and			

8. Exposure controls/personal protection

Other ExposureThese occup as lo be us chemin A tim [Meth STEL exposion not k minut TWA is calcu AbsoreAppropriate engineering controlsMainte proces at th Nered Avoid with Devid the fidust/ levelEye ProtectionThe transmission calcu Absore	<u>a</u>	S	TEL	5	rwa.	
Other ExposureThese occup as lo be us chemi A tim [Meth STEL expos not k minut TWA i calcu AbsorAppropriate engineering controlsMaint proce at th Nere ProtectionRespiratory ProtectionWhere at th uith uith Device the f dust/ levelEye ProtectionThe u occup occup at the count		mg/m3	ppm	mg/m3	ppm	Footnote
Information occup as lo be us chemi A tin [Meth STEL expos not k minut TWA i calcu Absor Appropriate Maint engineering controls at th Respiratory Where Protection Avoid with Device the f dust/ level	yl Alcohol	328	250	262	200	
engineering controls proce at th Respiratory Where Protection Avoid with with Devid the f dust/ level Eye Protection The u	These Workplace Exposure Standards are guides to be used in the control of occupational health hazards. All atmospheric contamination should be kept to as low a level as is workable. These workplace exposure standards should not be used as fine dividing lines between safe and dangerous concentrations of chemicals. They are not a measure of relative toxicity. A time weighted average (TWA) has been established for Methyl alcohol [Methanol] (Safe Work Australia) of 262 mg/m ³ , (200 ppm). The corresponding STEL level is 328 mg/m ³ , (250 ppm). The STEL (Short Term Exposure Limit) is an exposure value that should not be exceeded for more than 15 minutes and should not be repeated for more than 4 times per day. There should be at least 60 minutes between successive exposures at the STEL. The exposure value at the TWA is the average airborne concentration of a particular substance when calculated over a normal 8 hour working day for a 5 day working week. Note: Absorption through the skin may be a significant source of exposure.					
Protection Avoic with with Devic the f dust/ level Eye Protection The u	Maintain the concentrations values below the TWA. This may be achieved by process modification, use of local exhaust ventilation, capturing substances at the source, or other methods.					
	Where ventilation is not adequate, respiratory protection may be required. Avoid breathing vapours or mists. Select and use respirators in accordance with AS 1716 - Respiratory Protective Devices and be selected in accordance with AS 1715 - Selection, Use and Maintenance of Respiratory Protective Devices. When mists or vapours exceed the exposure standards then the use of the following is recommended: Approved respirator with organic vapour and dust/mist filters. Filter capacity and respirator type depends on exposure levels.					
	The use of a face shield, chemical goggles or safety glasses with side shield protection as appropriate. Must comply with Australian Standards AS 1337 and be selected and used in accordance with AS 1336.					
prote	e gloves of impervious r ective gloves - Select copriate glove type will	ion, use a	and maint	enance.	Final cho	pice of



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	can include methods of handling, and engineering controls as determined by appropriate risk assessments. Avoid skin contact when removing gloves from hands, do not touch the gloves outer surface. Dispose of gloves as hazardous waste.
Personal Protective Equipment Footwear	Personal protective equipment should not solely be relied upon to control risk and should only be used when all other reasonably practicable control measures do not eliminate or sufficiently minimise risk. Guidance in selecting personal protective equipment can be obtained from Australian, Australian/New Zealand or other approved standards. Rubber boots.
Footwear	
Body Protection	Flame retardant antistatic protective clothing. Clean clothing or protective clothing should be worn, preferably with an apron. Clothing for protection against chemicals should comply with AS 3765 Clothing for Protection Against Hazardous Chemicals.
Hygiene Measures	Always wash hands before smoking, eating or using the toilet. Wash contaminated clothing and other protective equipment before storing or re-using.

9. Physical and chemical properties

Form	Liquid
Appearance	Clear, mobile, volatile, highly polar liquid.
Odour	Mild, characteristic alcohol odour, when pure. Crude methanol may have a repulsive, pungent odour.
Melting Point	-97.8 °C
Boiling Point	64.7 °C
Solubility in Water	Miscible in water in all proportions.
Solubility in Organic Solvents	Miscible with other alcohols, esters, ketones, ethers and most other organic solvents.
Specific Gravity	0.791 at 20 °C
рН	Not available. Methanol is both a weak acid and a weak base.
Vapour Pressure	128 hPa (96 mm Hg) at 20 °C
Vapour Density (Air=1)	1.1 (air = 1)
Evaporation Rate	4.1 $(n-butyl acetate = 1)$
Odour Threshold	Reported values vary widely; 4.2-5960 ppm (geometric mean: 160 ppm) (detection); 53-8940 ppm (geometric mean: 690 ppm) (recognition). 100 %
Volatile Component	
Partition Coefficient: n-octanol/water	Log P(oct) = -0.77
Surface Tension	22.5 mN/m (22.5 dynes/cm) at 20 °C
Flash Point	9.7 °C (closed cup)
Flammability	HIGHLY FLAMMABLE. Keep away from heat, sparks or naked flames. Use flameproof equipment and fittings to prevent flammability risk. Electrically link and ground metal containers for transfer of the product to prevent accumulation of static electricity. Ensure adequate ventilation to prevent an explosive vapour-air mixture. Vapours will travel considerable distances to sources of ignition.
Auto-Ignition Temperature	Reported values vary: 385 °C; 455 °C; 464-470 °C
Flammable Limits - Lower	5.5 vol%
Flammable Limits - Upper	36.5 vol%



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Explosion Properties	air, at or a a source of agents may r explosion. O heat. Mixtur evolution of than militar due to the f Mixtures wit bromide. Mix shock-sensit beryllium hy (tetraphosph	bove 11 °C over a ignition of suffi react violently or concentrated perox tes with mineral a theat. Mixtures w ty explosives. Mix formation of hydro th acetyl bromide tures with perchl tive or explosive dride, cyanuric c forus hexaoxide) m th diethyl zinc re	wide concentrat cient energy. Mi explosively; in tide and methanol cids may react v with powdered met tures with alkal gen-air mixtures react violently, oric acid or met compounds. Mixtu chloride, isocyan	y form explosive mixtures with ion range, and may be ignited by xtures with strong oxidizing creased risk of fire and can be detonated by shock or igorously or violently, with the als can detonate, with more power i metals may react explosively , unless air is excluded. with the evolution of hydrogen al perchlorates may form res with alkyaluminium solutions, ates or phosphorus (III) oxide ly with generation of heat. with ignition.
Dynamic Viscosity	0.61 mPa.s a	t 20 °C		
Other Information	Refractive i	ndex: 1.329 @ 20	°C	
10. Stability and r	eactivity			

v	U U
Chemical Stability	Normally stable. Decomposes on heating to produce carbon monoxide and formaldehyde. Hygroscopic (absorbs moisture from the air).
Conditions to Avoid	Heat, high temperatures, flames, static discharge, sparks and other ignition sources, confined spaces, moisture and incompatibles.
Incompatible Materials	Acids (mineral acids, such as sulfuric acid, or organic acids), acid anhydrides, acid halides, alkali metals (e.g. sodium or potassium), alkaline earth metals, metals (such as metallic powdered aluminium, powdered magnesium and zinc), reducing agents, some forms of plastics, rubber, and coatings, oxidizing agents (such as perchloric acid, metal perchlorates, salts of oxyhalogenic acids, bromine, chlorine, chromium trioxide, halogen oxides, nitrates, nitric acid, nitrogen oxides, nonmetallic oxides, chromosulfuric acid, sodium hypochlorite), hydrides, zinc diethyl, halogens. hydrogen peroxide, carbon tetrachloride and metals, acetyl bromide, dichloromethane, potassium tert-butoxide, alkylaluminium solutions, beryllium hydride, cyanuric chloride, isocyanates or phosphorus (III) oxide (tetraphosphorus hexaoxide), diethyl zinc.
Hazardous	Carbon monoxide, carbon dioxide and formaldehyde.
Decomposition Products	
Possibility of hazardous reactions	Can react vigorously with oxidizers. Violent reaction with alkyl aluminium salts, acetyl bromide, chloroform + sodium methoxide, chromic anhydride, cyanuirc chlorite, lead perchlorate, phosphorous trioxide, nitric acid. Exothermic reaction with sodium hydroxide + chloroform. Incompatible with beryllium dihydride, metals (potassium and magnesium), oxidants (barium perchlorate, bromine, sodium hypochlorite, chlorine, hydrogen peroxide), potassium tert-butoxide, carbon tetrachloride, alkali metals, metals (aluminium, potassium magnesium, zinc), and dichlormethane. Rapid autocatalytic dissolution of aluminium, magnesium or zinc in 9:1 methanol + carbon tetrachloride - sufficiently vigorous to be rated as potentially hazardous. May attack some plastics, rubber, and coatings.
Hazardous Polymerization	Will not occur.

11. Toxicological Information

Toxicology Information	This substance should be treated with great care.
Ingestion	Toxic if Swallowed. Effects are the same as those described for 'Inhalation'. There is a wide range of individual susceptibility to the toxic effects of methanol (from a fatal dose of 15 mL of 40% methanol, to survival following ingestion of 500 mL of the same solution). In general, 300 to 1000 mg/kg is considered the range of minimum lethal dose for untreated cases of methanol



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Inhalation	lungs) during i comparison to r potentially fat Ingestion is no Toxic if inhale and can very re	hanol can probably be easily aspir ingestion or vomiting, based on it related alcohols. Aspiration of me cal accumulation of fluid in the l bt a typical route of occupational ed. A slight irritant to the mucou eadily form extremely high vapour	es physical properties and ethanol could cause a ungs (pulmonary edema). exposure. as membranes. Methanol is toxic concentrations at room
	first, methanol symptoms such a appearance of d (typically 8-24 period is then	hhalation is the most common route causes mild central nervous syst as nausea, headache, vomiting, diz drunkenness. A time period with no hours, but may last several hour followed by development of metabo oms such as headache, dizziness, n	tem (CNS) depression with sziness, incoordination and an o obvious symptoms follows as to 2 days). This latent plic acidosis and severe visual
	in more severe breathing have failure, may oc include reduced double and/or s poisoning and t	cases by abdominal and muscular p been observed. Coma and death, us ccur if medical treatment is not r d reactivity and/or increased sens snowy vision, and blindness. Depen the promptness of treatment, survi nent blindness, vision disturbance	pain and difficult periodic sually due to respiratory received. Visual effects may sitivity to light, blurred, adding on the severity of wors may recover completely or
Skin	skin, based on located. Methyl	ntact with skin. Methanol may be m unconfirmed animal information. N L alcohol is a defatting agent and kin absorption can occur; symptoms	No human information was W may cause skin to become dry
Eye	H370 Causes dam	mage to organs, eyes.	
Respiratory sensitisation Skin Sensitisation		based on available information.	
Germ cell		based on available information.	
mutagenicity	Not classified	based on available information.	
Carcinogenicity	Not classified	the IARC Monographs. based on available information.	
Reproductive Toxicity	Not classified	based on available information.	
STOT-single exposure		: organ toxicity - Single Exposure mage to organs, eyes.	e Category 1, Eyes
STOT-repeated exposure	Not classified	based on available information.	
Chronic Effects	contact may cau those of acute body. Because o cumulative pois	ent of vision has been reported. P use dermatitis. Chronic exposure m exposure. Methanol is only very s of this slow elimination, methanol son. Though a single exposure may result in the accumulation of a ha	hay cause effects similar to slowly eliminated from the should be regarded as a cause no effect, daily
Serious eye damage/irritation Mutageniaity		based on available information.	
Mutagenicity	NOU CLASSIFIED	Dased on available information.	

12. Ecological information

Ecotoxicity	Harmful effect on aquatic organisms. Risk of formation of explosive vapours above water surface. When used properly, no impairments in the function of waste-water-treatment plants are to be expected.
Persistence and degradability	Abiotic degradation: Slow degradation. (air) Biologic degradation: BOD 76 % von TOD /5 d (closed bottle test). Readily biodegradable (reduction: DOC >70 %; BOD >60 %; BOD5 to COD >50 %). Degradability: BOD5: 0.60 - 1.12 g/g; COD: 1.42 g/g; TOD: 1.5 g/g.



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Mobility	Distribution: log P(o/w): -0.74.
Bioaccumulative Potential	No bioaccumulation is to be expected (log P(o/w <1).
Environmental Protection	Do not allow to enter waters, waste water, or soil!

13. Disposal considerations

Disposal	Whatever cannot be saved for recovery or recycling should be disposed of
Considerations	according to relevant local, state and federal government regulations.

14. Transport information

Transport Information	Dangerous Goods of Class 3 Flammable Liquids, are incompatible in a placard load with any of the following: - Class 1, Class 2.1, if both the Class 3 and Class 2.1, dangerous goods are in bulk, Class 2.3, Class 4.2, Class 5, Class 6, if the Class 3 dangerous goods are nitromethane and Class 7.
U.N. Number	1230
UN proper shipping name	METHANOL
Transport hazard class(es)	3
Sub.Risk	6.1
Hazchem Code	•2WE
Packing Group	II
EPG Number	3A3
IERG Number	16
Environmental Hazards	Harmful to aquatic organisms. Risk of formation of explosive vapours above water surface. When used properly, no impairments in the function of waste-water-treatment plants are to be expected.

15. Regulatory information

Regulatory Information	All of the significant ingredients in this formulation are compliant with Australian Industrial Chemicals Introduction Scheme (AICIS) regulations. Not listed under WHS Regulation 2011, Schedule 10 - Prohibited carcinogens, restricted carcinogens and restricted hazardous chemicals.
Poisons Schedule	S6

16. Other Information

Literature	'Standard for the Uniform Scheduling of Medicines and Poisons .', Commonwealth
References	of Australia. National Road Transport Commission, 'Australian Code for the Transport of Dangerous Goods by Road and Rail 7th. Ed.'.
	Safe Work Australia, 'National Code of Practice fot the Preparation of Safety Data Sheets for Hazardous Chemicals'.
	Standards Australia, 'SAA/SNZ HB 76:2010 Dangerous Goods - Initial Emergency Response Guide', Standards Australia/Standards New Zealand.
	Safe Work Australia, 'Hazardous Chemical Information System'.
	Safe Work Australia, 'National Code of Practice for the Labelling of Safe Work Hazardous Substances'.
	Safe Work Australia, 'National Exposure Standards for Atmospheric Contaminants in the Occupational Environment'.
Empirical Formula & Structural Formula	CH3OH - Methanol
	End Of MSDS
	Convision Charical Safety International Div Itd

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