MONARCH®

Roof & Gutter - White

Australian Brushware Corporation Pty Ltd

Chemwatch Hazard Alert Code: 2

Chemwatch: 5578-38 Version No: 3.1

Safety Data Sheet according to WHS Regulations (Hazardous Chemicals) Amendment 2020 and ADG requirements

Issue Date: **28/09/2023** Print Date: **13/10/2023**

S.GHS.AUS.EN.E

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

Product Identifier

Product name	Roof & Gutter - White	
Chemical Name	Not Applicable	
Synonyms	Monarch Roof & Gutter White 300g,9320090032875,MS-3287	
Chemical formula	Not Applicable	
Other means of identification	Not Available	

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

Relevant identified uses Ideal for all roofing & outdoor applications. Use according to manufacturer's directions.

Details of the manufacturer or supplier of the safety data sheet

Registered company name	Australian Brushware Corporation Pty Ltd	
Address	143-147 National Blvd Campbellfield VICTORIA 3061 Australia	
Telephone	+61 1800 666 078	
Fax	Not Available	
Website	monarchpainting.com	
Email	Not Available	

Emergency telephone number

· · · · · · · · · · · · · · · · · · ·	
Association / Organisation	Not Available
Emergency telephone numbers	Not Available
Other emergency telephone numbers	Not Available

SECTION 2 Hazards identification

Classification of the substance or mixture

Poisons Schedule	Not Applicable	
Classification [1]	Classification [1] Sensitisation (Skin) Category 1, Specific Target Organ Toxicity - Repeated Exposure Category 2	
Legend: 1. Classified by Chernwatch; 2. Classification drawn from HCIS; 3. Classification drawn from Regulation (EU) No 1272/200		

Label elements

Hazard pictogram(s)		
Signal word Warning		
Hazard statement(s)		
H317	May cause an allergic skin reaction.	
H373	May cause damage to organs through prolonged or repeated exposure.	

Precautionary	statement(s	Prevention
I recautionally	Statement(S	

, , ,		
P260	Do not breathe mist/vapours/spray.	
P280 Wear protective gloves and protective clothing.		
P272 Contaminated work clothing should not be allowed out of the workplace.		

Precautionary statement(s) Response

IF ON SKIN: Wash with plenty of water and soap.
IF ON SKIN: Wash with plenty

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P314	P314 Get medical advice/attention if you feel unwell.	
P333+P313 If skin irritation or rash occurs: Get medical advice/attention.		
P362+P364	Take off contaminated clothing and wash it before reuse.	

Precautionary statement(s) Storage

Not Applicable

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
2224-33-1	0-10 vinyltris(methylethylketoxime)silane	
34036-80-1	0-10	2-butanone-O.O'.O"-(phenylsilylidene)trioxime
Legend: 1. Classified by Chemwatch; 2. Classification drawn from HCIS; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 4. Classification drawn from C&L * EU IOELVs available		

SECTION 4 First aid measures

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

Indication of any immediate medical attention and special treatment needed

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
lvice for firefighters	
Fire Fighting	 Alert Fire Brigade and tell them location and nature of hazard. Wear breathing apparatus plus protective gloves. Prevent, by any means available, spillage from entering drains or water courses. Use water delivered as a fine spray to control fire and cool adjacent area. 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. Equipment should be thoroughly decontaminated after use.
Fire/Explosion Hazard	 Combustible. Slight fire hazard when exposed to heat or flame. Heating may cause expansion or decomposition leading to violent rupture of containers. On combustion, may emit toxic fumes of carbon monoxide (CO).

Roof & Gut	ter - White
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	carbon monoxide (CO) carbon dioxide (CO2) nitrogen oxides (NOx) silicon dioxide (SiO2) other pyrolysis products typical of burning organic material. Contains low boiling substance: Closed containers may rupture due to pressure buildup under fire conditions. May emit poisonous fumes. May emit corrosive fumes.
HAZCHEM	Not Applicable

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. Avoid contact with skin and eyes. Wear impervious gloves and safety goggles. Trowel up/scrape up. Place spilled material in clean, dry, sealed container. Flush spill area with water.
Major Spills	 Clear area of personnel and move upwind. Alert Fire Brigade and tell them location and nature of hazard. Wear breathing apparatus plus protective gloves. Prevent, by any means available, spillage from entering drains or water course. Stop leak if safe to do so. Contain spill with sand, earth or vermiculite. Collect recoverable product into labelled containers for recycling. Neutralise/decontaminate residue (see Section 13 for specific agent). Collect solid residues and seal in labelled drums for disposal. Wash area and prevent runoff into drains. After clean up operations, decontaminate and launder all protective clothing and equipment before storing and re-using. 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	 Contains low boiling substance: Storage in sealed containers may result in pressure buildup causing violent rupture of containers not rated appropriately. Check for bulging containers. Vent periodically Always release caps or seals slowly to ensure slow dissipation of vapours 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. DO NOT allow material to contact humans, exposed food or food utensils. Avoid contact with incompatible materials. When handling, DO NOT eat, drink or smoke. Keep containers securely sealed when not in use. Avoid physical damage to containers. Always wash hands with soap and water after handling. Work clothes should be laundered separately. Launder contaminated clothing before re-use. 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 are maintained. Store in original containers. Keep containers securely sealed.
Other information	 Store in a cool, dry, well-ventilated area. Store away from incompatible materials and foodstuff containers. Protect containers against physical damage and check regularly for leaks. Observe manufacturer's storage and handling recommendations contained within this SDS.

Conditions for safe storage, including any incompatibilities

Suitable container	 Metal can or drum Packaging as recommended by manufacturer. Check all containers are clearly labelled and free from leaks.
Storage incompatibility	 Avoid strong acids, bases. Avoid reaction with oxidising agents

SECTION 8 Exposure controls / personal protection

Control parameters

Occupational Exposure Limits (OEL)

Emergency Limits

Ingredient	TEEL-1	TEEL-2		TEEL-3
Roof & Gutter - White	Not Available	Not Available		Not Available
Ingredient	Original IDLH		Revised IDLH	
vinyltris(methylethylketoxime)silane	Not Available		Not Available	
2-butanone-O,O',O"- (phenylsilylidene)trioxime	Not Available		Not Available	
Occupational Exposure Banding				
Ingredient	Occupational Exposure Band Rating		Occupational Expo	sure Band Limit
vinyltris(methylethylketoxime)silane	D		> 0.1 to ≤ 1 ppm	
2-butanone-O,O',O"-	E		≤ 0.1 ppm	
(phenylsilylidene)trioxime				

	be highly effective in protecting workers and will typically be in The basic types of engineering controls are: Process controls which involve changing the way a job activit Enclosure and/or isolation of emission source which keeps a "adds" and "removes" air in the work environment. Ventilation ventilation system must match the particular process and che Employers may need to use multiple types of controls to prev Local exhaust ventilation usually required. If risk of overexpo protection. Supplied-air type respirator may be required in sp An approved self contained breathing apparatus (SCBA) may Provide adequate ventilation in warehouse or closed storage velocities which, in turn, determine the "capture velocities" of Type of Contaminant: solvent, vapours, degreasing etc., evaporating from tank (ii	ty or process is done to reduce the risk. selected hazard "physically" away from the worker and ven in can remove or dilute an air contaminant if designed proper emical or contaminant in use. vent employee overexposure. sure exists, wear approved respirator. Correct fit is essentia becial circumstances. Correct fit is essential to ensure adequ y be required in some situations. area. Air contaminants generated in the workplace possess fresh circulating air required to effectively remove the conta	of protection. tilation that strategically ty. The design of a I to obtain adequate ate protection. s varying "escape" iminant. Air Speed: 0.25-0.5 m/s	
	aerosols, fumes from pouring operations, intermittent conta		(50-100 f/min.) 0.5-1 m/s (100-200	
Appropriate engineering controls	drift, plating acid fumes, pickling (released at low velocity in		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.)	
	grinding, abrasive blasting, tumbling, high speed wheel gen very high rapid air motion).	2.5-10 m/s (500-2000 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	ure 1: Disturbing room air currents		
	2: Contaminants of low toxicity or of nuisance value only.	y. 2: Contaminants of high toxicity		
	3: Intermittent, low production.	3: High production, heavy use		
	 Large hood or large air mass in motion 	4: Small hood-local control only		
	Simple theory shows that air velocity fells repidly with distance		ty gonorally degrades	
	Simple theory shows that air velocity falls rapidly with distanc with the square of distance from the extraction point (in simpl accordingly, after reference to distance from the contaminatir 1-2 m/s (200-400 f/min) for extraction of solvents generated i producing performance deficits within the extraction apparatu more when extraction systems are installed or used.	the away from the opening of a simple extraction pipe. Veloci le cases). Therefore the air speed at the extraction point sho ng source. The air velocity at the extraction fan, for example in a tank 2 meters distant from the extraction point. Other mo	buld be adjusted, , should be a minimum of echanical considerations,	
Individual protection measures, such as personal protective equipment	with the square of distance from the extraction point (in simple accordingly, after reference to distance from the contaminating 1-2 m/s (200-400 f/min) for extraction of solvents generated in producing performance deficits within the extraction apparatu	the away from the opening of a simple extraction pipe. Veloci le cases). Therefore the air speed at the extraction point sho ng source. The air velocity at the extraction fan, for example in a tank 2 meters distant from the extraction point. Other mo	buld be adjusted, , should be a minimum of echanical considerations,	
measures, such as personal	 with the square of distance from the extraction point (in simple accordingly, after reference to distance from the contaminatin 1-2 m/s (200-400 f/min) for extraction of solvents generated i producing performance deficits within the extraction apparatumore when extraction systems are installed or used. Image: A strand or a strain of the strand or used or used or used of the strand or used or used of the strand or used or	the away from the opening of a simple extraction pipe. Veloci le cases). Therefore the air speed at the extraction point sho ng source. The air velocity at the extraction fan, for example in a tank 2 meters distant from the extraction point. Other mu us, make it essential that theoretical air velocities are multiple	v document, describing iew of lens absorption is should be trained in ttion immediately and ens should be removed in	
measures, such as personal protective equipment	 with the square of distance from the extraction point (in simple accordingly, after reference to distance from the contaminatin 1-2 m/s (200-400 f/min) for extraction of solvents generated i producing performance deficits within the extraction apparatumore when extraction systems are installed or used. Image: A strand or a strain of the strand or used or used or used of the strand or used or used of the strand or used or	ee away from the opening of a simple extraction pipe. Veloci le cases). Therefore the air speed at the extraction point sho g source. The air velocity at the extraction fan, for example in a tank 2 meters distant from the extraction point. Other mu is, make it essential that theoretical air velocities are multipl of the extraction point. Other mu is, make it essential that theoretical air velocities are multipl of the extraction point. Other mu is, make it essential that theoretical air velocities are multipl of the extraction point. Other mu is, make it essential that theoretical air velocities are multipl of the extraction point. Other mu is, make it essential that theoretical air velocities are multipl extraction of the extraction of the	v document, describing iew of lens absorption is should be trained in ttion immediately and ens should be removed in	
measures, such as personal protective equipment	 with the square of distance from the extraction point (in simple accordingly, after reference to distance from the contaminatin 1-2 m/s (200-400 f/min) for extraction of solvents generated i producing performance deficits within the extraction apparatumore when extraction systems are installed or used. Image: A start of the extraction of solvents generated is producing performance deficits within the extraction apparatumore when extraction systems are installed or used. Safety glasses with side shields. Chemical goggles. [AS/NZS 1337.1, EN166 or national extra dispersion of the class of chemicals in use and an atheir removal and suitable equipment should be readily a remove contact lens as soon as practicable. Lens should bar a clean environment only after workers have washed hat See Hand protection below Wear chemical protective gloves, e.g. PVC. Wear safety footwear or safety gumboots, e.g. Rubber NOTE: 	ee away from the opening of a simple extraction pipe. Veloci le cases). Therefore the air speed at the extraction point sho g source. The air velocity at the extraction fan, for example in a tank 2 meters distant from the extraction point. Other mus, make it essential that theoretical air velocities are multipl with the extraction point. Other must are unable in the extraction point. Other must make it essential that theoretical air velocities are multipl with the extraction point. Other must are unable. In the extraction point and concentrate irritants. A written policy reated for each workplace or task. This should include a revi- account of injury experience. Medical and first-aid personnel available. In the event of chemical exposure, begin eye irriga d be removed at the first signs of eye redness or irritation - le inds thoroughly. [CDC NIOSH Current Intelligence Bulletin 59] seed individuals. Care must be taken, when removing gloves	v document, describing events absorption extension of the section	
measures, such as personal protective equipment Eye and face protection Skin protection	 with the square of distance from the extraction point (in simple accordingly, after reference to distance from the contamination 1-2 m/s (200-400 f/min) for extraction of solvents generated in producing performance deficits within the extraction apparatumore when extraction systems are installed or used. Image: Safety glasses with side shields. Chemical goggles. [AS/NZS 1337.1, EN166 or national extraction for the class of chemicals in use and and adsorption for the class of chemicals in use and and adsorption for the class of chemicals in use and and a clean environment only after workers have washed hard 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 predispose equipment, to avoid all possible skin contact. 	ee away from the opening of a simple extraction pipe. Veloci le cases). Therefore the air speed at the extraction point sho g source. The air velocity at the extraction fan, for example in a tank 2 meters distant from the extraction point. Other mus, make it essential that theoretical air velocities are multipl with the extraction point. Other must are unable in the extraction point. Other must make it essential that theoretical air velocities are multipl with the extraction point. Other must are unable. In the extraction point and concentrate irritants. A written policy reated for each workplace or task. This should include a revi- account of injury experience. Medical and first-aid personnel available. In the event of chemical exposure, begin eye irriga d be removed at the first signs of eye redness or irritation - le inds thoroughly. [CDC NIOSH Current Intelligence Bulletin 59] seed individuals. Care must be taken, when removing gloves	v document, describing events absorption e document, describing events absorption should be trained in tition immediately and ens should be removed in e).	

Eye wash unit.

Respiratory protection

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

Selection of the Class and Type of respirator will depend upon the level of breathing zone contaminant and the chemical nature of the contaminant. Protection Factors (defined as the ratio of contaminant outside and inside the mask) may also be important.

Required minimum protection factor	Maximum gas/vapour concentration present in air p.p.m. (by volume)	Half-face Respirator	Full-Face Respirator
up to 10	1000	A-AUS / Class1	-
up to 50	1000	-	A-AUS / Class 1
up to 50	5000	Airline *	-
up to 100	5000	-	A-2
up to 100	10000	-	A-3
100+			Airline**

* - Continuous Flow ** - Continuous-flow or positive pressure demand

A(AII 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.

SECTION 9 Physical and chemical properties

Information on basic physical and chemical properties

Appearance	White paste with slight odour, immiscible in water.		
Physical state	Non Slump Paste	Relative density (Water = 1)	0.93-1.13
Odour	Slight	Partition coefficient n-octanol / water	Not Available
Odour threshold	Not Available	Auto-ignition temperature (°C)	Not Available
pH (as supplied)	Not Applicable	Decomposition temperature (°C)	Not Available
Melting point / freezing point (°C)	Not Available	Viscosity (cSt)	Not Available
Initial boiling point and boiling range (°C)	>35	Molecular weight (g/mol)	Not Applicable
Flash point (°C)	>=93	Taste	Not Available
Evaporation rate	Not Available	Explosive properties	Not Available
Flammability	Not Applicable	Oxidising properties	Not Available
Upper Explosive Limit (%)	Not Applicable	Surface Tension (dyn/cm or mN/m)	Not Available
Lower Explosive Limit (%)	Not Applicable	Volatile Component (%vol)	Not Available
Vapour pressure (kPa)	Not Available	Gas group	Not Available
Solubility in water	Immiscible	pH as a solution (1%)	Not Applicable
Vapour density (Air = 1)	Not Available	VOC g/L	<30

SECTION 10 Stability and reactivity

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

SECTION 11 Toxicological information

 Information on toxicological effects

 Information on toxicological effects

 Inhaltion of vapours or aerosols (mists, fumes), generated by the material during the course of normal handling, may be damaging to the health of the individual.

 There is some evidence to suggest that the material can cause respiratory irritation in some persons. The body's response to such irritation can cause further lung damage.

 Material is highly volatile and may quickly form a concentrated atmosphere in confined or unventilated areas. The vapour may displace and replace air in breathing zone, acting as a simple asphyxiant. This may happen with little warning of overexposure.

 Ingestion
 Accidental ingestion of the material may be damaging to the health of the individual.

[•] 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

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Skin Contact	The material is not thought to produce adverse health effects or skin irritation following contact (as classified by EC Directives using animal models). Nevertheless, good hygiene practice requires that exposure be kept to a minimum and that suitable gloves be used in an occupational setting. 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	If applied to the eyes, this material causes severe eye damage.
Chronic	Repeated or long-term occupational exposure is likely to produce cumulative health effects involving organs or biochemical systems. Skin contact with the material is more likely to cause a sensitisation reaction in some persons compared to the general population. 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. There has been concern that this material can cause cancer or mutations, but there is not enough data to make an assessment.

	TOXICITY	IRRITATION	
Roof & Gutter - White	Not Available	Not Available	
	TOXICITY	IRRITATION	
vinyltris(methylethylketoxime)silane	dermal (rat) LD50: >2009 mg/kg ^[1]	Eye: adverse effect observed (irritating) ^[1]	
	Oral (Rat) LD50: >2000 mg/kg ^[1]	Skin: no adverse effect observed (not irritating) ^[1]	
	ΤΟΧΙΟΙΤΥ	IRRITATION	
	TUNICITY	INNITATION	
2-butanone-0,0',0"-	dermal (rat) LD50: >2000 mg/kg ^[1]	Not Available	
2-butanone-0,0',0"- (phenylsilylidene)trioxime			
(phenylsilylidene)trioxime Legend: 1. V	dermal (rat) LD50: >2000 mg/kg ^[1]	Not Available ity 2. Value obtained from manufacturer's SDS. Unless otherwise	

VINYLTRIS(METHYLETHYLKETOXIME)SILANE No significant acute toxicological data The material may cause skin irritation the production of vesicles, scaling and			n after prolonged or repeated exposur	e and may produce on contact skin redness, swelling,
2-BUTANONE-O,O',O"- (PHENYLSILYLIDENE)TRIOXIME * Sibond SDS				
VINYLTRIS(METHYLETHYLKETO & 2-BUTAN (PHENYLSILYLIDE	ONE-0,0',0"-	The following information refers to contact allergens as a group and may not be specific to this product. Contact allergies quickly manifest themselves as contact eczema, more rarely as urticaria or Quincke's oedema. The pathogenesis of contact eczema involves a cell-mediated (T lymphocytes) immune reaction of the delayed type. Other allergic skin reactions, e.g. contact urticaria, involve antibody-mediated immune reactions. The significance of the contact allergen is r simply determined by its sensitisation potential: the distribution of the substance and the opportunities for contact with it are equally important. A weakly sensitising substance which is widely distributed can be a more important allergen than one with stronger sensitising potential with which few individuals come into contact. From a clinical point of view, substances are potential they produce an allergic test reaction in more than 1% of the persons tested		rely as urticaria or Quincke's oedema. The immune reaction of the delayed type. Other allergic actions. The significance of the contact allergen is not tance and the opportunities for contact with it are ed can be a more important allergen than one with From a clinical point of view, substances are persons tested. of prohaptens. Three putative metabolites were beta-epoxy oximes and a nitro analogue. When ctivity of Skin Sensitizers.
Acute Toxicity	×		Carcinogenicity	×
Skin Irritation/Corrosion	×		Reproductivity	×
Serious Eye Damage/Irritation	age/Irritation 🗙		STOT - Single Exposure	×
Respiratory or Skin sensitisation			STOT - Repeated Exposure	✓
Mutagenicity	×		Aspiration Hazard	×

Legend:

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

SECTION 12 Ecological information

oxicity					
	Endpoint	Test Duration (hr)	Species	Value	Source
Roof & Gutter - White	Not Available	Not Available	Not Available	Not Available	Not Available
	Endpoint	Test Duration (hr)	Species	Value	Source
vinyltris(methylethylketoxime)silane	EC50	72h	Algae or other aquatic plants	6.1mg/l	2
	EC50	48h	Crustacea	201mg/l	2
	NOEC(ECx)	72h	Algae or other aquatic plants	1mg/l	2
	LC50	96h	Fish	>100mg/l	2
	Endpoint	Test Duration (hr)	Species	Value	Source
	EC50	72h	Algae or other aquatic plants	13.8mg/l	2
2-butanone-O,O',O"- (phenylsilylidene)trioxime	EC50	48h	Crustacea	>101mg/l	Continued.
(prieryssiyiidene)(rioxiiie	LC50	96h	Fish	>89.8mg/l	2
					_

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Lege		Extracted from 1. IUCLID Toxicity Data 2. Europe ECHA Registered Substances - Ecotoxicological Information - Aquatic Toxicity 4. US EPA, Ecotox database - Aquatic Toxicity Data 5. ECETOC Aquatic Hazard Assessment Data 6. NITE (Japan) - Bioconcentration Data 7. METI (Japan) - Bioconcentration Data 8. Vendor Data				
DO NOT discharge into sewe	r or waterways.					
Persistence and degradat	ility					
Ingredient	Persistence: Water/Soil	Persistence: Air				
	No Data available for all ingredients	No Data available for all ingredients				

Bioaccumulative potent	ial
Ingredient	Bioaccumulation
	No Data available for all ingredients
Mobility in soil	
Ingredient	Mobility
	No Data available for all ingredients

SECTION 13 Disposal considerations

Waste treatment methods	Containers may still present a chemical hazard/ danger when empty.
Product / Packaging disposal	 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. Recycle wherever possible or consult manufacturer for recycling options. Consult State Land Waste Authority for disposal. Bury or incinerate residue at an approved site. Recycle containers if possible, or dispose of in an authorised landfill.

SECTION 14 Transport information

Labels Required	
Marine Pollutant	NO
HAZCHEM	Not Applicable

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

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

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

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

Not Applicable

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

Product name	Group
vinyltris(methylethylketoxime)silane	Not Available
2-butanone-O,O',O"- (phenylsilylidene)trioxime	Not Available

14.7.3. Transport in bulk in accordance with the IGC Code

Product name	Ship Type
vinyltris(methylethylketoxime)silane	Not Available
2-butanone-O,O',O"- (phenylsilylidene)trioxime	Not Available

SECTION 15 Regulatory information

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

vinyltris(methylethylketoxime)silane is found on the following regulatory lists

Australian Inventory of Industrial Chemicals (AIIC)

2-butanone-O,O',O"-(phenylsilylidene)trioxime is found on the following regulatory lists

Australia Hazardous Chemical Information System (HCIS) - Hazardous Chemicals

Australian	Inventory	of	Industrial	Chemicals	(AIIC)	۱
Australian	memory	UI.	muusinai	Chemicais		,

National Inventory Status

National Inventory	Status
Australia - AIIC / Australia Non-Industrial Use	Yes Continued
Canada - DSL	Yes

National Inventory	Status
China - IECSC	Yes
Europe - EINEC / ELINCS / NLP	Yes
Japan - ENCS	Yes
Korea - KECI	Yes
New Zealand - NZIoC	Yes
Philippines - PICCS	No (2-butanone-O,O',O"-(phenylsilylidene)trioxime)
USA - TSCA	Yes
Taiwan - TCSI	Yes
Mexico - INSQ	No (vinyltris(methylethylketoxime)silane; 2-butanone-O,O',O"-(phenylsilylidene)trioxime)
Vietnam - NCI	Yes
Russia - FBEPH	No (2-butanone-O,O',O"-(phenylsilylidene)trioxime)
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	28/09/2023
Initial Date	17/11/2022

SDS Version Summary

Version	Date of Update	Sections Updated
3.1	01/03/2023	Physical and chemical properties - Appearance, Identification of the substance / mixture and of the company / undertaking - Synonyms, Name

Other information

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

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

Definitions and abbreviations

PC - TWA: Permissible Concentration-Time Weighted Average PC - STEL: Permissible Concentration-Short Term Exposure Limit IARC: International Agency for Research on Cancer ACGIH: American Conference of Governmental Industrial Hygienists STEL: Short Term Exposure Limit TEEL: Temporary Emergency Exposure Limit. IDLH: Immediately Dangerous to Life or Health Concentrations ES: Exposure Standard OSF: Odour Safety Factor NOAEL :No Observed Adverse Effect Level LOAEL: Lowest Observed Adverse Effect Level TLV: Threshold Limit Value LOD: Limit Of Detection OTV: Odour Threshold Value BCF: BioConcentration Factors **BEI: Biological Exposure Index** AIIC: Australian Inventory of Industrial Chemicals DSL: Domestic Substances List NDSL: Non-Domestic Substances List IECSC: Inventory of Existing Chemical Substance in China EINECS: European INventory of Existing Commercial chemical Substances ELINCS: European List of Notified Chemical Substances NLP: No-Longer Polymers ENCS: Existing and New Chemical Substances Inventory KECI: Korea Existing Chemicals Inventory NZIoC: New Zealand Inventory of Chemicals PICCS: Philippine Inventory of Chemicals and Chemical Substances TSCA: Toxic Substances Control Act TCSI: Taiwan Chemical Substance Inventory INSQ: Inventario Nacional de Sustancias Químicas NCI: National Chemical Inventory FBEPH: Russian Register of Potentially Hazardous Chemical and Biological Substances

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