

# **ARDEX E25**

## Ardex (Ardex NZ)

Chemwatch: 7933-92

Version No: 2.1 Safety Data Sheet according to the Health and Safety at Work (Hazardous Substances) Regulations 2017 Chemwatch Hazard Alert Code: 2

Issue Date: **31/01/2025** Print Date: **04/02/2025** L.GHS.NZL.EN.E

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

## **Product Identifier**

Product name	ARDEX E25
Chemical Name	Not Applicable
Synonyms	mortar levelling compounds admixture; Ardex 250
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	Admixture for improving mortars and levelling compounds for high loads.
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## Details of the manufacturer or supplier of the safety data sheet

Registered company name	Ardex (Ardex NZ)
Address	32 Lane Street Woolston Christchurch New Zealand
Telephone	+64 3384 3029 +64 3384 9779
Fax	+64 3384 9779
Website	www.ardex.co.nz
Email	info@ardexnz.com

#### Emergency telephone number

	Association / Organisation	Ardex (Ardex NZ)
	Emergency telephone number(s)	+64 3 373 6900
	Other emergency telephone number(s)	0800 764 766 (NZ NPC)

## **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. Not regulated for transport of Dangerous Goods.

Classification <sup>[1]</sup>	Sensitisation (Skin) Category 1
Legend:	1. Classified by Chernwatch; 2. Classification drawn from CCID EPA NZ; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI
Determined by Chemwatch using GHS/HSNO criteria	6.5B (contact)
abel elements	
Hazard pictogram(s)	
Signal word	Warning

May cause an allergic skin reaction.

H317

## Precautionary statement(s) Prevention

P280	Wear protective gloves and protective clothing.
P261	Avoid breathing mist/vapours/spray.
P272	Contaminated work clothing should not be allowed out of the workplace.

## Precautionary statement(s) Response

P302+P352	IF ON SKIN: Wash with plenty of water.
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

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
55965-84-9	<0.002	5-chloro-2-methyl-4-isothiazolin-3-one
Not Available	balance	Ingredients determined not to be hazardous
Legend: 1. Classified by Chemwatch; 2. Classification drawn from CCID EPA NZ; 3. Classification drawn from Regulation (EU) No 12; VI; 4. Classification drawn from C&L * EU IOELVs available		

## **SECTION 4 First aid measures**

## Description of first aid measures

Eye Contact	<ul> <li>If this product comes in contact with eyes:</li> <li>Wash out immediately with water.</li> <li>If irritation continues, seek medical attention.</li> <li>Removal of contact lenses after an eye injury should only be undertaken by skilled personnel.</li> </ul>
Skin Contact	<ul> <li>If skin contact occurs:</li> <li>Immediately remove all contaminated clothing, including footwear.</li> <li>Flush skin and hair with running water (and soap if available).</li> <li>Seek medical attention in event of irritation.</li> </ul>
Inhalation	<ul> <li>If fumes, aerosols or combustion products are inhaled remove from contaminated area.</li> <li>Other measures are usually unnecessary.</li> </ul>
Ingestion	<ul> <li>Immediately give a glass of water.</li> <li>First aid is not generally required. If in doubt, contact a Poisons Information Centre or a doctor.</li> </ul>

## Indication of any immediate medical attention and special treatment needed

Treat symptomatically.

## **SECTION 5 Firefighting measures**

## Extinguishing media

There is no restriction on the type of extinguisher which may be used.

• Use extinguishing media suitable for surrounding area.

## Special hazards arising from the substrate or mixture

Fire Incompatibility	None known.
Advice for firefighters	
	<ul> <li>Alert Fire Brigade and tell them location and nature of hazard.</li> </ul>

Fire Fighting	<ul> <li>Wear breathing apparatus plus protective gloves in the event of a fire.</li> <li>Prevent, by any means available, spillage from entering drains or water courses.</li> <li>Use fire fighting procedures suitable for surrounding area.</li> <li>DO NOT approach containers suspected to be hot.</li> <li>Cool fire exposed containers with water spray from a protected location.</li> <li>If safe to do so, remove containers from path of fire.</li> <li>Equipment should be thoroughly decontaminated after use.</li> </ul>
Fire/Explosion Hazard	<ul> <li>Non combustible.</li> <li>Not considered a significant fire risk, however containers may burn.</li> </ul>

## **SECTION 6 Accidental release measures**

Personal precautions, protective equipment and emergency procedures

May emit corrosive fumes.

See section 8

## **Environmental precautions**

See section 12

## Methods and material for containment and cleaning up

Minor Spills	<ul> <li>Clean up all spills immediately.</li> <li>Avoid breathing vapours and contact with skin and eyes.</li> <li>Control personal contact with the substance, by using protective equipment.</li> <li>Contain and absorb spill with sand, earth, inert material or vermiculite.</li> <li>Wipe up.</li> <li>Place in a suitable, labelled container for waste disposal.</li> </ul>
Major Spills	<ul> <li>Moderate hazard.</li> <li>Clear area of personnel and move upwind.</li> <li>Alert Fire Brigade and tell them location and nature of hazard.</li> <li>Wear breathing apparatus plus protective gloves.</li> <li>Prevent, by any means available, spillage from entering drains or water course.</li> <li>Stop leak if safe to do so.</li> <li>Contain spill with sand, earth or vermiculite.</li> <li>Collect recoverable product into labelled containers for recycling.</li> <li>Neutralise/decontaminate residue (see Section 13 for specific agent).</li> <li>Collect solid residues and seal in labelled drums for disposal.</li> <li>Wash area and prevent runoff into drains.</li> <li>After clean up operations, decontaminate and launder all protective clothing and equipment before storing and re-using.</li> <li>If contamination of drains or waterways occurs, advise emergency services.</li> </ul>

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

## SECTION 7 Handling and storage

## Precautions for safe handling

Safe handling	<ul> <li>Avoid all personal contact, including inhalation.</li> <li>Wear protective clothing when risk of exposure occurs.</li> <li>Use in a well-ventilated area.</li> <li>Avoid contact with moisture.</li> <li>Avoid contact with incompatible materials.</li> <li>When handling, DO NOT eat, drink or smoke.</li> <li>Keep containers securely sealed when not in use.</li> <li>Avoid physical damage to containers.</li> <li>Always wash hands with scap and water after handling.</li> <li>Work clothes should be laundered separately. Launder contaminated clothing before re-use.</li> <li>Use good occupational work practice.</li> <li>Observe manufacturer's storage and handling recommendations contained within this SDS.</li> <li>Atmosphere should be regularly checked against established exposure standards to ensure safe working conditions are maintained.</li> <li>DO NOT allow clothing wet with material to stay in contact with skin</li> </ul>
Other information	<ul> <li>Store in original containers.</li> <li>Keep containers securely sealed.</li> <li>Store in a cool, dry, well-ventilated area.</li> <li>Store away from incompatible materials and foodstuff containers.</li> <li>Protect containers against physical damage and check regularly for leaks.</li> <li>Observe manufacturer's storage and handling recommendations contained within this SDS.</li> </ul>

## Conditions for safe storage, including any incompatibilities

Suitable container	<ul> <li>Polyethylene or polypropylene container.</li> <li>Packing as recommended by manufacturer.</li> <li>Check all containers are clearly labelled and free from leaks.</li> </ul>
Storage incompatibility	None known

# SECTION 8 Exposure controls / personal protection

## **Control parameters**

## Occupational Exposure Limits (OEL)

## INGREDIENT DATA

Not Available		
Ingredient	Original IDLH	Revised IDLH
5-chloro-2-methyl-4- isothiazolin-3-one	Not Available	Not Available

## MATERIAL DATA

## Exposure controls

Appropriate engineering controls	Engineering controls are used to remove a hazard or place a barrier between the worker and the hazard. Well-designed engineering control 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. General exhaust is adequate under normal operating conditions. If risk of overexposure exists, wear SAA approved respirator. Correct fit is essential to obtain adequate protection. Provide adequate ventilation in warehouse or closed storage areas. Air contaminants generated in the workplace possess varying "escape" velocities which, in turn, determine the "capture velocities" of fresh circulating air required to effectively remove the contaminant.

Air Speed: 0.25-0.5 m/s (50-100 f/min) 0.5-1 m/s (100-

200 f/min.)

1-2.5 m/s (200-500 f/min) 2.5-10 m/s (500-

2000 f/min.)

	<u></u>	
Type of Contaminant:		
solvent, vapours, degreasing etc., evaporating from tank (i	n still air)	
aerosols, fumes from pouring operations, intermittent conta spray drift, plating acid fumes, pickling (released at low vel	<u>,</u>	fers, welding,
direct spray, spray painting in shallow booths, drum filling,	conveyer loading, crusher dusts, gas	discharge (active
generation into zone of rapid air motion)		
	nerated dusts (released at high initial	velocity into zone
generation into zone of rapid air motion) grinding, abrasive blasting, tumbling, high speed wheel ge of very high rapid air motion).	nerated dusts (released at high initial Upper end of the range	velocity into zone
generation into zone of rapid air motion) grinding, abrasive blasting, tumbling, high speed wheel ge of very high rapid air motion). Within each range the appropriate value depends on:		velocity into zone
generation into zone of rapid air motion) grinding, abrasive blasting, tumbling, high speed wheel ge of very high rapid air motion). Within each range the appropriate value depends on: Lower end of the range	Upper end of the range	velocity into zone
generation into zone of rapid air motion) grinding, abrasive blasting, tumbling, high speed wheel ge of very high rapid air motion). Within each range the appropriate value depends on: Lower end of the range 1: Room air currents minimal or favourable to capture	Upper end of the range 1: Disturbing room air currents	velocity into zone

Simple theory shows that air velocity falls rapidly with distance away from the opening of a simple extraction pipe. Velocity generally decreases with the square of distance from the extraction point (in simple cases). Therefore the air speed at the extraction point should be adjusted, accordingly, after reference to distance from the contaminating source. The air velocity at the extraction fan, for example, should be a minimum of 1-2 m/s (200-400 f/min.) for extraction of solvents generated in a tank 2 meters distant from the extraction point. Other mechanical considerations, producing performance deficits within the extraction apparatus, make it essential that theoretical air velocities are multiplied by factors of 10 or more when extraction systems are installed or used.

Individual protection measures, such as personal protective equipment

protective equipment	
Eye and face protection	<ul> <li>Safety glasses with side shields.</li> <li>Chemical goggles. [AS/NZS 1337.1, EN166 or national equivalent]</li> <li>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].</li> </ul>
Skin protection	See Hand protection below
Hands/feet protection	<ul> <li>Wear chemical protective gloves, e.g. PVC.</li> <li>Wear stelly lockwar or safety gumbots, e.g. Rubber</li> <li>Note:</li> <li>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.</li> <li>Contaminated leather items, such as shoes, bells and watch-bands should be removed and destroyed.</li> <li>The selection of suitable gloves does not only depend on the material, but also on further marks of quality which vary from manufacturer to manufacturer. Where the chemical is a preparation of several substances, the resistance of the glove material can not be calculated in advance and has therefore to be checked prior to the application.</li> <li>The exact break through time for substances has to be obtained from the manufacturer of the protective gloves and has to be observed where making a final choice.</li> <li>Personal hygiene is a key element of effective hand care. Gloves must only be worn on clean hands. After using gloves, hands should be washed and dried thoroughly. Application of a non-perfumed moisturiser is recommended.</li> <li>Suitability and durability of glove type is dependent on usage. Important factors in the selection of gloves include: <ul> <li>requency and duration of contact,</li> <li>elonemical resistance of glove material,</li> <li>glove thickness and</li> </ul> </li> <li>destribity</li> <li>Select gloves tested to a relevant standard (e.g. Europe EN 374, US F739, AS/NZS 2161.1 or national equivalent).</li> <li>When only brief contact is expected, a glove with a protection class of 5 or higher (breakthrough time greater than 240 minutes according to EN 374, AS/NZS 2161.10.1 or national equivalent) is recommended.</li> <li>Sorta gloves should be replaced.</li> <li>Sorta gloves should be replaced.&lt;</li></ul>
Body protection	See Other protection below
Other protection	► Overalls.

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P.V.C apron.Barrier cream.

Skin cleansing cream.

Eye wash unit.

## **SECTION 9** Physical and chemical properties

## Information on basic physical and chemical properties

Appearance	Pink liquid with a weak characteristic odour; mixes with water.		
Physical state	Liquid	Relative density (Water = 1)	Not Available
Odour	Not Available	Partition coefficient n-octanol / water	Not Available
Odour threshold	Not Available	Auto-ignition temperature (°C)	Not Applicable
pH (as supplied)	4-5	Decomposition temperature (°C)	Not Available
Melting point / freezing point (°C)	Not Available	Viscosity (cSt)	Not Available
Initial boiling point and boiling range (°C)	Not Available	Molecular weight (g/mol)	Not Applicable
Flash point (°C)	Not Applicable	Taste	Not Available
Evaporation rate	Not Applicable	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	Miscible	pH as a solution (1%)	Not Available
Vapour density (Air = 1)	Not Available	VOC g/L	Not Available
Heat of Combustion (kJ/g)	Not Available	Ignition Distance (cm)	Not Available
Flame Height (cm)	Not Available	Flame Duration (s)	Not Available
Enclosed Space Ignition Time Equivalent (s/m3)	Not Available	Enclosed Space Ignition Deflagration Density (g/m3)	Not Available

## **SECTION 10 Stability and reactivity**

Reactivity	See section 7
Chemical stability	<ul> <li>Unstable in the presence of incompatible materials.</li> <li>Product is considered stable.</li> <li>Hazardous polymerisation will not occur.</li> </ul>
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

sed on available data, the classification criteria are not met.
sed on available data, the classification criteria are not met.
sed on available data, the classification criteria are not met.
ere is sufficient evidence to classify this material as sensitising to skin or the respiratory system
sed on available data, the classification criteria are not met.
sed on available data, the classification criteria are not met.
sed on available data, the classification criteria are not met.
sed on available data, the classification criteria are not met.
sed on available data, the classification criteria are not met.
sed on available data, the classification criteria are not met.
e material is not thought to produce adverse health effects or irritation of the respiratory tract (as classified by EC Directives using animal dels). Nevertheless, good hygiene practice requires that exposure be kept to a minimum and that suitable control measures be used in an upational setting.
e material has <b>NOT</b> been classified by EC Directives or other classification systems as "harmful by ingestion". This is because of the lack corroborating animal or human evidence. The material may still be damaging to the health of the individual, following ingestion, especially ere pre-existing organ (e.g liver, kidney) damage is evident. Present definitions of harmful or toxic substances are generally based on ses producing mortality rather than those producing morbidity (disease, ill-health). Gastrointestinal tract discomfort may produce nausea I vomiting. In an occupational setting however, ingestion of insignificant quantities is not thought to be cause for concern.

Skin Contact		or skin irritation following contact (as classified by EC Directives using animal sposure be kept to a minimum and that suitable gloves be used in an	
Eye	Although the liquid is not thought to be an irritant (as classified by EC Directives), direct contact with the eye may produce transient discomfort characterised by tearing or conjunctival redness (as with windburn).		
Chronic	individuals, and/or of producing a positive response in experin Substances that can cause occupational asthma (also known airway hyper-responsiveness via an immunological, irritant or exposure to the substance, sometimes even to tiny quantities a runny nose to asthma. Not all workers who are exposed to a advance who are likely to become hyper-responsive. Substances than can cuase occupational asthma should be d people with pre-existing air-way hyper-responsiveness. The la Wherever it is reasonably practicable, exposure to substance possible the primary aim is to apply adequate standards of co Activities giving rise to short-term peak concentrations should surveillance is appropriate for all employees exposed or liable	as asthmagens and respiratory sensitisers) can induce a state of specific other mechanism. Once the airways have become hyper-responsive, further , may cause respiratory symptoms. These symptoms can range in severity from a sensitiser will become hyper-responsive and it is impossible to identify in istinguished from substances which may trigger the symptoms of asthma in atter substances are not classified as asthmagens or respiratory sensitisers s that can cuase occupational asthma should be prevented. Where this is not	
	ΤΟΧΙΟΙΤΥ	IRRITATION	
ARDEX E25	Not Available	Not Available	
	dermal (rat) LD50: >1008 mg/kg <sup>[2]</sup>	Eye: adverse effect observed (irreversible damage) <sup>[1]</sup>	
5-chloro-2-methyl-4-	Inhalation (Rat) LC50: 1.23 mg/l4h <sup>[2]</sup>	Skin (Human - woman): 0.01%	
isothiazolin-3-one	Oral (Rat) LD50: 53 mg/kg <sup>[2]</sup>	Skin (Human): 0.01% - Severe	
		Skin (Human): 0.1%/48H	
		Skin: adverse effect observed (corrosive) <sup>[1]</sup>	
		Skin: adverse effect observed (irritating) <sup>[1]</sup>	
	potential: the distribution of the substance and the opportuniti which is widely distributed can be a more important allergen t contact. From a clinical point of view, substances are notewor tested. Asthma-like symptoms may continue for months or even year condition known as reactive airways dysfunction syndrome (R compound. Main criteria for diagnosing RADS include the abs of persistent asthma-like symptoms within minutes to hours or include a reversible airflow pattern on lung function tests, mor and the lack of minimal lymphocytic inflammation, without eos disorder with rates related to the concentration of and duration is a disorder that occurs as a result of exposure due to high o reversible after exposure ceases. The disorder is characterizz Exposure to the material may result in a possible risk of irreve concern is raised, generally, on the basis of appropriate studies with similar materials using mammalian su in vitro mutagenicity studies. In light of potential adverse effects, and to ensure a harmonis biocides has been established with the objective of ensuring a this aim, it is required that risk assessment of biocidal product the risk assessment of the biocidal products are the utilization applications and thus the exposure of humans and the enviro Humans may be exposed to biocidal products in different way intended for industrial sectors or professional uses only, wher professional users. In addition, potential exposure of non-use environment, for example through drinking water, the food cha attention should be paid to the exposure of vulnerable sub-poo other domestic animals can be exposed indirectly following th in terms of route (inhalation, dermal contact, and ingestion) ar frequency and duration. No significant acute toxicological data identified in literature s The European Union has reclassified several formaldehyde-ref (MBO) and hydroxypropylamine (HPT) as category 1B carcin formaldehyde-releasing agents were not. This is no longer the theoretical concentration of releasable formaldehyde is	Presible effects. The material may produce mutagenic effects in man. This ormatic cells in vivo. Such findings are often supported by positive results from ed risk assessment and management, the EU regulatory framework for a high level of protection of human and animal health and the environment. To is is carried out before they can be placed on the market. A central element in instructions that defines the dosage, application method and amount of nment to the biocidal substance. Is in both occupational and domestic settings. Many biocidal products are eas other biocidal products are commonly available for private use by non-ris of biocidal products (i.e. the general public) may occur indirectly via the ain, as well as through atmospheric and residential exposure. Particular pulations, such as the elderly, pregnant women, and children. Also pets and e application of biocidal products. Furthermore, exposure to biocides may vary and pathway (food, drinking water, residential, occupational) of exposure, level, esarch. eleasing agents (FRAs) such as methylenedimorpholine (MBM), oxazolidine or case. Based on this regulation, formulations for which the maximum than > 1000 ppm (>0.1%), have to be labelled as carcinogenic.	
	potentially harmful microbes that could cause health problems A large proportion of bactericides on the market today are cla conditions they release small amounts of formaldehyde – this as a biocide their use may become restricted or unfavourable	ssed as formaldehyde releasing biocides which means that under specific is their mode of action in the presence of bacteria. Although they are effective	

	It has also been proposed by the ECHA Risk Assessin same as formaldehyde because formaldehyde is relea- interaction with microorganisms). Formaldehyde generators (releasers) are often used a generated following hydrolysis. The most widely used cell. Some release detectable levels of formaldehyde Many countries are placing regulatory pressure on su Formaldehyde generators are a diverse group of cher prepared by reacting an amino alcohol with formaldeh There is concern that when formaldehyde-releasing p triethanolamine (TEA), diethanolamine (DEA), or mon substances that can potentially penetrate skin. One widely-discussed hypothesis states that formaldeh imbalance in the microbial flora of in-use metalworking favours the proliferation of certain nontuberculosis my aerosols can cause hypersensitivity pneumonitis (HP) workers. Symptoms of HP include flu-like illness acco According to Annex VI of the Cosmetic Directive 76/76 ppm). In addition, the provisions of Annex VI state tha <i>All finished products containing formaldehyde or subs</i> <i>warning "contains formaldehyde"</i> where the concentrer Formaldehyde-releasing preservatives have the ability releasing preservatives ensures that the actual level of ensure absence of microbial growth. The formaldehyde electron-rich groups to disrupt metabolic processes, e The material may be irritating to the eye, with prolong produce conjunctivitis. The material may cause skin irritation after prolonged dermatitis is often characterised by skin redness (cryt spongy layer (spongiosis) and intracellular oederma of	ased when these substances come in as preservatives (antimicrobials, bioc antimicrobial compounds function by into the air space, above working sol ppliers and users to replace formalde micals that can be recognised by a sr nyde ("formaldehyde-condensates"), reservatives are present in a formula toethanolamine (MEA), nitrosamines ender (MWFs). The hypothesis furth robacteria (NTM) in MWFs and that a laso known as extrinsic allergic alve mpanied by chronic dyspnea, i.e., dif 68/EC, the maximum authorised con- t, stances in this Annex and which releat ation of formaldehyde in the finished j y to release formaldehyde in the groutes i fere formaldehyde in the groutes i de reacts most rapidly with organic are eventually causing death of the organ ed contact causing inflammation. Rep or repeated exposure and may prod hema) and swelling epidermis. Histol	The contact under favorable conditions (i.e. bides, microbiocides). Formaldehyde may be y releasing formaldehyde once inside the microbe lutions, especially when pH has dropped. shyde generators. mall, easily detachable formaldehyde moiety, tition that also includes amines, such as can be formed,; nitrosamines are carcinogenic triazines and oxazolidines, may cause an the asserts that this putative microbial imbalance the subsequent inhalation of NTM-containing eolitis, in a small percentage of susceptible ficult or laboured respiration centration of free formaldehyde is 0.2% (2000 ase formaldehyde must be labelled with the product exceeds 0.05%. Iall amounts over time. The use of formaldehyde- is always very low but at the same time sufficient to nd inorganic anions, amino and sulfide groups and ism. peated or prolonged exposure to irritants may uce a contact dermatitis (nonallergic). This form of
Acute Toxicity	×	Carcinogenicity	×
Skin Irritation/Corrosion	×	Reproductivity	×
Serious Eye Damage/Irritation	×	STOT - Single Exposure	×
Respiratory or Skin sensitisation	*	STOT - Repeated Exposure	×

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SECTION 12 Ecological information
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Mutagenicity

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## Toxicity

ARDEX E25	Endpoint	Test Duration (hr)	Species	Value	Source
	Not Available	Not Available	Not Available	Not Available	Not Available
	Endpoint	Test Duration (hr)	Species	Value	Source
5-chloro-2-methyl-4-	EC50	96h	Algae or other aquatic plants	0.03- 0.13mg/L	4
	EC50	72h	Algae or other aquatic plants	0.018- 0.026mg/L	4
isothiazolin-3-one	NOEC(ECx)	504h	Crustacea	0.172mg/l	1
	EC50	48h	Crustacea	4.71mg/l	1
	LC50	96h	Fish	0.13- 0.31mg/L	4
Legend:	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				

Legend:

Aspiration Hazard

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

## DO NOT discharge into sewer or waterways.

## Persistence and degradability

Ingredient	Persistence: Water/Soil	Persistence: Air
5-chloro-2-methyl-4- isothiazolin-3-one	HIGH	HIGH
Bioaccumulative potential		
Ingredient	Bioaccumulation	
5-chloro-2-methyl-4- isothiazolin-3-one	LOW (LogKOW = 0.0444)	
Mobility in soil		
Ingredient	Mobility	

5-chloro-2-methyl-4- isothiazolin-3-one	LOW (Log KOC = 45.15)

## **SECTION 13 Disposal considerations**

Waste treatment methods	
Product / Packaging disposal	<ul> <li>Containers may still present a chemical hazard/ danger when empty.</li> <li>Return to supplier for reuse/ recycling if possible.</li> <li>Otherwise: <ul> <li>If container can not be cleaned sufficiently well to ensure that residuals do not remain or if the container cannot be used to store the same product, then puncture containers, to prevent re-use, and bury at an authorised landfill.</li> <li>Where possible retain label warnings and SDS and observe all notices pertaining to the product.</li> <li>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.</li> <li>A Hierarchy of Controls seems to be common - the user should investigate:</li> <li>Reduction</li> <li>Reuse</li> <li>Recycling</li> <li>Disposal (if all else fails)</li> </ul> </li> <li>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 precycled 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. Shell 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.</li> <li><b>DO NOT allow wash water form cleaning or process equipment to enter drains.</b></li> <li>It may be necessary to collect all wash water for treatment before disposal.</li> <li>It may be necessary to collect all wash water for treatment before disposal.</li> <li>It may be necessary to collect all wash water for treatment before disposal.</li> <li>It may be necessary to collect all wash or to regional waste management authority for disposal if no suitable treatment or disposal facility can be identified.</li> <li>Obispose of by: burial in a land-</li></ul>

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.

Only dispose to the environment if a tolerable exposure limit has been set for the substance.

Only deposit the hazardous substance into or onto a landfill or sewage facility or incinerator, where the hazardous substance can be handled and treated appropriately.

## **SECTION 14 Transport information**

#### Labels Required

Marine Pollutant	NO
HAZCHEM	Not Applicable

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

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

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

## 14.7. Maritime transport in bulk according to IMO instruments

## 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
5-chloro-2-methyl-4- isothiazolin-3-one	Not Available

## 14.7.3. Transport in bulk in accordance with the IGC Code

Product name	Ship Type
5-chloro-2-methyl-4- isothiazolin-3-one	Not Available

#### **SECTION 15 Regulatory information**

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

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

HSR Number	Group Standard
HSR002503	Additives Process Chemicals and Raw Materials Subsidiary Hazard Group Standard 2020

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

#### 5-chloro-2-methyl-4-isothiazolin-3-one is found on the following regulatory lists

New Zealand Approved Hazardous Substances with controls

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

## Additional Regulatory Information

Not Applicable

## **Hazardous Substance Location**

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

Hazard Class	Quantities
Not Applicable	Not Applicable

## **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	Gas (aggregate water capacity in mL)	Liquid (L)	Solid (kg)	Maximum quantity per package for each classification
6.5A or 6.5B	120	1	3	

## Tracking Requirements

Not Applicable

#### **National Inventory Status**

National Inventory	Status
Australia - AIIC / Australia Non- Industrial Use	Yes
Canada - DSL	Yes
Canada - NDSL	No (5-chloro-2-methyl-4-isothiazolin-3-one)
China - IECSC	Yes
Europe - EINEC / ELINCS / NLP	Yes
Japan - ENCS	Yes
Korea - KECI	Yes
New Zealand - NZIoC	Yes
Philippines - PICCS	Yes
USA - TSCA	All chemical substances in this product have been designated as TSCA Inventory 'Active'
Taiwan - TCSI	Yes
Mexico - INSQ	Yes
Vietnam - NCI	Yes
Russia - FBEPH	Yes
Legend:	Yes = All CAS declared ingredients are on the inventory No = One or more of the CAS listed ingredients are not on the inventory. These ingredients may be exempt or will require registration.

## **SECTION 16 Other information**

Revision Date	31/01/2025
Initial Date	31/01/2025

#### 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

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- 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
   MARPOL: International Convention for the Prevention of Pollution from Ships
- IMSBC: International Maritime Solid Bulk Cargoes Code
- IGC: International Gas Carrier Code
- IBC: International Bulk Chemical Code
- 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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end of SDS