

SDI Limited

Version No: 7.1 Safety Data Sheet according to the Health and Safety at Work (Hazardous Substances) Regulations 2017 Issue Date: 23/12/2022 Print Date: 21/11/2023 L.GHS.NZL.EN

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

Product Identifier		
Product name	Pola Professional 37.5%	
Chemical Name	Not Applicable	
Synonyms	Not Available	
Proper shipping name	HYDROGEN PEROXIDE, AQUEOUS SOLUTION with not less than 20% but not more than 60% hydrogen peroxide (stabilized as necessary)	
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 Professional Dental use: To medically bleach endodontically treated teeth, to be performed by a dentist.

Details of the manufacturer or supplier of the safety data sheet

Registered company name	SDI Limited	SDI (North America) Inc.	SDI HOLDINGS PTY LTD DO
Address	3-15 Brunsdon Street Bayswater VIC 3153 Australia	1279 Hamilton Parkway Itasca IL 60143 United States	Rua Dr. Reinaldo Schmithausen 3141 – Cordeiros Itajaí – SC – CEP 88310-004 Brazil
Telephone	+61 3 8727 7111	+1 630 361 9200	+55 11 3092 7100
Fax	+61 3 8727 7222	Not Available	Not Available
Website	www.sdi.com.au	www.sdi.com.au	http://www.sdi.com.au/
Email	info@sdi.com.au	USA.Canada@sdi.com.au	Brasil@sdi.com.au
Registered company name	Registered company name SDI Germany GmbH		
Address	Hansestrasse 85 Cologne D-51149 Germany		
Telephone	+49 0 2203 9255 0		
Fax	+49 0 2203 9255 200		
Website	www.sdi.com.au		
Email	germany@sdi.com.au		

Emergency telephone number

Association / Organisation	SDI Limited	CHEMWATCH EMERGENCY RESPONSE (24/7)
Emergency telephone numbers	131126 Poisons Information Centre	+64 800 700 112
Other emergency telephone numbers	+61 3 8727 7111	+61 3 9573 3188

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

SECTION 2 Hazards identification

Classification of the substance or mixture

Classification ^[1]	Acute Toxicity (Oral) Category 4, Skin Corrosion/Irritation Category 2, Serious Eye Damage/Eye Irritation Category 1, Specific Target Organ Toxicity - Single Exposure (Respiratory Tract Irritation) Category 3
Legend:	1. Classification by vendor; 2. Classification drawn from CCID EPA NZ; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI
Determined by using GHS/HSNO criteria	6.1D (oral), 6.3A, 8.3A, 6.1E (respiratory tract irritant)

Label elements

Hazard pictogram(s)



Signal word Danger

Hazard statement(s)	
H302	Harmful if swallowed.
H315	Causes skin irritation.
H318	Causes serious eye damage.
H335	May cause respiratory irritation.

Precautionary statement(s) Prevention

P271	Use only outdoors or in a well-ventilated area.
P280	Wear protective gloves, protective clothing, eye protection and face protection.
P261	Avoid breathing mist/vapours/spray.
P264	Wash all exposed external body areas thoroughly after handling.
P270	Do not eat, drink or smoke when using this product.

Precautionary statement(s) Response

IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.
Immediately call a POISON CENTER/doctor/physician/first aider.
IF SWALLOWED: Call a POISON CENTER/doctor/physician/first aider if you feel unwell.
IF ON SKIN: Wash with plenty of water.
IF INHALED: Remove person to fresh air and keep comfortable for breathing.
Rinse mouth.
If skin irritation occurs: Get medical advice/attention.
Take off contaminated clothing and wash it before reuse.

Precautionary statement(s) Storage

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

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
7722-84-1	30-37.5	hydrogen peroxide
1310-73-2	<0.5	sodium hydroxide
Legend:	1. Classification by vendor; 2. Classification drawn from CCID EPA NZ; 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. Seek medical attention. 	
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

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

Special hazards arising from the substrate or mixture

Fire Incompatibility	Avoid any contamination of this material as it is very reactive and any contamination is potentially hazardous	
dvice for firefighters		
Fire Fighting	 Alert Fire Brigade and tell them location and nature of hazard. May be violently or explosively reactive. Wear full body protective clothing with breathing apparatus. Prevent, by any means available, spillage from entering drains or water courses. Fight fire from a safe distance, with adequate cover. Extinguishers should be used only by trained personnel. Use water delivered as a fine spray to control fire and cool adjacent area. DO NOT approach containers suspected to be hot. Cool fire exposed containers with water spray from a protected location. If safe to do so, remove containers from path of fire. If fire gets out of control withdraw personnel and warn against entry. Equipment should be thoroughly decontaminated after use. 	
Fire/Explosion Hazard	 Will not burn but increases intensity of fire. Heating may cause expansion or decomposition leading to violent rupture of containers. Heat affected containers remain hazardous. Contact with combustibles such as wood, paper, oil or finely divided metal may produce spontaneous combustion or violent decomposition May emit irritating, poisonous or corrosive fumes. 	

SECTION 6 Accidental release measures

Personal precautions, protective equipment and emergency procedures

See section 8

Environmental precautions

See section 12

Methods and material for containment and cleaning up

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

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

SECTION 7 Handling and storage

Precautions for safe handling	
Safe handling	For oxidisers, including peroxides.

	Avoid personal contact and inhalation of dust, mist or vapours.
	Provide adequate ventilation.
	Always wear protective equipment and wash off any spillage from clothing.
	· Keep material away from light, heat, flammables or combustibles.
	Keep cool, dry and away from incompatible materials.
	Avoid physical damage to containers.
	DO NOT repack or return unused portions to original containers. Withdraw only sufficient amounts for immediate use.
	Use only minimum quantity required.
	· Avoid using solutions of peroxides in volatile solvents. Solvent evaporation should be controlled to avoid dangerous concentration of the
	peroxide.
	Do NOT allow oxidisers to contact iron or compounds of iron, cobalt, or copper, metal oxide salts, acids or bases.
	Do NOT use metal spatulas to handle oxidisers
	Do NOT use glass containers with screw cap lids or glass stoppers.
	Store peroxides at the lowest possible temperature, consistent with their solubility and freezing point.
	CAUTION: Do NOT store liquids or solutions of peroxides at a temperature below that at which the oxidiser freezes or precipitates. Peroxides
	in particular, in this form are extremely shock and heat-sensitive. Refrigerated storage of peroxides must ONLY be in explosion-proof units.
	. The hazards and consequences of fires and explosions during synthesis and use of oxidisers is widely recognised; spontaneous or induced
	decomposition may culminate in a variety of ways, ranging from moderate gassing to spontaneous ignition or explosion. The heat released fro
	spontaneous decomposition of an energy-rich compound causes a rise in the surrounding temperature; the temperature will rise until thermal
	balance is established or until the material heats to decomposition,
	The most effective means for minimising the consequences of an accident is to limit quantities to a practical minimum. Even gram-scale
	explosions can be serious. Once ignited the burning of peroxides cannot be controlled and the area should be evacuated.
	Unless there is compelling reason to do otherwise, peroxide concentration should be limited to 10% (or less with vigorous reactants). Peroxid
	concentration is rarely as high as 1% in the reaction mixture of polymerisation or other free-radical reactions,
	Oxidisers should be added slowly and cautiously to the reaction medium. This should be completed prior to heating and with good agitation.
	Addition oxidisers to the hot monomer is extremely dangerous. A violent reaction (e.g., fire or explosion) can result from inadvertent mixing of
	promoters (frequently used with peroxides in polymerisation systems) with full-strength oxidisers
	Organic peroxides are very sensitive to contamination (especially heavy-metal compounds, metal oxide salts, alkaline materials including
	amines, strong acids, and many varieties of dust and dirt). This can initiate rapid, uncontrolled decomposition of peroxides and possible
	generation of intense heat, fire or explosion The consequences of accidental contamination from returning withdrawn material to the storage
	generation of mense react, me of explosion the consequences of accidental containing with a with material to the storage container can be disastrous.
	· When handling NEVER smoke, eat or drink.
	Always wash hands with soap and water after handling.
	Vise only good occupational work practice.
	Observe manufacturer's storage and handling recommendations contained within this MSDS.
	Store in a dry and well ventilated-area, away from heat and sunlight.
Other information	Store between 2 and 8 deg C.
	Do not store in direct sunlight.

Conditions for safe storage, including any incompatibilities

Suitable container	DO NOT repack. Use containers supplied by manufacturer only.
Storage incompatibility	Avoid strong acids, bases.

SECTION 8 Exposure controls / personal protection

Control parameters

Occupational Exposure Limits (OEL)

INGREDIENT DATA

Source	Ingredient	Material name	TWA	STEL	Peak	Notes
New Zealand Workplace Exposure Standards (WES)	hydrogen peroxide	Hydrogen peroxide	1 ppm / 1.4 mg/m3	Not Available	Not Available	Not Available
New Zealand Workplace Exposure Standards (WES)	sodium hydroxide	Sodium hydroxide	Not Available	Not Available	2 mg/m3	Not Available

Emergency Limits

Ingredient	TEEL-1	TEEL-2		TEEL-3
hydrogen peroxide	Not Available	Not Available		Not Available
sodium hydroxide	Not Available	Not Available		Not Available
Ingredient	Original IDLH		Revised IDLH	
hydrogen peroxide	75 ppm		Not Available	
sodium hydroxide	10 mg/m3		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 controls can be highly effective in protecting workers and will typically be independent of worker interactions to provide this high level of protection. The basic types of engineering controls are: Process controls which involve changing the way a job activity or process is done to reduce the risk. Enclosure and/or isolation of emission source which keeps a selected hazard "physically" away from the worker and ventilation that strategically "adds" and "removes" air in the work environment. Ventilation can remove or dilute an air contaminant if designed properly. The design of a ventilation system must match the particular process and chemical or contaminant in use. Employers may need to use multiple types of controls to prevent employee overexposure. Local exhaust ventilation usually required. If risk of overexposure exists, wear approved respirator. Correct fit is essential to obtain adequate protection. Supplied-air type respirator may be required in special circumstances. Correct fit is essential to ensure adequate protection. An approved self contained breathing apparatus (SCBA) may be required in some situations. Provide adequate ventilation in warehouse or closed storage area. Air contaminants generated in the workplace possess varying "escape"
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	velocities which, in turn, determine the "capture velocities" of	fresh circulating air required to effectively remove the conta	minant.
	Type of Contaminant:		Air Speed:
	solvent, vapours, degreasing etc., evaporating from tank (ir	n still air).	0.25-0.5 m/s (50-100 f/min.)
	aerosols, fumes from pouring operations, intermittent conta drift, plating acid fumes, pickling (released at low velocity ir		0.5-1 m/s (100-200 f/min.)
	direct spray, spray painting in shallow booths, drum filling, or 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 ger very high rapid air motion).	nerated dusts (released at high initial velocity into zone of	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	1: Disturbing room air currents	
	2: Contaminants of low toxicity or of nuisance value only.	2: Contaminants of high toxicity	
	3: Intermittent, low production.	3: High production, heavy use	
	4: Large hood or large air mass in motion	4: Small hood-local control only	
	accordingly, after reference to distance from the contaminatin 1-2 m/s (200-400 f/min) for extraction of solvents generated in producing performance deficits within the extraction apparatu more when extraction systems are installed or used.	n a tank 2 meters distant from the extraction point. Other me	echanical considerations
Individual protection measures, such as personal protective equipment			
Eye and face protection	the wearing of lenses or restrictions on use, should be cr and adsorption for the class of chemicals in use and an a their removal and suitable equipment should be readily a remove contact lens as soon as practicable. Lens should		ew of lens absorption should be trained in tion immediately and ens should be removed
Skin protection	See Hand protection below		
Hands/feet protection	 Wear chemical protective gloves, e.g. PVC. Wear safety footwear or safety gumboots, e.g. Rubber 		
Body protection	See Other protection below		
Other protection	 Overalls. PVC Apron. PVC protective suit may be required if exposure severe. Evewash unit. 		

Respiratory protection

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

Where the concentration of gas/particulates in the breathing zone, approaches or exceeds the "Exposure Standard" (or ES), respiratory protection is required. Degree of protection varies with both face-piece and Class of filter; the nature of protection varies with Type of filter.

Required Minimum Protection Factor	Half-Face Respirator	Full-Face Respirator	Powered Air Respirator
up to 10 x ES	B-AUS	-	B-PAPR-AUS / Class 1
up to 50 x ES	-	B-AUS / Class 1	-
up to 100 x ES	-	B-2	B-PAPR-2 ^

^ - Full-face

A(All classes) = Organic vapours, B AUS or B1 = Acid gasses, B2 = Acid gas or hydrogen cyanide(HCN), B3 = Acid gas or hydrogen cyanide(HCN), E = Sulfur dioxide(SO2), G = Agricultural chemicals, K = Ammonia(NH3), Hg = Mercury, NO = Oxides of nitrogen, MB = Methyl bromide, AX = Low boiling point organic compounds(below 65 degC)

SECTION 9 Physical and chemical properties

Information on basic physical and chemical properties

Appearance	Clear, blue gel, mixes with water.		
Physical state	Gel	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 Available
pH (as supplied)	6.5-8	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 Available	Taste	Not Available
Evaporation rate	Not Available	Explosive properties	Not Available
Flammability	Not Available	Oxidising properties	Not Available
Upper Explosive Limit (%)	Not Available	Surface Tension (dyn/cm or mN/m)	Not Available
Lower Explosive Limit (%)	Not Available	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

SECTION 10 Stability and reactivity

Reactivity	See section 7
Chemical stability	 Unstable in the presence of incompatible materials. Product is considered stable under normal handling conditions. Prolonged exposure to heat. Hazardous polymerisation will not occur.
Possibility of hazardous reactions	See section 7
Conditions to avoid	See section 7
Incompatible materials	See section 7
Hazardous decomposition products	See section 5

SECTION 11 Toxicological information

Information on toxicological effects

Inhaled	individuals, following inhalation. In contrast to most organs irritant and then repairing the damage. The repair process may however, produce further lung damage resulting in th	e material produces irritation of the respiratory system, in a substantial number of s, the lung is able to respond to a chemical insult by first removing or neutralising the s, which initially evolved to protect mammalian lungs from foreign matter and antigens, ne impairment of gas exchange, the primary function of the lungs. Respiratory tract ng the recruitment and activation of many cell types, mainly derived from the vascular
Ingestion	Accidental ingestion of the material may be harmful; anim produce serious damage to the health of the individual.	al experiments indicate that ingestion of less than 150 gram may be fatal or may
Skin Contact	following direct contact, and/or produces significant inflam inflammation being present twenty-four hours or more after repeated exposure; this may result in a form of contact de and swelling (oedema) which may progress to blistering (' may be intercellular oedema of the spongy layer of the sk The material may accentuate any pre-existing dermatitis of Skin contact will result in rapid drying, bleaching, leading of Open cuts, abraded or irritated skin should not be expose	condition to chemical burns on prolonged contact d to this material vrasions, puncture wounds or lesions, may produce systemic injury with harmful effects
Eye	When applied to the eye(s) of animals, the material produ	ces severe ocular lesions which are present twenty-four hours or more after instillation
_)··		
Chronic		isease of the airways involving difficult breathing and related systemic problems. supational exposure may produce cumulative health effects involving organs or
Chronic	Limited evidence suggests that repeated or long-term occ	isease of the airways involving difficult breathing and related systemic problems.
	Limited evidence suggests that repeated or long-term occ biochemical systems.	isease of the airways involving difficult breathing and related systemic problems. supational exposure may produce cumulative health effects involving organs or
Chronic	Limited evidence suggests that repeated or long-term occ biochemical systems.	isease of the airways involving difficult breathing and related systemic problems. supational exposure may produce cumulative health effects involving organs or IRRITATION
Chronic Pola Professional 37.5%	Limited evidence suggests that repeated or long-term occ biochemical systems. TOXICITY Not Available	isease of the airways involving difficult breathing and related systemic problems. upational exposure may produce cumulative health effects involving organs or IRRITATION Not Available
Chronic	Limited evidence suggests that repeated or long-term occ biochemical systems. TOXICITY Not Available TOXICITY	isease of the airways involving difficult breathing and related systemic problems. supational exposure may produce cumulative health effects involving organs or IRRITATION Not Available IRRITATION
Chronic Pola Professional 37.5%	Limited evidence suggests that repeated or long-term occ biochemical systems. TOXICITY Not Available TOXICITY Dermal (rabbit) LD50: >2000 mg/kg ^[1]	isease of the airways involving difficult breathing and related systemic problems. upational exposure may produce cumulative health effects involving organs or IRRITATION Not Available IRRITATION
Chronic Pola Professional 37.5%	Limited evidence suggests that repeated or long-term occ biochemical systems. TOXICITY Not Available TOXICITY Dermal (rabbit) LD50: >2000 mg/kg ^[1] Inhalation(Mouse) LC50; 2800 mg/L4h ^[2]	isease of the airways involving difficult breathing and related systemic problems. upational exposure may produce cumulative health effects involving organs or IRRITATION Not Available IRRITATION
Chronic Pola Professional 37.5%	Limited evidence suggests that repeated or long-term occ biochemical systems. TOXICITY Not Available TOXICITY Dermal (rabbit) LD50: >2000 mg/kg ^[1] Inhalation(Mouse) LC50; 2800 mg/L4h ^[2] Oral (Rat) LD50: >225 mg/kg ^[2]	isease of the airways involving difficult breathing and related systemic problems. upational exposure may produce cumulative health effects involving organs or IRRITATION Not Available IRRITATION Not Available
Chronic Pola Professional 37.5% hydrogen peroxide	Limited evidence suggests that repeated or long-term occ biochemical systems. TOXICITY Not Available TOXICITY Dermal (rabbit) LD50: >2000 mg/kg ^[1] Inhalation(Mouse) LC50; 2800 mg/L4h ^[2] Oral (Rat) LD50: >225 mg/kg ^[2] TOXICITY	isease of the airways involving difficult breathing and related systemic problems. upational exposure may produce cumulative health effects involving organs or IRRITATION Not Available IRRITATION Not Available IRRITATION IRRITATION IRRITATION IRRITATION
Chronic Pola Professional 37.5%	Limited evidence suggests that repeated or long-term occ biochemical systems. TOXICITY Not Available TOXICITY Dermal (rabbit) LD50: >2000 mg/kg ^[1] Inhalation(Mouse) LC50; 2800 mg/L4h ^[2] Oral (Rat) LD50: >225 mg/kg ^[2] TOXICITY Dermal (rabbit) LD50: >25 mg/kg ^[2]	isease of the airways involving difficult breathing and related systemic problems. upational exposure may produce cumulative health effects involving organs or IRRITATION Not Available IRRITATION Not Available IRRITATION Eye (rabbit): 0.05 mg/24h SEVERE

		Skin (rabbit): 50	0 mg/24h SEVERE		
		Skin: adverse ef	fect observed (corrosive) ^[1]		
Legend:	Legend: 1. Value obtained from Europe ECHA Registered Substances - Acute toxicity 2. Value obtained from manufacturer's SDS. Unles specified data extracted from RTECS - Register of Toxic Effect of chemical Substances				
HYDROGEN PEROXIDE	No significant acute toxicological data identified in literature search. For hydrogen peroxide: Hazard increases with peroxide concentration, high concentrations contain an additive stabiliser. Pharmacokinetics Hydrogen peroxide is a normal product of metabolism. It is readily decomposed by catalase in normal cells. In experimental animals exposed hydrogen peroxide is a normal product of metabolism. It is readily decomposed by catalase in normal cells. In experimental animals exposed hydrogen peroxide has been detected in breath. • Absorption: Hydrogen peroxide is decomposed in the bowel before absorption. When applied to tissue, solutions of hydrogen peroxide in the bowel perior by the period of the second of the bowel before absorption. When applied to tissue, solutions of hydrogen peroxide in two-electron transfer reaction, often catalysed by flavoproteins, or by an initial one-electron step to O2 followed by dismutation to hydrogen peroxide. • Hydrogen peroxide has been detected in serum and in intact liver, based on the results of toxicity studies, the lungs, intestine, thymus, lin and kidney may be distribution sites. In rabbits and cast that died after intravenous administration of hydrogen peroxide has been detected in a real tubular epithelial tissue was observed following oral administration of hydrogen peroxide to mice. • Metabolism Glutathione peroxides, responsible for decomposing hydrogen peroxide, is present in normal human tissues (IARC 1985). When hydrogen peroxide has been detected in human breath at levels ranging from 1.0+/-5 g/L to 0.34+/-0.17 g/L. Carcinogenicity Gastric and duodenal lesions including adenomas, carcinomas, and adenocarcinomas have been observed in mice treated orally with hydrogen peroxide induced DNA damage, sister chromatid exchanges and chromosomal aberrations in marmalian cells <i>in vitro</i> . Hydrogen peroxide induced DNA damage, sister chromatid exchanges and chromosomal aberrations in marmalian cells <i>in vitro</i> . Hydrogen peroxide induced DNA damage, sister chromatid				
	The material may produce severe irritation to the eye causing pronounced inflammation. Repeated or prolonged exposure to irritants may produce conjunctivitis. The material may produce severe skin irritation after prolonged or repeated exposure, and may produce a contact dermatitis (nonallergic). This form of dermatitis is often characterised by skin redness (erythema) thickening of the epidermis. Histologically there may be intercellular oedema of the spongy layer (spongiosis) and intracellular oedema of the epidermis. Prolonged contact is unlikely, given the severity of response, but repeated exposures may produce severe ulceration.				
HYDROGEN PEROXIDE & SODIUM HYDROXIDE	Asthma-like symptoms may continue for months or even years after exposure to the material ends. This may be due to a non-allergic condition known as reactive airways dysfunction syndrome (RADS) which can occur after exposure to high levels of highly irritating compound. Main criteria for diagnosing RADS include the absence of previous airways disease in a non-atopic individual, with sudden onset of persistent asthma-like symptoms within minutes to hours of a documented exposure to the irritant. Other criteria for diagnosis of RADS include a reversible airflow pattern on lung function tests, moderate to severe bronchial hyperreactivity on methacholine challenge testing, and the lack of minimal lymphocytic inflammation, without eosinophilia. RADS (or asthma) following an irritating inhalation is an infrequent disorder with rates related to the concentration of and duration of exposure to the irritating substance. On the other hand, industrial bronchitis is a disorder that occurs as a result of exposure due to high concentrations of irritating substance (often particles) and is completely reversible after exposure ceases. The disorder is characterized by difficulty breathing, cough and mucus production.				
Acute Toxicity	~	Carcinogenicity	×		
Skin Irritation/Corrosion	 ✓ 	Reproductivity	×		
Serious Eye Damage/Irritation	✓	STOT - Single Exposure	×		
Respiratory or Skin sensitisation	×	STOT - Repeated Exposure	×		

SECTION 12 Ecological information

 Toxicity
 Endpoint
 Test Duration (hr)
 Species
 Value
 Source

	Not Available	Not Available		Not Available		Not Available	Not Available
	Endpoint	Test Duration (hr)		Species		Value	Source
	EC50	72h		Algae or other aquatic plants		0.69mg/l	4
hydrogen peroxide	EC50	48h		Crustacea		2mg/l	2
	EC50	96h		Algae or other aquatic plants		2.27mg/l	4
	NOEC(ECx)	72h		Algae or other aquatic plants		0.1mg/l	1
	LC50	96h		Fish		16.4mg/l	2
	Endpoint	Test Duration (hr)	s	pecies	Valu	e	Source
	EC50	48h	С	rustacea	34.5	9-47.13mg/l	4
sodium hydroxide	LC50	96h	Fish		144-	267mg/l	4
	EC50(ECx)	48h	С	rustacea	34.5	9-47.13mg/l	4
Legend:	Ecotox databas	1. IUCLID Toxicity Data 2. Europe E e - Aquatic Toxicity Data 5. ECETO on Data 8. Vendor Data	•	•			

DO NOT discharge into sewer or waterways.

Persistence and degradability

Ingredient	Persistence: Water/Soil	Persistence: Air
hydrogen peroxide	LOW	LOW
sodium hydroxide	LOW	LOW

Bioaccumulative potential

Ingredient	Bioaccumulation
hydrogen peroxide	LOW (LogKOW = -1.571)
sodium hydroxide	LOW (LogKOW = -3.8796)

Mobility in soil

Ingredient	Mobility
hydrogen peroxide	LOW (KOC = 14.3)
sodium hydroxide	LOW (KOC = 14.3)

SECTION 13 Disposal considerations

Waste treatment methods	
Product / Packaging disposal	 DO NOT allow wash water from cleaning or process equipment to enter drains. It may be necessary to collect all wash water for treatment before disposal. In all cases disposal to sewer may be subject to local laws and regulations and these should be considered first. Where in doubt contact the responsible authority. Recycle wherever possible. Special hazard may exist - specialist advice may be required. Consult manufacturer for recycling options. Consult State Land Waste Management Authority for disposal. Bury or incinerate residue at an approved site. Decontaminate empty containers. Observe all label safeguards until containers are cleaned and destroyed. Puncture containers to prevent re-use and bury at an authorised landfill.

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

Land transport (UN)		
14.1. UN number or ID number	2014	
14.2. UN proper shipping name	HYDROGEN PEROXI	DE, AQUEOUS SOLUTION with not less than 20% but not more than 60% hydrogen peroxide (stabilized as necessary)
14.3. Transport hazard class(es)	Class Subsidiary Hazard	5.1 8
14.4. Packing group	Ш	
14.5. Environmental hazard	Not Applicable	
14.6. Special precautions for user	Special provisions Limited quantity	Not Applicable

Air transport (ICAO-IATA / DGR)

14.1. UN number	2014			
14.2. UN proper shipping name	Hydrogen peroxide, aqueous solution with more than 40% but 60% or less hydrogen peroxide (stabilized as necessary); Hydrogen peroxide, aqueous solution with 20% or more but 40% or less hydrogen peroxide (stabilized as necessary)			
	ICAO/IATA Class	5.1		
14.3. Transport hazard class(es)	ICAO / IATA Subsidiary Hazard	8		
01255(85)	ERG Code	5C		
14.4. Packing group	II			
14.5. Environmental hazard	Not Applicable			
14.6. Special precautions for user	Special provisions		A2 A75	
	Cargo Only Packing Instructions		554; Forbidden	
	Cargo Only Maximum Qty / Pack		5 L; Forbidden	
	Passenger and Cargo Packing Instructions		550; Forbidden	
	Passenger and Cargo Maximum Qty / Pack		1 L; Forbidden	
	Passenger and Cargo Limited Qu	antity Packing Instructions	Y540; Forbidden	
	Passenger and Cargo Limited Ma	ximum Qtv / Pack	0.5 L; Forbidden	

Sea transport (IMDG-Code / GGVSee)

14.1. UN number	2014			
14.2. UN proper shipping name	HYDROGEN PEROXIDE	HYDROGEN PEROXIDE, AQUEOUS SOLUTION with not less than 20% but not more than 60% hydrogen peroxide (stabilized as necessary)		
14.3. Transport hazard class(es)	IMDG Class IMDG Subsidiary Haza	5.1 ard 8		
14.4. Packing group	II			
14.5 Environmental hazard	Not Applicable			
14.6. Special precautions for user	Special provisions	F-H, S-Q Not Applicable		

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
hydrogen peroxide	Not Available
sodium hydroxide	Not Available

14.7.3. Transport in bulk in accordance with the IGC Code

Product name	Ship Type
hydrogen peroxide	Not Available
sodium hydroxide	Not Available

If packed as Chemical kits the following classification may be considered if all ICAO/IATA transport requirements are met: Chemical Kit UN3316 - Class 9, SP A44 & A163.

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			
HSR002521	Animal Nutritional and Animal Care Products Group Standard 2020			
HSR002530	Cleaning Products Subsidiary Hazard Group Standard 2020			
HSR002535	Gases under Pressure Mixtures Subsidiary Hazard Group Standard 2020			
HSR002503	Additives Process Chemicals and Raw Materials Subsidiary Hazard Group Standard 2020			
HSR002606	Lubricants Lubricant Additives Coolants and Anti freeze Agents Subsidiary Hazard Group Standard 2020			
HSR002612	Metal Industry Products Subsidiary Hazard Group Standard 2020			
HSR002624	N.O.S. Subsidiary Hazard Group Standard 2020			
HSR002638	Photographic Chemicals Subsidiary Hazard Group Standard 2020			
HSR002644	Polymers Subsidiary Hazard Group Standard 2020			
HSR002647	Reagent Kits Group Standard 2020			
HSR002648	Refining Catalysts Group Standard 2020			
HSR002653	Solvents Subsidiary Hazard Group Standard 2020			
HSR002670	Surface Coatings and Colourants Subsidiary Hazard Group Standard 2020			
HSR002684	Water Treatment Chemicals Subsidiary Hazard Group Standard 2020			
HSR100425	Pharmaceutical Active Ingredients Group Standard 2020			
HSR002600	Leather and Textile Products Subsidiary Hazard Group Standard 2020			
HSR002544	Construction Products Subsidiary Hazard Group Standard 2020			
HSR002549	Corrosion Inhibitors Subsidiary Hazard Group Standard 2020			
HSR002552	Cosmetic Products Group Standard 2020			
HSR002558	Dental Products Subsidiary Hazard Group Standard 2020			
HSR002565	Embalming Products Subsidiary Hazard Group Standard 2020			
HSR002571	Fertilisers Subsidiary Hazard Group Standard 2020			
HSR002573	Fire Fighting Chemicals Group Standard 2021			
HSR002578	Food Additives and Fragrance Materials Subsidiary Hazard Group Standard 2020			
HSR002585	Fuel Additives Subsidiary Hazard Group Standard 2020			
HSR002596	Laboratory Chemicals and Reagent Kits Group Standard 2020			
HSR100580	Tattoo and Permanent Makeup Substances Group Standard 2020			
HSR100757	Veterinary Medicines Limited Pack Size Finished Dose Group Standard 2020			
HSR100758	Veterinary Medicines Non dispersive Closed System Application Group Standard 2020			
HSR100759	Veterinary Medicines Non dispersive Open System Application Group Standard 2020			

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

hydrogen peroxide is found on the following regulatory lists

International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Monographs - Not Classified as Carcinogenic

- New Zealand Approved Hazardous Substances with controls
- New Zealand Hazardous Substances and New Organisms (HSNO) Act Classification of Chemicals
- New Zealand Hazardous Substances and New Organisms (HSNO) Act Classification of Chemicals Classification Data
- New Zealand Inventory of Chemicals (NZIoC)
- New Zealand Workplace Exposure Standards (WES)

sodium hydroxide 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)

New Zealand Workplace Exposure Standards (WES)

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

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Pola Professional 37.5%

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
Not Applicable	Not Applicable	Not Applicable	Not Applicable	Not Applicable

Tracking Requirements

Not Applicable

National Inventory Status

National Inventory	Status			
Australia - AIIC / Australia Non-Industrial Use	Yes			
Canada - DSL	Yes			
Canada - NDSL	No (hydrogen peroxide; sodium hydroxide)			
China - IECSC	Yes			
Europe - EINEC / ELINCS / NLP	Yes			
Japan - ENCS	Yes			
Korea - KECI	Yes			
New Zealand - NZIoC	Yes			
Philippines - PICCS	Yes			
USA - TSCA	Yes			
Taiwan - TCSI	Yes			
Mexico - INSQ	Yes			
Vietnam - NCI	Yes			
Russia - FBEPH	Yes			
Legend:	Yes = All CAS declared ingredients are on the inventory No = One or more of the CAS listed ingredients are not on the inventory. These ingredients may be exempt or will require registration.			

SECTION 16 Other information

Revision Date	23/12/2022
Initial Date	10/11/2015

SDS Version Summary

Version	Date of Update	Sections Updated
6.1	01/11/2019	One-off system update. NOTE: This may or may not change the GHS classification
7.1	23/12/2022	Classification review due to GHS Revision change.

Other information

Classification of the preparation and its individual components has drawn on official and authoritative sources as well as independent review by SDI Limited 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
- ٠ STEL: Short Term Exposure Limit
- ► TEEL: Temporary Emergency Exposure Limit.
- ۲ IDLH: Immediately Dangerous to Life or Health Concentrations
- ٠ ES: Exposure Standard
- OSF: Odour Safety Factor
- NOAEL: No Observed Adverse Effect Level ٠
- LOAEL: Lowest Observed Adverse Effect Level ۲ TLV: Threshold Limit Value
- ٠ LOD: Limit Of Detection
- OTV: Odour Threshold Value
- ٠ BCF: BioConcentration Factors
- ٠
- BEI: Biological Exposure Index DNEL: Derived No-Effect Level ۲
- ۲ PNEC: Predicted no-effect concentration
- AIIC: Australian Inventory of Industrial Chemicals
- DSL: Domestic Substances List

- NDSL: Non-Domestic Substances List
- IECSC: Inventory of Existing Chemical Substance in China
- ▶ EINECS: European INventory of Existing Commercial chemical Substances ELINCS: European List of Notified Chemical Substances
- NLP: No-Longer Polymers
 ENCS: Existing and New Chemical Substances Inventory
- KECI: Korea Existing Chemicals Inventory
- NZIoC: New Zealand Inventory of Chemicals
 PICCS: Philippine Inventory of Chemicals and Chemical Substances
- TSCA: Toxic Substances Control Act
- TCSI: Taiwan Chemical Substance Inventory
- INSQ: Inventario Nacional de Sustancias Químicas
- NCI: National Chemical Inventory
- + FBEPH: Russian Register of Potentially Hazardous Chemical and Biological Substances

The information contained in the Safety Data Sheet is based on data considered to be accurate, however, no warranty is expressed or implied regarding the accuracy of the data or the results to be obtained from the use thereof.

Other information:

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