

# Section 1 – Identification of Chemical Product and Company

01468     Gorilla Power Fixer Automotive Part B     10 gm     Grey       01460     Gerilla Power Fixer Automotive Part B     20 gm     Grey	Code	Description	Size	Colour
01460 Covilla Dover Eiver Automotiva Dart P. 20 am Crav	01468	Gorilla Power Fixer Automotive Part B	10 gm	Grey
Grina Power Fixer Automotive Part B So gin Grey	01469	Gorilla Power Fixer Automotive Part B	30 gm	Grey

Recommended use:	Filler			
Supplier contact details:	Soudal Ltd	Freephone: 0800 70 10 80		
	14 Avalon Drive	Phone: (07) 847 5540		
	Nawton	Fax: (07) 847 0324		
	Hamilton 3200	Email: sales@soudal.co.nz		
	New Zealand	Website: <u>www.soudal.co.nz</u>		
POISON CENTRE NUMBER: 0800 764 766 (24 hours)				

# Section 2 – Hazard Identification

#### **Statement of Hazardous Nature**

This product is classified as:

**HAZARDOUS SUBSTANCE** according to the criteria of HSNO.

**REGULATED** under NZS5433:2007 Transport of Dangerous Goods on Land

# Hazardous Substances and New Organisms (HSNO) classification:

Classification		Hazard	statements
Organic Peroxide Type E 5.2E		H242	Heating may cause a fire
Acute Oral Toxicity Category 4 6.1D		H302	Harmful if swallowed
Acute Inhalation Toxicity Category 4	6.1D	H332	Harmful if inhaled
Skin Effects Category 2	6.3A	H315	Causes skin irritation
Eye Effects Category 2	6.4A	H319	Causes serious eye irritation
Skin Sensitisation Category 1 6.5B		H317	May cause an allergic skin reaction
Chronic Aquatic Hazard Category 1	9.1A	H410	Very toxic to aquatic life with long lasting effects
Vertebrate hazard Category 3 9.3C		H433	harmful to terrestrial vertebrates

**HSNO Signal Word:** 



#### **Precautionary Statements:**

Read label before use.

Keep out of reach of children.

Ensure all safety directions are read and understood before use Keep away from heat, sparks, open flames and hot surfaces No smoking Keep only in original packaging Keep Cool Ground/ bond container and receiving equipment Use only outdoors or in a well-ventilated place Avoid breathing fumes/ mists/ vapours/ sprays



Wear protective gloves/ protective clothing/ eye protection/ face Contaminated work clothing should not be allowed out of the workplace

Do not eat, drink or smoke while handling this product

In case of fire Use Carbon Dioxide, Dry powder to extinguish

#### Section 3 - Composition/Information on Ingredients

Avoid release to the environment

Store in a well-ventilated place. Store at temperatures not exceeding 30°C Store separately Protect from sunlight

Ingredient	CAS No.	Individual HSNO classification	Concentration (% by Wt.)
Dibenzoyl peroxide	94-36-0	Organic Peroxide Type B; Eye Effects Category 2; Skin Sensitsation Category 1; Chronic Aquatic hazard Category 1	40 – 60
Dimethyl Phthalate	131-11-3	Acute Oral Toxicity Category 4; Acute Inhalation Toxicity Category 3; Eye Effects Category 2; Chronic Aquatic hazard Category 4; Vertebrate Hazard Category 3	20 – 40
Ethylene glycol	107-21-1	Acute Oral Toxicity Category 4; Eye Effects Category 2; STOT – SE Category 1; STOT – RE Category 1; Vertebrate Hazard Category 3	1 - 10

This is a commercial product whose exact ratio of components may vary slightly. Minor quantities of other non hazardous ingredients are also possible.

# Section 4 – First Aid Measures

# NZ Poisons Centre 0800 POISON (0800 764 766) | NZ Emergency Services: 111

#### Eye contact:

Immediately hold the eyelids apart and flush the eye with 2% sodium carbonate solution or 5% sodium ascorbate solution then wash continuously for at

least 15 minutes with fresh running water. Ensure complete irrigation of the eye by keeping eyelids apart and away from eye and moving the eyelids by occasionally lifting the upper and lower lids. Transport to hospital (or doctor) without further delay. Removal of contact lenses should only be undertaken by trained personnel.

# Skin or hair contact:

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, without delay.

#### 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.

## General advice and advice for physicians:

Treat symptomatically.

You should call The Poisons Information Centre if you feel that you may have been poisoned, burned or irritated by this product. The number is 0800 764766 from anywhere in New Zealand (13 1126 in Australia) and is available at all times. Have this SDS or product label with you when you call.

Section 5 - Fire-Fighting Measures





# Extinguishing media:

Foam, Carbon Dioxide, Dry Powder, water fog

## Fire/ Explosion Hazard

Liquid and vapour are flammable. Moderate fire hazard when exposed to heat or flame. Vapour forms an explosive mixture with air. Moderate explosion hazard when exposed to heat or flame. Vapour may travel a considerable distance to source of ignition. Heating may cause expansion or decomposition leading to violent rupture of containers. On combustion, may emit toxic fumes of carbon monoxide (CO).

# Advice for fire-fighters:

Alert Fire & Emergency New Zealand and tell them location and nature of hazard. May be violently or explosively reactive. Wear breathing apparatus plus protective gloves in the event of a fire. Prevent, by any means available, spillage from entering drains or water course. Consider evacuation (or protect in place). Fight fire from a safe distance, with adequate cover. If safe, switch off electrical equipment until vapour fire hazard removed. Use water delivered as a fine spray to control the fire and cool adjacent area. Avoid spraying water onto liquid pools. Cool fire exposed containers with water spray from a protected location. If safe to do so, remove containers from path of fire.

# Section 6 - Accidental Release Measures

# **Minor Spills**

Environmental hazard - contain spillage. Remove all ignition sources. Clean up all spills immediately. Avoid breathing vapours and contact with skin and eyes. Control personal contact with the substance, by using protective equipment. Contain and absorb small quantities with vermiculite or other absorbent material. Wipe up. Collect residues in a flammable waste container. Slippery when spilt.

# **Major Spills:**

Environmental hazard – contain spillage. Slippery when spilt. Clear area of personnel. Alert Fire & Emergency New Zealand and tell them location and nature of hazard. Control personal contact with the substance, by using protective equipment as required. Prevent spillage from entering drains or water ways. Contain spill with sand, earth or vermiculite. Collect recoverable product into labelled containers for recycling. Absorb remaining product with sand, earth or vermiculite and place in appropriate containers for disposal. Wash area and prevent runoff into drains or waterways. If contamination of drains or waterways occurs, advise emergency services.

## Section 7 - Handling and Storage

#### Handling:

Mix only as much as is required. DO NOT return the mixed material to original containers. 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 peroxides to contact iron or compounds of iron, cobalt, or copper, metal oxide salts, acids or bases. Do NOT use metal spatulas to handle peroxides 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 peroxide freezes or precipitates. Peroxides in this form are extremely shock and heat-sensitive. Refrigerated storage of peroxides must ONLY be in explosionproof units. The hazards and consequences of fires and explosions during synthesis and use of peroxides is widely recognised; spontaneous or induced decomposition may culminate in a variety of ways, ranging from moderate gassing to spontaneous ignition or explosion. The heat released from spontaneous 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). Peroxide concentration is rarely as high as 1% in the reaction mixture of polymerisation or other free-radical reactions, Peroxides should be added slowly and cautiously to the reaction medium. This should be completed prior to heating and with good agitation. Addition of peroxide 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 peroxide 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 container can be disastrous. When handling NEVER smoke, eat or drink. Always wash hands with soap and water after handling. Use only good occupational work practice. Observe manufacturer's storage and handling recommendations contained within this SDS.

#### Storage:

Store in original containers in an isolated approved flammable materials storage area. Keep containers securely sealed as supplied. WARNING: Gradual decomposition during storage in sealed containers may lead to a large pressure build-up and subsequent explosion. No smoking, naked lights, heat or ignition sources. Store in a cool, dry, well ventilated area. Store under cover and away from sunlight. Store below safe storage (control) temperature. Always store below 30 °C. Store away from flammable or combustible materials, debris and waste. Contact may cause fire or violent reaction. Store away from incompatible materials. Store away from foodstuff containers DO NOT stack on wooden floors or wooden



pallets. Protect containers against physical damage. Check regularly for spills and leaks. Observe manufacturer's storage and handling recommendations contained within this SDS. Keep locked up.

Restrictions may apply on quantities and to other materials permitted in the same location.

#### FOR MINOR QUANTITIES:

Ensure that: packages are not opened in storage area, the goods are kept at least 3 metres from sources of heat as well as all other dangerous goods and all other materials which might react with this material might react to cause a fire, a chemical reaction or explosion, materials for absorbing and neutralising spills are kept near the storage; procedures are displayed at the storage describing actions to be taken in the event of a spill or fire. adequate numbers and types of portable fire extinguisher are provided in or near the storage area.

#### **Section 8 - Exposure Controls/Personal Protection**

#### **Exposure limits:**

CAS no.	Substance or ingredient	WES-TWA	WES-STEL
94-36-0	Dibenzoyl peroxide	5 mg/m <sup>3</sup>	

The TWA exposure value is the average airborne concentration of a particular substance when calculated over a normal 8 hour working day for a 5-day working week. The STEL (Short Term Exposure Limit) is an exposure value that may be equalled (but should not be exceeded) longer than 15 minutes and should not be repeated more than 4 times per day. There should be at least 60 minutes between successive exposures at the STEL. The term "peak "is used when the TWA limit, because of the rapid action of the substance, should never be exceeded, even briefly.

## **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. General exhaust is adequate under normal operating conditions. Local exhaust ventilation may be required in specific circumstances. If risk of overexposure exists, wear 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.

Control	Protective measure
Eye	Safety glasses with side shields. Chemical goggles. 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], [AS/NZS 1336 or national equivalent]
Respiratory	Type A organic vapour is reommended (AS/NZS 1716 & 1715, EN 143:2000 & 149:001, ANSI Z88 or national equivalent)
Skin	Wear chemical protective gloves, e.g. Natural+Neoprene; Natrural rubber; neoprene; Nitrile; PVC; Teflon Wear safety footwear or safety gumboots, e.g. Rubber NOTE: The material may produce skin sensitisation in predisposed individuals. Care must be taken, when removing gloves and other protective equipment, to avoid all possible skin contact. Contaminated leather items, such as shoes, belts and watch-bands should be removed and destroyed.

### Section 9 - Physical and Chemical Properties

#### General substance properties:



Property	Details
Appearance	Paste
Odour	Characteristic
рН	No data.
Vapour pressure	No data
Vapour Density	> 1 heavier than air
Viscosity	Viscous
Boiling Point	°C
Volatile materials	%
Water solubility	immiscible
Freezing/melting point	No data.
Solubility	Immiscible
Specific gravity/density	115 – 1.25 g/ml
Flash point	34 °C
Auto-ignition temperature	No Data
SADT	50 °C
Upper and lower flammability limits	Lower % Upper %
Corrosiveness	No data.

# Section 10 - Stability and Reactivity

# Stability:

Unstable in the presence of incompatible substances. Stable under normal conditions.

# Conditions to avoid:

Avoid heat, sparks, flames and any other sources of ignition.

# Incompatible materials to avoid:

Avoid oxidising agents (nitrates, oxidising acids, chlorine bleaches, pool chlorine etc) as ignition may result

# Hazardous decomposition products:

Combustion will result in the release of carbon monoxide; carbon dioxide and other toxic vapours

# Section 11 - Toxicological Information

# Acute toxicity:

Test	Data and symptoms of exposure
Inhaled	The material can cause respiratory irritation in some persons. The body's response to such irritation can cause further lung damage. The inhalation of organic peroxide dusts or vapours can produce throat and lung irritation and cause an asthma-like effect. Over-exposure can cause tears, salivation, lethargy, slow breathing,



	breathing difficulties, headache, weakness, tremor, stupor and swelling of the lung. Inhalation hazard is increased at higher temperatures.
Oral	Accidental ingestion of the material may be damaging to the health of the individual. Ingestion of organic peroxides may produce nausea, vomiting, abnormal pain, stupor, bluish discoloration of skin and mucous membranes. Inflammation of the heart muscle may also occur. Swallowing dimethyl phthalate may cause central nervous system depression. The toxicity of phthalates is not excessive due to slow oral absorption and metabolism. Absorption is affected by fat in the diet. Repeated doses can cause cumulative toxic effects, and symptoms include an enlarged liver which often reverses if exposure is maintained. Carbohydrate metabolism is disrupted, and cholesterol and triglyceride levels in the blood falls. In rats, there is also strong evidence of withering of the testicles. Some phthalates can increase the effects of antibiotics, thiamine (vitamin B1) and sulfonamides. For ethylene glycol: Symptoms following swallowing ethylene glycol include failure of breathing, central nervous system depression, cardiovascular collapse, lung swelling, acute kidney failure, and even brain damage. Swallowing 100 millilitres has caused death. There are three stages of ethylene glycol poisoning. The severity of each stage depends upon the amount of ethylene glycol swallowed. There is usually minimal damage to the liver. In the first 12 hours, central nervous system depression is necession is actidosis, coma, convulsions and seizures may also occur. Disorders in eye movements may occur, although otherwise eye examination usually remains normal. At 12-24 hours after swallowing, effects on the lung and heart appear. These are characterized by fast heart rate, fast breathing, and mildly high blood pressure. Congestive heart failure and circulatory collapse may occur. In severe poisonings. Effects on the kidney are seen 24-72 hours post-ingestion and are characterized by reduced urine output, flank pain, death of kidney are seen 24-72 hours post-ingestion and are characterized by reduced urine output, flank pain, death of kidney are see
Dermal	The material may accentuate any pre-existing dermatitis condition Skin contact with the material may damage the health of the individual; systemic effects may result following absorption. All organic peroxides are irritating to the skin and if allowed to remain on the skin, may produce inflammation; some are allergenic. Dimethyl phthalate, used as an insect repellent, has not been known to cause irritation or sensitization. Most liquid alcohols appear to act as primary skin irritants in humans. Significant percutaneous absorption occurs in rabbits but not apparently in man. Open cuts, abraded or irritated skin should not be exposed to this material Entry into the blood-stream, through for example, cuts, abrasions or lesions, may produce systemic injury with harmful effects. Examine the skin prior to the use of the material and ensure that any external damage is suitably protected. There is some evidence to suggest that the material may cause mild but significant inflammation of the skin either following direct contact or after a delay of some time. Repeated exposure can cause contact dermatitis which is characterised by redness, swelling and blistering.
Eye	Animal testing shows dimethyl phthalate only causes slight irriation when applied undiluted to the eye. Eye contact with organic peroxides can cause clouding, redness, swelling and burns of the eye on prolonged contact. Evidence exists, or practical experience predicts, that the material may cause eye irritation in a substantial number of individuals. Prolonged eye contact may cause inflammation characterised by a temporary redness of the conjunctiva (similar to windburn).
Chronic	Long-term exposure to respiratory irritants may result in airways disease, involving difficulty breathing and related whole-body problems. Strong evidence exists that this substance may cause irreversible mutations (though not lethal) even following a single exposure. Skin contact with the material is more likely to cause a sensitisation reaction in some persons compared to the general population. Laboratory (in vitro) and animal studies show, exposure to the material may result in a possible risk of irreversible effects, with the possibility of producing mutation. There has been some concern that this material can cause cancer or mutations but there is not enough data to make an assessment. Substance accumulation, in the human body, may occur and may cause some concern following repeated or long-term occupational exposure. Based on experience with similar materials, there is a possibility that exposure to the material may reduce fertility in humans at levels which do not cause other toxic effects. Prolonged or repeated skin contact with benzoyl peroxide may result in allergic skin reactions even at diluted concentrations. Ingestion results in abdominal pain, low body oxygen and severe depression. Chronic effects of exposure include allergic reactions characterised by redness, itching, oozing, crusting, and scaling of the skin and asthmatic wheezing. Although it does not exhibit complete carcinogenic or tumour-initiating activity, it has been associated with certain tumours of like papillomas and squamous cell carcinomas. Because it is soluble in fat but much less so in water, dimethyl phthalate can accumulate in body tissue. Therefore chronic exposure tends to be more important than acute exposure. Persistent exposure over a long period of time to peroxides produces allergic skin reactions (redness and scaling of the skin) and asthmatic wheezing. Exposure to ethylene glycol over a period of several weeks may





cause throat irritation, mild headache and low backache. These may worsen with increasing concentration of the substance. They may progress to a burning sensation in the throat, a burning cough, and drowsiness.

	Oral LD <sub>50</sub> mg/m <sup>3</sup>	Dermal LD <sub>50</sub> mg/m <sup>3</sup>	Inhalation LC <sub>50</sub> mg/L
Dibenzoyl Peroxide	6400	> 1000	
Dimethyl Phthalate	200 – 2000	> 2000	
Ethylene Glycol	3.58 – 12.7	9530	100.2 /8h

## Section 12 - Ecological Information

Very toxic to aquatic life with long lasting effects. Harmful to terrestrial vertebrates. Wastes resulting from use of the product must be disposed of on site or at approved waste sites. Do NOT discharge to sewer or waterway

	Fish mg/L	Crustacea mg/L	Algae mg/L
Dibenzoyl Peroxide	LC 50 0.06	EC <sub>50</sub> 0.11	EC <sub>50</sub> 0.042
			NOEC 0.02
Dimethyl Phthalate	LC 50 16	EC <sub>50</sub> 33	EC <sub>50</sub> 0.96
			BCF 100
			EC <sub>10</sub> 0.28
			NOEC 0.2
Ethylene Glycol	LC 50 72 - 860	EC <sub>50</sub> 100	EC <sub>50</sub> 3 - 536
		NOEC 1	

Persistance	Persistance	Bioaccumulation	Mobility
			LOW
_	_	_	LOW
_	_	_	HIGH
	Persistance H <sub>2</sub> O/ Soil LOW LOW LOW	H₂O/ Soil         Air           LOW         LOW           LOW         LOW	H2O/ Soil         Air           LOW         LOW         LOW           LOW         LOW         LOW

# Section 13 - Disposal Considerations

Containers may still present a chemical hazard/ danger when empty. Return to supplier for reuse/ recycling if possible. Otherwise: If container cannot 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.

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 or consult manufacturer for recycling options. Consult Land Waste Authority for disposal. Bury or incinerate residue at an approved site. 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



HAZCHEM

1W

# SOUDAL

# SAFETY DATASHEET

5.2							
None							
3108							
ORGANIC PEROXIDE, TYPE E, SOLID							
122 274 323							
500 g							
5.2							
None							
3108							
A20							
570							
25 Kg							
570							
10 Kg							
Passenger & Cargo Limited Quantity							
Forbidden							
Forbidden							
ORGANIC PEROXIDE, TYPE E, SOLID							
5.2							
None							
3108							
F-J, S-R							
122 274							
500 g							

Yes

**ORGANIC PEROXIDE, TYPE E, SOLID** 

# Section 15 - Regulatory Information

Marine pollutant

Shipping Name

HSNO approval number and Group Standard: HSR002629 Organic Peroxides

# Group Standard conditions and other regulations:

Condition	Requirement	
SDS	Safety data sheet must be available to a person handling the substance within 10 minutes.	
Emergency plan	Required when quantities exceed 500Lt	
Certified handler	Not required	
Tracking	Not applicable	
Bunding and secondary containment	Secondary containment is required based on pack size and total volume	
Signage	Required when present in quantity 250 L.	
Compliance certificate	Not required	



Hazardous Atmosphere zone			Not required	
Fire extinguisher			2 required when quantities exceed 250 Lt	
National Invent	ories			
Australia	AICS	Y	es	
Canada	DSL	Y	es	
Canada	NDSL	N	No	
China	IESCS	Y	Yes	
Europe	EINECS	Y	es	
Japan	ENCS	Y	es	
Korea	KECI	Y	es	
New Zealand	NZIoC	Y	es	

# Section 16 – Other Information

PICCS

TSCA

#### **Revision History**

Philippines

USA

March 2019

Origination

Yes

Yes

# Abbreviations:

Abbreviation	Description		
CAS number	Number assigned to chemical in the Chemical Abstracts Service registry		
HAZCHEM code	Code used by fire-fighters to determine correct method of action in the case of fire		
HSNO	Hazardous Substances and New Organisms (Act)		
ICAO Technical Instructions	International Civil Aviation Organization Technical Instructions		
IMDG code	International Maritime Dangerous Goods code controlled by the International Maritime Organization (IMO)		
LC <sub>50</sub>	Lethal concentration 50% - concentration fatal to 50% of the tested population		
LD <sub>50</sub>	Lethal dose 50% - dose fatal to 50% of the tested population		
NZS 5433	New Zealand Standard 5433 (Standard for the Transport of Dangerous Goods on Land)		
SDS	Safety data sheet		
STEL	Short term exposure limit		
TWA	Time weighted average (typically measured as 8 hours)		
UN number	United nations number		
WES	Workplace exposure standard		

#### References

Chemical properties and HSNO classifications derived from the New Zealand chemical classification information database (CCID).www.epa.govt.nz.

Workplace exposure limits derived from Workplace Exposure Standards and Biological Exposure Indices 9th Edition.

The information provided on this SDS is correct to the best of our knowledge, information and belief at the date of its publication. The information given is designed only as a guidance for safe handling, use, processing, storage, transportation, disposal and release and is not to be considered as a warranty or quality specification. The information relates only to the specific material designated and may not be valid for such material in combination with any other material or in any process, unless specified in the text.



# This SDS was prepared by Collievale Enterprises Ltd in accord with the Hazardous Substances (Safety Data Sheets) Notice 2017 <a href="http://www.collievale.com">http://www.collievale.com</a> Phone +64 7 5432428 <a href="http://www.collievale.com">End of MSDS</a>

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