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1. PRODUCT IDENTIFICATION

Product Name Other Names	Battery Fluid, Acid Battery Fluid, Acid, Sulphuric Acid 1260, Electrolyte, Sulphuric Acid, Battery Acid, Sulphuric acid (Electrolyte)
Use	Electrolyte for lead-acid batteries
Supplier Name and Address	Century Yuasa Batteries 259 Church St, Onehunga, Auckland 1643
Telephone	0800 93 93 93
Emergency (24 Hours)	0274 686 673
Relevant identified uses	Electrolyte for lead-acid batteries

2. HAZARD(S) IDENTIFICATION

Considered a Hazardous Substance according to the criteria of the New Zealand Hazardous Substances New Organisms (HSNO) legislation. Classified as Dangerous Goods for transport purposes.

Signal Word DANGER

GHS Classification Metal Corrosion Category 1, Acute Toxicity (Inhalation) Category 2, Skin Corrosion/Irritation Category 1A, Serious Eye Damage Category 1

HSNO Classification GHS Label Elements

6.1D (inhalation), 6.1E (Oral), 6.7A, 6.9A (Inhalation), 8.1A, 8.2B, 8.3A, 9.1C, 9.1D



Corrosive Acute toxicity

IN THE EVENT OF EXPOSURE TO BATTERY FLUID, ACID

Hazard Statements	H290	May be corrosive to metals	H330	Fatal if inhaled
	H302	Harmful if swallowed		
	H314	Causes severe skin burns and eye damage		

IN THE EVENT OF EXPOSURE TO INTERNAL COMPONENTS

Precautionary Statements	Prevention		Response	
	P101	If medical advice is needed, have product container or label at hand.	P301+P330+P331	IF SWALLOWED: Rinse mouth. Do NOT induce vomiting.
	P102	Keep out of reach of children	P303+P361+P353	IF ON SKIN (or hair): Take off immediately
	P103	Read label before use.		all contaminated clothing. Rinse skin with water/ shower.
	P234	Keep only in original container.	P305+P351+P338	IF IN EYES: Rinse cautiously with water for
	P260	Do not breathe dust / fume / gas / mist / vapours / spray.		several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.
	P271	Use only outdoors or in a well- ventilated area.	P304+P340	IF INHALED: Remove person to fresh air and keep comfortable for breathing.
	P280	Wear protective gloves / protective clothing / eye protection / face	P390	Absorb spillage to prevent material damage.
		protection	<u>Storage</u>	
	Disposal		P406	Store in a corrosion resistant container with resistant inner Liner
	P501	Dispose of contents, container to authorised chemical landfill or if organic, to high temperature incineration	P403+P233	Store in a well-ventilated place. Keep container tightly closed.
	<u>Recycle</u>			
		Refer to section 13		



3. COMPOSITION, INFORMATION ON INGREDIENTS

Ingredient	Identification	Content % weight
Sulphuric Acid <51% (H ₂ SO ₄)	CAS 7664-93-9	33-36%
Water	-	64-67%
de		

4. FIRST AID MEASURES

DESCRIPTION OF FIRST AID MEASURES

Eye contact	 If, Sulphuric acid 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, Sulphuric acid comes in contact with skin or hair: Immediately flush body and clothes with large amounts of water, using safety shower if available. Quickly remove all contaminated clothing, including footwear. Wash skin and hair with running water. Continue flushing with water until advised to stop by the Poisons Information Centre. Transport to hospital, or doctor.
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. Inhalation of vapours or aerosols (mists, fumes) may cause lung oedema. Corrosive substances may cause lung damage (e.g. lung oedema, fluid in the lungs). As this reaction may be delayed up to 24 hours after exposure, affected individuals need complete rest (preferably in semi-recumbent posture) and must be kept under medical observation even if no symptoms are (yet) manifested. Before any such manifestation, the administration of a spray containing a dexamethasone derivative or beclomethasone derivative may be considered. This must definitely be left to a doctor or person authorised by him/her.
Ingestion	 For advice, contact a Poisons Information Centre or a doctor at once. Urgent hospital treatment is likely to be needed. 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. Transport to hospital or doctor without delay.
MEDICAL ATTENTION A	ND SPECIAL TREATMENT. Indication of any immediate medical attention and special treatment needed
Treat symptomatically.	 For acute or short term repeated exposures to strong acids: Airway problems may arise from laryngeal edema and inhalation exposure. Treat with 100% oxygen initially. Respiratory distress may require cricothyroidotomy if endotracheal intubation is contraindicated by excessive swelling Intravenous lines should be established immediately in all cases where there is evidence of circulatory compromise. Strong acids produce a coagulation necrosis characterised by formation of a coagulum (eschar) as a result of the desiccating action of the acid on proteins in specific tissues.
Ingestion:	 Immediate dilution (milk or water) within 30 minutes post ingestion is recommended. DO NOT attempt to neutralise the acid since exothermic reaction may extend the corrosive injury. Be careful to avoid further vomit since re-exposure of the mucosa to the acid is harmful. Limit fluids to one or two glasses in an adult. Charcoal has no place in acid management. Some authors suggest the use of lavage within 1 hour of ingestion.
Skin:	 Skin lesions require copious saline irrigation. Treat chemical burns as thermal burns with non-adherent gauze and wrapping. Deep second-degree burns may benefit from topical silver sulphadiazine.
Eye:	• Eye injuries require retraction of the eyelids to ensure thorough irrigation of the conjuctival cul-de-sacs. Irrigation



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should last at least 20-30 minutes. DO NOT use neutralising agents or any other additives. Several litres of saline are required.

- Cyclopaedic drops, (1% cyclopentolate for short-term use or 5% homatropine for longer term use) antibiotic
- drops, vasoconstrictive agents or artificial tears may be indicated dependent on the severity of the injury.
- Steroid eye drops should only be administered with the approval of a consulting ophthalmologist).

5. FIRE FIGHTING MEASURES







7. HANDLING AND STORAGE

Safe Handling	 DO NOT a Use in a w Handle an When han Always wa Always wa Use good 	Avoid all personal contact and wear protective clothing when risk of exposure occurs. DO NOT allow clothing wet with material to stay in contact with skin Use in a well-ventilated area, avoid generating and breathing mist Handle and open container with care and keep containers securely sealed when not in use When handling, DO NOT eat, drink or smoke. Always wash hands with soap and water after handling. Work clothes should be laundered separately. Always wash hands with soap and water after handling. Work clothes should be laundered separately. Use good occupational work practice. Observe manufacturer's storage and handling recommendations contained within this SDS.				
Storage	foodstuff c Protect con Floors sho DO NOT s	Store in original containers and store in a cool, dry, well-ventilated area away from incompatible materials and foodstuff containers. Protect containers against physical damage and check regularly for leaks Floors should be covered or coated with acid resistant material. DO NOT store in pits, depressions, basements or areas where vapours may be trapped Attacks some plastics, rubber and coatings				
Suitable container	Check that	as recommended by containers are clearl yethylene, Polypropyl	y labelled	tainers are suitable		
Storage incompatibili	 Glass, Polyethylene, Polypropylene or Poly-lined containers are suitable torage incompatibility Is a strong oxidiser Reacts with water or steam Reacts violently with many substances including reducing agents, combustible materials, organic substances, alkalis, ammonium tetraperoxochromate, aniline, 1,2-ethanediamine, ethanolamine, isoprene, mesityl oxide, endo-norbanecarboxylic acid ethyl ester, perchlorates, sodium carbonate, zinc chlorate Reacts, possibly causing ignition or explosion, with many substances, including non-oxidising mineral acids, organ acids, bases, reducing agents, acetic anhydride, acetone cyanohydrin, acetonitrile, acrolein, acrylates, acrylonitrile alcohols, aldehydes, alkylene oxides, allyl alcohol, allyl chloride, substituted, allyls, 2-aminoethanol, ammonium hydroxide, bromine pentafluoride, n-butyraldehyde, caprolactam solution, carbides, caesium acetylene carbide, chlorine trifluoride, chlorosulfonic acid, cresols, cuprous nitride, diisobutylene, ethylene carbide, inchorate, potrassium permanganate, beta-propiolactone, proylene oxide, pyridine, rubidium acetylene, solium chlorate, sodium hydroxide, styrene monomer, zinc phosphorus, picrates, potassium chlorate, sodium, sodium chlorate, sodium hydroxide, styrene monomer, zinc phosphide Increases the explosive sensitivity of nitromethane Incompatible with 2-amino-5-nitrothiazole, 2-aminothiazole, ammonia, aliphatic amines, alkanolamines, amides, organic anhydrides, isocyanate, vinyl acetate, alkylene oxides, epichlorohydrin. Attacks some plastics, rubber and coatings Reacts with metals to produce flammable hydrogen gas 					mesityl oxide, endo nineral acids, organic rylates, acrylonitrile, anol, ammonium cetylene carbide, ne cyanohydrin, heptafluoride, iron, tric acid, p- crates, potassium etylene, silver
= May be stor	red together	() = May be stor	ed together with spo	ecific preventions	×= Must not b	e stored together
E 3	×		V V V			
FLAMMABLES	EXPLOSIVES		OXIDISERS	HARMFUL	IRRITANT	CORROSIVE

8. EXPOSURE CONTROLS, PERSONAL PROTECTION

AUSTRALIAN EXPOSURE STANDARDS (Occupational Exposure Limits)

Ingredient	Material name	TWA	STEL
Sulphuric Acid (H2SO4)	Sulphuric acid	1 mg/m3	3 mg/m3

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.



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PERSONAL PROTECTION: Not normally required; however if in contact with internal components:-



Respirator Type Where the concentration of gas / particulates in the breathing zone, approaches or exceeds the "Exposure Standard" (or ES), respiratory protection is required. ~. .ff:

Required Minimum Protection Factor	Half-Face Respirator	Full-Face Respirator	Powered Air Respirator
up to 10 x ES	P1 Air-line*	-	PAPR-P1 -
up to 50 x ES	Air-line**	P2	PAPR-P2
up to 100 x ES	-	P3	-
		Air-line*	-
100+ x ES	-	Air-line**	PAPR-P3

Negative pressure demand Continuous flow



Clothing

- When handling corrosive liquids, wear trousers or
- overalls outside of boots, to avoid spills entering boots. Overalls or PVC protective suit may be required if exposure severe.



Other Protection Eyewash unit.

- Barrier cream.
- Skin cleansing cream.

PHYSICAL AND CHEMICAL PROPERTIES



Eye Protection

- Safety glasses with unperforated side shields may be used where continuous eye protection is desirable, as in laboratories;
- Spectacles are not sufficient where complete eye protection is needed such as when handling bulkquantities, where there is a danger of splashing, or if the material may be under pressure.
- Chemical goggles whenever there is a danger of the material coming in contact with the eyes; goggles must be properly fitted.
- Contact lenses may pose a special hazard; soft contact lenses may absorb and concentrate irritants. Medical and first-aid personnel should be trained in their removal and suitable equipment should be readily available.

Glove Type

Wear chemical protective gloves, e.g. PVC

Foot wear

Wear safety footwear or safety gumboots e.g. Rubber

Appearance	Clear colourless, mobile liquid that mixes with water.			
Odour	Not Available	Lower explosive limits	Not Available	
Odour threshold	Not Available	Vapour pressure (kPa)	13 to 22 mmHg @ 25 °C	
рН	<1	Vapour density (Air = 1)	Not Available	
Melting point / freezing point (°C)	95 °C / -7 to -70 °C	Relative density (Water = 1)	1.2-1.3 (Sulphuric acid electrolyte) @ 25 °C	
Initial boiling point and boiling range (°C)	95 °C (Sulphuric acid electrolyte)	Solubility in water (g,L)	Immiscible	
Flash point	Non-flammable	Partition coefficient: n- octanol/water	Not Available	
Evaporation rate	Not Available	Auto-ignition temperature	Not Available	
Flammability	Not Applicable	Decomposition temperature (°C)	Not Available	
Upper, lower flammability or explosive limits	Not Applicable	Viscosity	Not Available	

10. STABILITY AND REACTIVITY Reactivity See section 7 and this section under Chemical stability Is a strong oxidiser Reacts violently with many substances including reducing agents, combustible materials, organic substances, alkalis Acids often catalyse (increase the rate of) chemical reactions. Inorganic acids are generally soluble in water with the release of hydrogen ions. The resulting solutions have pH of less than 7.0. The dissolution of inorganic acids in water or the dilution of their concentrated solutions with additional water may generate significant heat. The addition of water to inorganic acids often generates sufficient heat in the small region of mixing to cause some of the water to boil explosively. The resulting "bumping" can spatter the acid. Possibility of hazardous See section 5 & 7 reactions Reacts, possibly causing ignition or explosion, with many substances, including non-oxidising mineral acids, phosphorus, picrates, potassium chlorate, potassium permanganate, beta-propiolactone, propylene oxide. pyridine, rubidium acetylene, silver permanganate, sodium, sodium chlorate, sodium hydroxide, styrene monomer, zinc phosphide Reacts with mild steel, galvanised steel / zinc, active metals, including such structural metals as aluminium and iron, to release hydrogen, a flammable gas. Inorganic acids neutralise chemical bases (for example: amines and inorganic hydroxides) to form salts neutralisation can generate dangerously large amounts of heat in small spaces. Inorganic acids generate flammable and/or toxic gases in contact with dithiocarbamates, isocyanates, mercaptans, nitrides, nitriles, sulphides, and strong reducing agents. Additional gas-generating reactions occur with sulphites, nitrites, thiosulphates (to give H2S and SO3), dithionites (SO2), and even carbonates. Reacts with cyanide compounds to release gaseous hydrogen cyanide



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Incompatible materials	 See section 7 Avoid heat, sparks , open flame, and other ignition sources Avoid storage with oxidisers, alkalis, reducing agents, common metals and their alloys Avoid any contamination of this material as it is very reactive and any contamination is potentially hazardous
Chemical stability	 Unstable in the presence of incompatible materials. Product is considered stable. Hazardous polymerisation will not occur Contact with alkaline material liberates heat
Hazardous decomposition products	 See section 5 Sulphuric acid may decompose to sulphur trioxide, carbon monoxide, sulphuric acid mist, sulphur dioxide and hydrogen.

11. TOXICOLOGICAL INFORMATION ACUTE EFFECTS

No adverse health effects expected if the product is handled in accordance with this safety Data sheet and the product Label. Symptoms or effects that may arise if the product is mishandled and overexposure occurs are:-

Inhaled	Corrosive acids can cause irritation of the respiratory tract, with coughing, choking and mucous membrane damage. There may be dizziness, headache, nausea and weakness. Inhalation of aerosols (mists, fumes), generated by the material during the course of normal handling, may produce toxic effects; these may be fatal. The material can cause respiratory irritation in some persons.
Ingestion	Considered an unlikely route of entry in commercial/industrial environments Ingestion of acidic corrosives may produce burns around and in the mouth, the throat and oesophagus. Immediate pain and difficulties in swallowing and speaking may also be evident. The material is considered to be harmful by all exposure routes The liquid is highly discomforting and corrosive if swallowed Ingestion may result in nausea, abdominal irritation, pain and vomiting
Skin contact	Skin contact with acidic corrosives may result in pain and burns; these may be deep with distinct edges and may heal slowly with the formation of scar tissue. Skin contact is not thought to produce harmful health effects (as classified under EC Directives using animal models). Systemic harm, however, has been identified following exposure of animals by at least one other route and the material may still produce health damage following entry through wounds, lesions or abrasions. Open cuts, abraded or irritated skin should not be exposed to this material Entry into the blood-stream, via, cuts, abrasions or lesions, may produce systemic injury with harmful effects. The liquid is highly discomforting and corrosive to the skin and is capable of causing ulceration and severe burns if exposure is prolonged, even minor exposure is highly discomforting The material may cause skin irritation after prolonged or repeated exposure and may produce on contact skin redness, swelling, the production of vesicles, scaling and thickening of the skin.
Eye	This material can cause eye irritation and damage in some persons If applied to the eyes, this material causes severe eye damage. Direct eye contact with acid corrosives may produce pain, tears, sensitivity to light and burns. The material may produce moderate eye irritation leading to inflammation. Repeated or prolonged exposure to irritants may produce conjunctivitis. Mild burns of the epithelia generally recover rapidly and completely.
Chronic effects	Repeated or prolonged exposure to acids may result in the erosion of teeth, swelling and/or ulceration of mouth lining. Irritation of airways to lung, with cough, and inflammation of lung tissue often occurs. Substance accumulation, in the human body, may occur and may cause some concern following repeated or long-term occupational exposure. Principal routes of exposure are usually by skin contact with the material, eye contact and inhalation of vapour. The material is considered to be harmful by all exposure routes and contact may cause rapid tissue destruction As with any chemical product, contact with unprotected bare skin; inhalation of vapour, mist or dust in work place atmosphere; or ingestion in any form, should be avoided by observing good occupational work practice
Sulphuric Acid <i>(undiluted)</i>	Asthma-like symptoms may continue for months or even years after exposure to the material ceases. This may be due to a non-allergenic condition known as reactive airways dysfunction syndrome (RADS) which can occur following exposure to high levels of highly irritating compound. Key criteria for the diagnosis of RADS include the absence of preceding respiratory disease, in a non-atopic individual, with abrupt onset of persistent asthma-like symptoms within minutes to hours of a documented exposure to the irritant. A reversible airflow pattern, on spirometry, with the presence of moderate to severe bronchial hyper-reactivity on methacholine challenge testing and the lack of minimal lymphocytic inflammation, without eosinophilia, have also been included in the criteria for diagnosis of RADS. 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. Industrial bronchitis, on the other hand, is a disorder that occurs as result of exposure due to high concentrations of irritating substance (often particulate in nature) and is completely reversible after exposure ceases. The disorder is characterised by dyspnoea, cough and mucus production
	WARNING: For inhalation exposure ONLY: This substance has been classified by the IARC as Group 1: CARCINOGENIC TO HUMANS Occupational exposures to strong inorganic acid mists of sulphuric acid:



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Acute Toxicity	Skin Irritation / Corrosion	Serious Eye Damage / Irritation	Respiratory Or Ski Sensitisation	n Mutagenicity	Carcinogenicity	Reproductivity	Stot - Single Exposure	Stot - Repeated Exposure	Aspiration Hazard
✓	✓	 ✓ 	0	1	 Image: A set of the set of the	 ✓ 	✓	✓	Û
		required to mak	() =	n available 🛛 🗙 Data Not Avai			ot fill the cri	teria for classifi	cation
12. EC	OLOGICAL	INFORMATIO	N						
Ecotoxicity		 Prevent, by any means available, spillage from entering drains or water courses. DO NOT discharge into sewer or waterways. DO NOT discharge into sewer or waterways. May cause long term adverse effects in the environment Avoid contaminating waterways. The product is highly acidic. If large spills occurred a water pH drop could be responsible for an environmental effect on aquatic organisms. Ecotoxicity data for Sulphuric Acid Mosquito fish LC50 42mg/L/96hr Shore crab LC50 70-80mg/L/48hr 							
		Hooknose fish		-C50 80-90mg	J/L/4011	Cockle		LC30 200-	500mg/L/48hr
Degradability Bio-accumula		No Data availabl No Data availabl	-						
Potential			o for an ingrou						
Mobility in So	bil	 During trans materials 	sport through t	he soil, sulfuri	c acid can dis	solve some of	the soil ma	terial, in particu	lar carbonate-bas
Other Advers	e Effects	No Data availabl	e for all ingred	lients					
13. DIS	POSAL CO	NSIDERATION	IS						
Safe Handling Disposal	g &	Dispose in accordance with federal, state or local regulations.							
		 Return to supplier for reuse/ recycling if possible. Use soda ash or slaked lime to neutralise <u>Otherwise:</u> 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, and 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. 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 possible to reclaim the product by filtration, distillation or some other means. Shelf 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. 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. Consult manufacturer for recycling options or consult local or regional waste management authority for disposal if no suitable treatment or disposal facility can be identified. Treat and neutralise at an approved treatment plant. Treatment should involve: Neutralisation with soda-ash or soda-lime followed by: burial in a land-fill specifically licenced to accept chemical and / or pharmaceutical wastes or lncineration in a licenced apparatus (after admixture with suitable combustible material). Decontaminate empty containers with 5% aqueous sodium hydroxide or soda ash, followed by water. Observe all label safeguards until containers are cleaned and destroyed. 							
Environmenta Regulations	al	Dispose in aRefer to see	accordance wi ction 15	th federal, sta	te or local reg	ulations.			
14. TR	ANSPORT II	NFORMATION							
JN Number		2796							
Proper Shipp	ing Name	Battery fluid, acid	d or Sulfuric ad	id with not mo	ore than 51 pe	rcent acid 🗸			

Proper Shipping Name	Battery fluid, acid or Sulfuric acid with not more than 51 percent acid				
Transport Hazard Class	Class: 8	Sub risk: Not Applicable			
Packing group	II		5. 11 13		
Environmental Hazards	No relevant data				
Special Precautions	Not Applicable Limited quantity	1L	CORROSIVE		
Additional Information	Marine Pollutant:	Yes	0		
Hazchem Code	2R				



15. REGULATORY INFORMATION

SAFETY, HEALTH AND ENVIRONMENTAL REGULATIONS, SPECIFIC FOR THE SUBSTANCE OR MIXTURE

This substance is to be managed using the conditions specified in the applicable Group Standard					
HSR002493 Additiv	ditives, Process Chemicals and Raw Materials (Corrosive) Group Standard 2006 ditives, Process Chemicals and Raw Materials (Corrosive, Toxic [6.7]) Group Standard 2006 ditives, Process Chemicals and Raw Materials (Toxic [6.1 + 6.7]) Group Standard 2006 ditives, Process Chemicals and Raw Materials (Toxic [6.1]) Group Standard 2006				
Sulphuric Acid CAS 7664-93-9 is found on the following regulatory Lists		"International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Monographs", "International Air Transport Association (IATA) Dangerous Goods Regulations - Prohibited List Passenger and Cargo Aircraft", "New Zealand Inventory of Chemicals (NZIoC), New Zealand Workplace Exposure Standards", New Zealand Hazardous and New Organisms (HSNO) Act – Classification of Chemicals"			
Location Test Certificate		Subject to Regulation 55 of the Hazardous Substances (Classes 1 to 5 Controls) Regulations, a location test certificate is required when quantity greater than or equal to those indicated below are present			
Hazard Class		Not applicable			
Quantity beyond which controls apply for closed containers		Not applicable			
Quantity beyond which controls apply when use occurring in open containers		Not applicable			
Approved Handler	must be	Subject to Regulation 9 of the Hazardous Substances (Classes 6, 8, and 9 Controls) Regulations, the substance must be under the personal control of an Approved Handler when present in a quantity greater than or equal to those indicated below			
Class of Substance	Quanti	Quantities			
6.1	, ,	Any quantity			
6.7A	10 kg or more, if solid 10 L or more, if liquid				
8.1A	N/A				
8.2B	Any qua	Any quantity			
9.1C, 9.2D	9.1C, 9.2D Any quantity				
16. OTHER RELEVANT INFORMATION					

Revision Information	Revision No	Date	Description	
	1	09/02/2016	Initial SDS creation	
Abbreviations	CAS #	Chemical Abstract Service Number – used to uniquely identify chemical compounds		
	IARC	International Agency for Research on Cancer		
	HSNO	Hazardous Substances and New Organisms ((HSNO) Act		
	LC50	 LC50 Lethal Concentration- toxicity of the surrounding medium that will kill half of the sample population of a spec animal in a specified period through exposure via inhalation (respiration) SDS Safety Data Sheet- (SDS), previously called a Material Safety Data Sheet (SDS), 		
	SDS			
	TGA	Therapeutic	Goods Administration	