

SDS no. 0DH7TSDY • Version 1.0 • Date of issue: 2023-05-31

SECTION 1: Identification

GHS Product identifier

Product name

SULFURIC ACID 15-51%

Recommended use of the chemical and restrictions on use

Fertilizers, chemicals, dyes and pigments, etchant, alkylation catalyst, electroplating baths, iron and steel, rayon and film, industrial explosives, non-ferrous metallurgy, analytical reagent and laboratory reagent.

Supplier's details

Name Address	ChemSupply Australia Pty Ltd 38-50 Bedford Street 5013 Gillman South Australia Australia
Telephone email	08 8440 2000 www.chemsupply.com
Emergency phone number	
	CHEMCALL 1800 127 406 (Australia) / +64-4-917-9888 (International)

SECTION 2: Hazard identification

General hazard statement

Classified as dangerous goods according to the Australian Dangerous Goods Code (ADG).

Classified as Hazardous according to the Globally Harmonised System of classification and labelling of Chemicals (GHS) including Work, Health and Safety regulations, Australia.

Classification of the substance or mixture

GHS classification in accordance with: UN GHS revision 7

- Serious eye damage/eye irritation, Cat. 1
- Corrosive to metals, Cat. 1
- Skin corrosion/irritation, Cat. 1A

GHS label elements, including precautionary statements

Pictograms



Signal word Danger Hazard statement(s) H290 May be corrosive to metals H314 Causes severe skin burns and eye damage Precautionary statement(s) P260 Do not breathe dust/fume/gas/mist/vapors/spray. P280 Wear protective gloves/protective clothing/eye protection/face protection. P301+P330+P331 IF SWALLOWED: Rinse mouth. Do NOT induce vomiting. P303+P361+P353 IF ON SKIN (or hair): Take off immediately all contaminated clothing. Rinse skin with water [or shower]. P304+P340 IF INHALED: Remove person to fresh air and keep comfortable for breathing. P305+P351+P338 IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing. P310 Immediately call a POISON CENTER/doctor/physcian P363 Wash contaminated clothing before reuse. P390 Absorb spillage to prevent material-damage. P405 Store locked up. P406 Store in a corrosive resistant/... container with a resistant inner liner. P501 Dispose of contents/container to an approved waste disposal facility

SECTION 3: Composition/information on ingredients

Mixtures

Molecular weight: 98.08

Components		
Component	Concentration	
Water (CAS no.: 7732-18-5; EC no.: 231-791-2)	49 - 85 % (weight)	
CLASSIFICATIONS: No data available. HAZARDS: No data available.		
Sulfuric acid (CAS no.: 7664-93-9; EC no.: 231-639-5; Index no.: 016-020-00-8)	15 - 51 % (weight)	
CLASSIFICATIONS: Skin corrosion/irritation, Cat. 1A. HAZARDS: H314 - Causes severe skin burns and eye damage. [SCLs/M-factors/ATEs]: Skin Corr. 1A; H314: C ≥		
15 %; Skin Irrit. 2; H315: 5 % \leq C $<$ 15 %; Eye Irrit. 2; H319: 5 % \leq C $<$ 15 %		

SECTION 4: First-aid measures

Description of necessary first-aid measures

General advice	First Aid Facilities: Maintain eyewash fountain and drench facilities and normal washroom facilities in work area.
	For advice, contact a Poisons Information Centre (e.g. phone Australia 13 11 26; New Zealand 0800 764 766) or a doctor (at once).
If inhaled	If inhaled, remove from contaminated area. Apply artificial respiration if not breathing.
In case of skin contact	Immediately remove contaminated clothing and wash affected area with water for at least 15 minutes. Ensure contaminated clothing is washed before re-use. Seek immediate medical advice /attention.

In case of eye contactf in eyes, hold eyelids apart and flush eye continuously with running water. Continue
flushing until advised to stop by a Poisons Information Centre (e.g. phone Australia 13
11 26; New Zealand 0800 764 766) or a doctor, or for at least 15 minutes.

If swallowed If swallowed, do NOT induce vomiting.

Personal protective equipment for first-aid responders

WARNING: It may be hazardous to the person providing aid to give mouth-to-mouth resuscitation when the inhaled material is toxic, infectious or corrosive.

No action shall be taken involving any personal risk or without suitable training. If it is suspected that fumes are still present, the rescuer should wear an appropriate mask or self-contained breathing apparatus. It may be dangerous to the person providing aid to give mouth-to-mouth resuscitation. Wash contaminated clothing thoroughly with water before removing or wear gloves.

Most important symptoms/effects, acute and delayed

The most important known symptoms and effects are described in the labelling (see section 2.2) and/or in section 11

Indication of immediate medical attention and special treatment needed, if necessary

For advice in an emergency, contact a Poisons Information Centre (Phone Australia 131 126) or a doctor at once.

SECTION 5: Fire-fighting measures

Suitable extinguishing media

When material is not involved in fire: Do not use water on material itself. Use fire extinguishing media appropriate for surrounding environment. Use water spray, dry chemical, carbon dioxide, or appropriate foam.

Specific hazards arising from the chemical

Irritating and highly toxic fumes and gases, including oxides of sulfur. Reaction with water or steam may generate much heat which will increase the concentration of fumes in the air, and may produce toxic and corrosive fumes.

Material does not burn. Fire or heat will produce irritating, poisonous and/or corrosive gases. Containers may explode when heated. Contact with metals may evolve flammable hydrogen gas.

Special protective actions for fire-fighters

Fire fighters should wear full protective clothing and self-contained breathing apparatus (SCBA) operated in positive pressure mode. Fight fire from safe location.

SECTION 6: Accidental release measures

Personal precautions, protective equipment and emergency procedures

Wear respiratory protection. Avoid breathing vapours, mist or gas. Ensure adequate ventilation. Evacuate personnel to safe areas. For personal protection see section 8.

Methods and materials for containment and cleaning up

Neutralise with lime or sodium carbonate, adjust the pH to 6-10. For larger spills notify Emergency Services.

Small Spillages: Absorb or contain liquid with sand, earth or spill control material, or neutralize with sodium carbonate or other alakli material. Shovel up using non sparking tools and place in a labelled, sealable container for subsequent safe disposal. Put leaking containers in a labelled drum or overdrum.

SECTION 7: Handling and storage

Precautions for safe handling

Corrosive liquid. May produce severe burns. Avoid ingestion and inhalation of gas/fumes/vapour/spray mist. Attacks skin and eyes. Avoid contact with eyes, skin, or clothing. Keep locked up. Keep containers tightly closed when not in use. Use only with adequate ventilation. If ingested, seek medical advice immediately and show the container or the label. Wear suitable protective clothing, gloves and eye/face protection when mixing and using. Wash thoroughly after handling. Remove contaminated clothing and wash before reuse. Discard contaminated shoes. Ensure a high level of personal hygiene is maintained when using this product, that is, always wash hands before eating, drinking, smoking or using the toilet facilities. Keep away from incompatibles such as oxidizing agents, combustible materials, organic materials, metals, acids, alkalis, moisture/water. Contact with water will generate heat. When diluting, always add the acid to water; never add water to the acid. May corrode metallic surfaces.

Conditions for safe storage, including any incompatibilities

Store in tightly closed containers, in a cool, dry, well-ventilated corrosives area with acid resistant floors. Store away from incompatible substances, such as water, alkaline substances, oxidizing agents, and reducing agents. Store away from combustible substances, sources of ignition and heat.

Extremely corrosive in presence of aluminium, of zinc (50% sulfuric acid). Concentrated acid is non-corrosive to lead and mild steel, but diluted acid attacks most metals. Highly corrosive in presence of steel, of copper (20% sulfuric acid). Slightly corrosive to extremely corrosive in presence of stainless steel(304), of stainless steel(316) (20-50% sulfuric acid). Corrosive to most metals in the presence of moisture, liberating explosive hydrogen gas.

Store at room temperature (15 to 23 °C recommended). Protect from freezing.

SECTION 8: Exposure controls/personal protection

Control parameters

CAS: 7664-93-9 (EC: 231-639-5)

Sulfuric acid

ACGIH (USA): 0.2 mg/m3, (Thor.) TLV® inhalation; 0.2 mg/m3 TWA inhalation; AU/SWA (Australia): 3 mg/m3 STEL inhalation; 1 mg/m3 TWA inhalation; NIOSH (USA): 1 mg/m3 REL inhalation;

Appropriate engineering controls

Use ventilation adequate to keep exposures (airborne levels of dust, fume, vapor, gas, etc.) below recommended exposure limits.

Individual protection measures, such as personal protective equipment (PPE)

Eye/face protection

The use of a face shield, chemical goggles or safety glasses with side shield protection as appropriate. Must comply with Australian Standards AS 1337 and be selected and used in accordance with AS 1336.

Skin protection

Clean impervious clothing should be worn. Clothing for protection against chemicals should comply with AS 3765 Clothing for Protection Against Hazardous Chemicals.

Hand Protection: Normally not required but if in doubt ensure hand protection should complies with AS 2161, Occupational protective gloves - Selection, use and maintenance.

Body protection

Suitable protective workwear, e.g. cotton overalls buttoned at neck and wrist is recommended. Chemical resistant apron is recommended where large quantities are handled.

Respiratory protection

If engineering controls are not effective in controlling airborne exposure then an approved respirator with a replaceable vapor/ mist filter should be used. Refer to relevant regulations for further information concerning respiratory protective requirements. Reference should be

made to Australian Standards AS/ NZS 1715, Selection, Use and Maintenance of Respiratory Protective Devices; and AS/NZS 1716, Respiratory Protective Devices, in order to make any necessary changes for individual circumstances.

SECTION 9: Physical and chemical properties

Basic physical and chemical properties

Physical state Appearance Color Odor Odor threshold Melting point/freezing point Boiling point or initial boiling point and boiling range Flammability Lower and upper explosion limit/flammability limit Flash point Explosive properties

Auto-ignition temperature Decomposition temperature Oxidizing properties pH

Kinematic viscosity Solubility

Partition coefficient n-octanol/water (log value) Vapor pressure

Evaporation rate Density and/or relative density

Relative vapor density

Particle characteristics No data available. Liquid Clear, colourless liquid. No data available. Odourless. >1 ppm (Sulfuric Acid). -8.6 °C (15%); -14 °C (20%); -46 °C (33%); -33 °C (51%). 102.6 °C (15%); 104 °C (20%); 109 °C (33%); 125 °C (51%). No data available. No data available. No data available. Contact with most metals causes formation of flammable and explosive hydrogen gas. However, the risk is reduced due to the weaker concentration of Sulfuric Acid present. Exothermic reaction with water. Containers may explode when heated or if contaminated with water. Slightly explosive in presence of oxidizing materials. Mixtures of sulfuric acid and any of the following can explode: p-nitrotoluene, pentasilver trihydroxydiaminophosphate, perchlorates, alcohols with strong hydrogen peroxide, ammonium tetraperoxychromate. mercuric nitrite, potassium chlorate, potassium permanganate with potassium chloride. Nitramide decomposes explosively on contact with concentrated sulfuric acid. 1,3,5-Trinitrosohexahydro-1,3,5-triazine + sulfuric acid causes explosive decomposition. No data available. 340 °C (sulfuric acid) No data available. <0.3 [Acidic].
pH of 1.0 N solution (~5.0%): 0.3;
pH of 0.1 N solution (~0.5%): 1.2;
pH of 0.01 N solution (~0.05%): 2.1. Viscosity: 21 mPas @ 25 °C (Sulfuric Acid). Solubility in Water: Soluble in water in all proportions. CAUTION: Always add the acid to water. Heat evolution due to mixing may cause explosive spattering. Solubility in Organic Solvents: Insoluble in methanol, diethyl ether, n-octanol. No data available. The highest known values are 2.3 kPa (17.535 mmHg) (@ 20 °C) (Water) and 0.1 kPa (1 mmHg) (@ 20 °C) (sulfuric acid). Weighted average: 14.03 mmHg (@ 20 °C) (20%). No data available. Specific Gravity: 1.1 (15%); 1.142 (20%); 1.18 (25%); 1.25 (33.33%); 1.30 (40%); 1.40 (50%); 1.41 (51%). The highest known value is 3.4 (Air = 1) (Sulfuric acid). Weighted average: 1.18 (Air = 1) (20%); 2.01 (Air = 1) (50%).

Supplemental information regarding physical hazard classes No data available.

Further safety characteristics (supplemental)

No data available.

SECTION 10: Stability and reactivity

Reactivity

Stable under normal conditions of storage and handling.

Reacts with incompatible materials

Chemical stability

Stable under normal temperatures pressures and conditions of storage and handling. Concentrated solutions react violently with water, spattering and liberating heat.

Possibility of hazardous reactions

Concentrated solutions react violently with water, spattering and liberating heat. Corrosively attacks most metals liberating hydrogen gas, (potential explosion). Sulfuric acid reacts vigorously, violently or explosively with many organic and inorganic chemicals. Reacts with carbonates to generate carbon dioxide gas. Reacts with cyanides to form toxic hydrogen cyanide. Reacts with sulfides to form toxic hydrogen sulfide.

Conditions to avoid

Exposure to moist air, moisture, or water (Note: Use great caution in mixing with water due to heat evolution that causes explosive spattering. Always add the acid to water, never the reverse.), metals, excess heat, combustible materials, organic materials, oxidizers, amines, bases and incompatible materials.

Incompatible materials

Water, combustible materials, oxidizing agents, reducing agents, metals as powders, metals as non powders (yields hydrogen gas), metal alloys, metal compounds, acids, alkalis, organic materials, organic solvents, alkali metals, alkaline earth metals, alkaline earth compounds, alkali hydroxides solutions, chlorates, perchlorates, permanganates, carbides, cyanides, nitrides, sulfides, fulminates, picrates, nitrates, nitrites, halogens, halogen-halogen compounds, salts of oxyhalogenic acids, acetylides, oxides and hydrides, anilines, organic nitro compounds, peroxi compounds, acetic anhydride, acetone cyanhydrin, acetone + nitric acid, acetone + potassium dichromate, acrolein, allyl alcohol, allyl chloride, 2-aminoethanol, ammonia, ammonium triperchromate, n-butyraldehyde, diisobutylene, epichlorohydrin, ethylene diamine, ethylene glycol, ethylenimine, isoprene, lithium silicide, pentasilver trihydroxydiaminophosphate, phosphorus, phosphorus isocyanate, beta-propiolactone, and pyridine.

Hazardous decomposition products

Irritating and highly toxic fumes and gases, including oxides of sulfur. Reaction with water or steam may generate much heat which will increase the concentration of fumes in the air, and may produce toxic and corrosive fumes. Contact with most metals causes formation of flammable and explosive hydrogen gas.

SECTION 11: Toxicological information

Information on toxicological effects

Acute toxicity

Ingestion: Corrosive. Harmful if swallowed. Ingestion of liquid or spray mist may produce severe burns to the mouth, throat and stomach, resulting in sore throat, immediate severe burning pain in the mouth, throat, abdomen, general feeling of sickness, vomiting, diarrhoea, and the risk of perforation of oesophagus and stomach, leading to death. Ingestion can cause severe swelling of the larynx and skeletal paralysis affecting the ability to breathe, circulatory shock and convulsions. Circulatory collapse with clammy skin, weak and rapid pulse, shallow respirations, and scanty urine may follow ingestion. Circulatory shock is often the immediate cause of death. Ingestion can possibly cause pyloric stenosis after a latency period of some weeks.

Inhalation: Harmful if inhaled. Inhalation of the vapours/aerosols, spray mist or fumes may produce severe irritation and chemical burns to the nose, throat and respiratory tract, with burning sensation, sore throat, coughing, wheezing, choking, laboured breathing or shortness of breath, headache, nausea, and vomiting. Inhalation may result in dental erosion, laryngitis, bronchitis, spasm, inflammation and oedema of the larynx and bronchi, chemical pneumonitis, and delayed pulmonary oedema. Severe over-exposure can result in death.

Respiratory Irritation: Human volunteers exposed to sulfuric acid for 5-15 minutes noticed no odour, or irritation below 1 mg/m³. All volunteers noticed the exposure at 3 mg/m³ and at 5 mg/m³ some people found it objectionable. A deep breath usually produced coughing and respiratory changes were reported. Tolerance to sulfuric acid can occur.

In another study, volunteers exposed to high levels (39 mg/m³ dry mist and 21 mg/m³ wet mist sulfuric acid) for 1/2-1 hour reported severe symptoms of irritation of the upper airways and signs of bronchial obstruction. These symptoms persisted for several days in two volunteers. Occupational exposure to sulfuric acid fumes in a closed space, produced injury to the upper

airways, and fluid accumulation and bleeding in the lungs to one worker. Most lung function tests had returned to normal after 6 weeks.

Skin corrosion/irritation

Skin contact may produce severe skin burns. Symptoms of redness, irritation, pain, scaling, or, occasionally, blistering can occur. May be harmful if absorbed through the skin. Circulatory collapse with clammy skin, weak and rapid pulse, shallow respirations, and scanty urine may follow skin contact. Circulatory shock is often the immediate cause of death.

Serious eye damage/irritation

Liquid or spray mist may produce severe tissue burns. Eye contact can cause watering, blurred vision, redness, irritation and pain. It can cause permanent eye/corneal damage and blindness.

Respiratory or skin sensitization

No data available.

Germ cell mutagenicity

No data available

Carcinogenicity

Occupational exposure to strong-inorganic-acid mists containing sulfuric acid is evaluated in the IARC Monographs (Vol. 54; 1992) as Group 1: Carcinogenic to humans.

Reproductive toxicity

No data available

Specific target organ toxicity (STOT) - single exposure

No data available

Specific target organ toxicity (STOT) - repeated exposure No data available

Aspiration hazard

No data available

Additional information

Repeated or prolonged contact with liquid, vapour or spray mist may produce harmful corrosive effects to skin and respiratory system, chronic eye irritation and severe skin irritation. Prolonged or repeated inhalation may cause nosebleeds, nasal congestion, erosion of the teeth, perforation of the nasal septum, chest pain and respiratory tract irritation leading to frequent attacks of bronchial infection. Prolonged or repeated eye contact may cause conjunctivitis. Long-term exposure to mist or vapours may cause damage to teeth. May cause adverse reproductive effects. Chronic exposure to mists containing sulfuric acid is a cancer hazard.

SECTION 12: Ecological information

Toxicity

Damage of aquatic organisms. Harmful effect due to pH shift. Toxic effect for fishes and algeal. Caustic even in diluted form. Does not cause biological oxygen deficit. Endangers drinking-water supplies if allowed to enter soil and/or waters in large quantities. Neutralization possible in waste water treatment plants.

Persistence and degradability

No data available.

Bioaccumulative potential

No data available.

Mobility in soil No data available.

Results of PBT and vPvB assessment

No data available.

Endocrine disrupting properties No data available.

Other adverse effects

No data available.

SECTION 13: Disposal considerations

Disposal methods

Product disposal

Waste material must be disposed of in accordance with the national and local regulations. Leave chemicals in original containers.

Waste treatment

Neutralise remaining product with lime, soda ash or sodium bicarbonate, adjusting pH to 6-8. Flush to sewer as greatly diluted solution.

SECTION 14: Transport information

ADG (Road and Rail) UN Number: 2796 Class: 8 Packing Group: II Proper Shipping Name: SULFURIC ACID

Environmental Hazards: Damage of aquatic organisms. Harmful effect due to pH shift. Toxic effect for fishes and algeal. Caustic even in diluted form. Does not cause biological oxygen deficit. Endangers drinking-water supplies if allowed to enter soil and/or waters in large quantities. Neutralization possible in waste water treatment plants.

Hazchem emergency action code (EAC)

2R

IMDG

UN Number: 2796 Class: 8 Packing Group: II EMS Number: Proper Shipping Name: SULFURIC ACID

IATA

UN Number: 2796 Class: 8 Packing Group: II Proper Shipping Name: SULFURIC ACID

SECTION 15: Regulatory information

Safety, health and environmental regulations specific for the product in question

Australia SUSMP Poison Schedule: S6

SECTION 16: Other information

Further information/disclaimer

ChemSupply Australia Pty Ltd does not warrant that this product is suitable for any use or purpose. The user must ascertain the suitability of the product before use or application intended purpose. Preliminary testing of the product before use or application is recommended. Any reliance or purported reliance upon ChemSupply Australia Pty Ltd with respect to any skill or judgement or advice in relation to the suitability of this product of any purpose is disclaimed. Except to the extent prohibited at law, any condition implied by any statute as to the merchantable quality of this product or fitness for any purpose is hereby excluded. This product is not sold by description. Where the provisions of Part V, Division 2 of the Trade Practices Act apply, the liability of ChemSupply Australia Pty Ltd is limited to the replacement of supply of equivalent goods or payment of the cost of replacing the goods or acquiring equivalent goods.

Preparation information

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Standard for the Uniform Scheduling of Medicines and Poisons, Commonwealth of Australia National Road Transport Commission, 'Australian Code for the Transport of Dangerous Goods by Road and Rail 7th. Ed.' Safe Work Australia, 'National Code of Practice fot the Preparation of Safety Data Sheets for Hazardous Chemicals', July 2020. Safe Work Australia, 'National Guide for Classifying Hazardous Chemicals', July 2020. Safe Work Australia, Workplace Exposure Standards for Airbourne Contaminants, December 2019 Safe Work Australia, Hazardous Chemical Information System (HCIS), hcis.safeworkaustralia.gov.au IATA, Dangerous Goods Regulations (DGR) IMO, International Maritime Dangerous Goods Code (IMDG)