

# 3 2 Revision

2

# **Material Safety Data Sheets**

# 1. IDENTIFICATION

Product Name Sulphuric Acid 95-98% Extra pure

Other Names Dihydrogen Sulfate; Dipping Acid; Fertilizer Acid; Hydrogen Sulphate; Oil of Vitriol; SULFURIC ACID; Sulfuric Acid

Code No 100-SA-2

Uses Chemical Reagent, Electroplating, Fertilizer Manufacture, laboratory reagent, pickling and anodizing metals.

Chemical Family No Data Available

Chemical Formula H<sub>2</sub>SO<sub>4</sub>

Chemical Name Sulphuric Acid

Product Description No Data Available

Company Arman sina.co

Contact Information info@armansina.com

www.armansina.com

# 2. HAZARD IDENTIFICATION

Hazard Categories Corrosive

Risk Phrases Causes severe burns.

Risk of serious eye damage.

Safety Phrases Avoid contact with skin and eyes.

In case of contact with eyes, rinse immediately with plenty of water and seek

medical advice.

Never add water to this product.

Wear suitable protective clothing, gloves and eye/face protection.

In case of accident or if you feel unwell, seek medical advice immediately (show

Symbol



# 3. COMPOSITION/INFORMATION ON INGREDIENTS

### Ingredients

Chemical Entity	Formula	CAS Number	Proportion
Sulphuric Acid	H <sub>2</sub> SO <sub>4</sub>	7664-93-9	95-98%
Water	H <sub>2</sub> O	7732-18-5	BALANCE %

#### 4. FIRST AID MEASURES

#### Description of necessary measures according to routes of exposure

Swallowed If swallowed, do NOT induce vomiting. Seek medical attention immediately. For advice, contact a Poison Information

Centre on 13 11 26 (Australia Wide) or a doctor (at once).

Eye If in eyes, hold eyelids apart and flush the eye continuously with running water. Continue flushing until advised to stop

by the Poisons Information Centre or a doctor, or for at least 15 minutes.

Skin If skin or hair contact occurs, remove contaminated clothing and flush skin and hair with running water. Continue

flushing until advised to stop by the Poisons Information Centre or a doctor.

Inhaled If inhaled, remove from contaminated area. To protect rescuer, use a Full-face Type B (Inorganic and acid gas)

respirator or an Air-line respirator (in poorly ventilated areas). Apply artificial respiration if not breathing.

Advice to Doctor Treat symptomatically based on judgement of doctor and individual reactions of patient. Can cause corneal burns.

**Medical Conditions Aggravated** 

by Exposure

No information available on medical conditions aggravated by exposure to this product.

#### 5. FIRE FIGHTING MEASURES

General Measures If safe to do so, remove containers from the path of fire. Evacuate area and contact emergency services.

Flammability Conditions Product is a non-flammable liquid

Extinguishing Media For large fires, flood fire area with large quantities of water while knocking down vapours with water fog. If there is

insufficient water supply, knock down vapours only. For small fires, use Carbon dioxide, dry chemical, dry sand or flooding quantities of water. Toxic gases may be evolved in a fire situation. Remain upwind and notify those downwind of hazard. If tanks or containers are involved in the fire, cool them with copious quantities of water until well after the fire is out. Do not allow water to get inside tanks or containers. Withdraw immediately from the fire area if the tanks discolour or there is a rising sound from the safety vents. Stay away from tank ends. Use an extinguishing

agent suitable for the surrounding fire including water spray, foam, carbon dioxide or dry chemical powder.

Fire and Explosion Hazard

The product is non-combustible but will support combustion of other materials and may emit toxic fumes including those of sulphuric acid fumes and sulphur dioxide. The packaging material may burn to emit noxious fumes. Reacts

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with most common metals to liberate hydrogen which can form explosive mixtures with air.

Hazardous Products of

Combustion

May evolve toxic gases (sulphur oxides) when heated to decomposition. May evolve flammable hydrogen gas in

contact with some metals.

Special Fire Fighting Instructions Clear fire area of all non-emergency personnel. Stay upwind. Keep out of low areas. Eliminate ignition sources. Move

fire exposed containers from fire area if it can be done without risk. Do NOT allow fire fighting water to reach

waterways, drains or sewers. Store fire fighting water for treatment.

Personal Protective Equipment Fire fighters should wear a positive-pressure self-contained breathing apparatus (SCBA) and protective fire fighting

clothing (includes fire fighting helmet, coat, trousers, boots and gloves) or chemical splash suit. Please note:

Structural fire fighters uniform will provide limited protection.

Flash Point No Data Available
Lower Explosion Limit No Data Available
Upper Explosion Limit No Data Available

Auto Ignition Temperature No Data Available

Hazchem Code 2P

#### **6. ACCIDENTAL RELEASE MEASURES**

General Response Procedure Avoid accidents, clean up immediately. May be slippery when spilt. Eliminate all sources of ignition. Increase

ventilation. Isolate the danger area. Use clean, non-sparking tools and equipment. Shut off all possible sources if ignition.

ignitio

Clean Up Procedures

affected area with copious (flooding) quantities of water, so as to rapidly dilute the acid. Collect rinse water. Contact emergency services if appropriate for large spills, e.g. where acid has pooled. Do not enter area, walk though spilled material or touch damaged containers. Restrict access downwind and for at least 25 metres in other directions,

unless appropriate PPE is worn. Ventilate area where possible. If safe to do so, stop the leak. If the spill is uncontained, consider constructing an earthen bund to prevent loss to waterways and drains. Pump up spilt acid using acid resistant equipment into suitable dry containers. Vapour suppressing foam may be used to control vapours, water spray to knock down or divert vapour clouds. Do not direct streams of water at spilt acid or allow

water to enter containers. On impervious floors and in bunded areas, neutralize residual acid with Soda Ash. The resultant reaction product, sodium sulfate, is water soluble. The use of liming materials, e.g. calcium carbonate, oxide or hydroxide, is generally not recommended as the reaction product, calcium sulfate or gypsum, is sparingly soluble. The slurry is difficult to hose, pump and remove. Where Soda Ash is used, it may also be left to dry and be scraped up. Liming materials may be preferred to neutralize spills in the open to soil, as calcium sulfate has less impact on the soil and local environment than sodium sulfate. Depending on the magnitude of the spill, the neutralization of spills to soil may not be necessary if the soil is alkaline, e.g. calcareous soils in semi-arid areas. As an alternative to neutralization, residual acid can be soaked up with dry earth, sand, vermiculite or other inert absorbent material. Do not use sawdust. Where appropriate, cover with plastic sheeting to minimize spreading or contact with rain and collect for complete neutralisation and disposal.

Containment Stop leak if safe to do so.

Decontamination To avoid violent reactions, always add acid to water and never water to acid. When cleaning up residual acid after a

spill, use copious (flooding) quantities of water from the outset, to provide rapid dilution.

**Environmental Precautionary** 

Measures

Container

Do not allow product to reach drains, sewers or waterways. If product does enter a waterway, advise the Environmental Protection Authority or your local Waste Authority.

Evacuation Criteria Do not enter area, walk though spilled material or touch damaged containers. Restrict access downwind and for at

least 25 metres in other directions, unless appropriate PPE is worn. Ventilate area where possible.

#### 7. HANDLING AND STORAGE

Handling This acid is very reactive, and any contamination, including water, is potentially hazardous. The addition of water to

concentrated inorganic acids generates heat, causing the water to boil violently in the area of mixing, splattering the acid. Ensure an eye bath and safety shower are available and ready for use. Observe good personal hygiene practices and recommended procedures. Wash thoroughly after handling. Take precautionary measures against static discharges by bonding and grounding equipment. Avoid contact with eyes, skin and clothing. Do not inhale product vapours. Avoid prolonged or repeated exposure. Do not combine part drums of the same product, as this may be a source of contamination. Do not mix with other chemicals. Keep dry reacts with water; may lead to drum

rupture.

Storage Store in a cool, dry, well-ventilated area. Keep containers tightly closed when not in use. Inspect regularly for deficiencies such as damage or leaks. Protect against physical damage. Store away from incompatible materials as listed in section 10. Store in secured, cool, dry, well ventilated area, removed from oxidising agents, alkalis (bases), most metals, sunlight, heat or ignition sources and foodstuffs. Do not place in aluminium or galvanized containers. Ensure containers are adequately labelled and protected from physical damage when not in use. Check regularly for

leaks or spills. Large storage areas should have appropriate fire protection and ventilation systems. This product has a UN classification of 1830 and a Dangerous Goods Class 8 (Corrosive) according to The Australian Code for the Transport of Dangerous goods By Road and Rail.

Container type/packaging must comply with all applicable local legislation. Store in original packaging as approved by

manufacturer.

# 8. EXPOSURE CONTROLS / PERSONAL PROTECTION

General The following exposure standard has been established by The Safe Work Australia (SWA);

Sulphuric Acid CAS 7664-93-9: TWA = 1 mg/m3 STEL = 3 mg/m3

NOTE: The exposure value at the TWA is the average airborne concentration of a particular substance when

calculated over a normal 8 hour working day for a 5 day working week.

These exposure standards are guides to be used in the control of occupational health hazards. All atmospheric contamination should be kept to as low a level as is workable. These exposure standards should not be used as fine dividing lines between safe and dangerous concentrations of chemicals. They are not a measure of relative toxicity.

Exposure Limits No Data Available

Biological Limits No information available on biological limit values for this product.

Engineering Measures A system of local and/or general exhaust is recommended to keep employee exposures as low as possible. Local exhaust ventilation is generally preferred because it can control the emissions of the contaminant at its source.

preventing dispersion of it into the general work area. Adequate ventilation should be provided so that exposure limits are not exceeded. Use a corrosion-resistant ventilation system separate from other exhaust ventilation systems. Exhaust directly to the outside. Use local exhaust ventilation, and process enclosure if necessary, to control airborne

spray/ mists. Supply sufficient air to make up for air removed by exhaust systems.

Personal Protection Equipment

RESPIRATOR: Where risk assessment shows air-purifying respirators are appropriate use a Full Facepiece Gas

Mask/Chemical Cartridge Respirators as a backup to engineering controls. If the respirator is the sole means of

Mask/Chemical Cartridge Respirators as a backup to engineering controls. If the respirator is the sole means of protection, use a full-face supplied air respirator. Use respirators and components tested and approved under appropriate government standards (AS1715/1716).

EYES: Tightly fitting safety goggles. Faceshield (8-inch minimum). Use equipment for eye protection tested and approved under appropriate government standards (AS1336/1337).

HANDS: Handle with gloves. Gloves must be inspected prior to use. Use proper glove removal technique (without touching glove's outer surface) to avoid skin contact with this product (AS2161).

HANDS: Handle with gloves. Gloves must be inspected prior to use. Use proper glove removal technique (without touching glove's outer surface) to avoid skin contact with this product. (AS2161).

CLOTHING: Complete suit protecting against chemicals, The type of protective equipment must be selected according to the concentration and amount of the dangerous substance at the specific workplace. (AS3765/2210).

**Full contact** 

Material: Fluorinated rubber Minimum layer thickness: 0.7 mm Break through time: 480 min

890 / Aldrich Z677698, Size M)

Splash contact

Material: Nitrile rubber

Minimum layer thickness: 0.2 mm Break through time: 30 min

743 / Aldrich Z677388, Size M)

**Work Hygienic Practices** 

Use of safe work practices are recommended to avoid eye or skin contact and inhalation. Observe good personal hygiene, including washing hands before eating. Prohibit eating, drinking and smoking in contaminated areas.

#### 9. PHYSICAL AND CHEMICAL PROPERTIES

Physical State Liquid

Appearance Clear Liquid
Odour Slight odour

Colour Colorless to brown
pH No Data Available
Vapour Pressure No Data Available

Relative Vapour Density  $3.39 \, \text{Air} = 1$ Boiling Point  $310^{\circ}\text{C}$ Melting Point  $-15^{\circ}\text{C}$ 

Freezing Point No Data Available

Solubility Soluble Specific Gravity 1.84

Flash Point No Data Available **Auto Ignition Temp** No Data Available **Evaporation Rate** No Data Available **Bulk Density** No Data Available **Corrosion Rate** No Data Available **Decomposition Temperature** No Data Available Density 1.84 g/ml Relative Specific Heat No Data Available Molecular Weight No Data Available **Net Propellant Weight** No Data Available **Octanol Water Coefficient** No Data Available Particle Size No Data Available **Partition Coefficient** No Data Available **Saturated Vapour Concentration** No Data Available **Vapour Temperature** No Data Available Viscosity No Data Available **Volatile Percent** No Data Available **VOC Volume** No Data Available

**Additional Characteristics** 

Surface tension 55.1 mN/m at 20 Relative vapour density 3.39 - (Air = 1.0)

When mixing acids with water (diluting), caution must be taken as heat will be generated which causes violent

spattering. Always add a small volume of acid to a large volume of water, NEVER the reverse. RESPIRATORS: In general the use of respirators should be limited and engineering controls employed to avoid

exposure. If respiratory equipment must be worn ensure correct respirator selection and training is undertaken. Remember that some respirators may be extremely uncomfortable when used for long periods. The use of air powered or air supplied respirators should be considered where prolonged or repeated use is necessary. PERSONAL PROTECTIVE EQUIPMENT GUIDELINES:

The recommendation for protective equipment contained within this report is provided as a guide only. Factors such as method of application, working environment, quantity used, product concentration and the availability of engineering controls should be considered before final selection of personal protective equipment is made. **HEALTH EFFECTS FROM EXPOSURE:** 

It should be noted that the effects from exposure to this product will depend on several factors including: frequency and duration of use; quantity used; effectiveness of control measures; protective equipment used and method of application. Given that it is impractical to prepare a ChemAlert report which would encompass all possible scenarios, it is anticipated that users will assess the risks and apply control methods where appropriate.

Potential for Dust Explosion

Product is a liquid. Fast or Intensely Burning No Data Available

Characteristics

Flame Propagation or Burning Rate of Solid Materials

Non-Flammables That Could Contribute Unusual Hazards to a

Properties That May Initiate or Contribute to Fire Intensity

No Data Available

No Data Available

No Data Available

Reactions That Release Gases or Reacts with most metals generating flammable/explosive hydrogen gas.

**Vapours** 

Release of Invisible Flammable

Vapours and Gases

No Data Available

#### 10. STABILITY AND REACTIVITY

General Information Corrosive liquid. Corrosive to many metals with the liberation of extremely flammable hydrogen gas.

Chemical Stability Potential for exothermic hazard. Can react violently, releasing heat, when mixed with water and strong alkalis (bases).

May evolve flammable, and when confined, explosive hydrogen gas in contact with some metals.

Conditions to Avoid Can act as an oxidizer with some organic compounds. Has a strong affinity for water. Reacts violently with water

generating large quantities of heat. Contact with hypochlorites (swimming pool sodium or calcium hypochlorite) liberates toxic chlorine gas. contact with cyanide's releases toxic hydrogen cyanide gas. Contact with sulphides and

carbides releases toxic and corrosive gases. Reacts violently with strong bases, amines and other alkaline.

Bases, Halides, Organic materials, Carbides, fulminates, Nitrates, picrates, Cyanides, Chlorates, alkalihalides, Zinc salts, permanganates, e.g. potassium permanganate, Hydrogen peroxide, Azides, Perchlorates., Nitromethane, phosphorous, Reacts violently with:, cyclopentadiene, cyclopentanone oxime, nitroaryl amines, hexalithium disilicide,

phosphorous(III) oxide, Powdered metals.

Hazardous Decomposition

Materials to Avoid

**Products** 

May evolve toxic gases (sulphur oxides) when heated to decomposition. May evolve flammable hydrogen gas in contact with some metals. Corrosive to many metals with the liberation of extremely flammable hydrogen gas.

Polymerization is not expected to occur. Exothermic reaction with water may result in violent spattering. Corrosive to **Hazardous Polymerisation** 

most metals liberating flammable hydrogen gas.

## 11. TOXICOLOGICAL INFORMATION

Oral LD50 Rat: 2140 mg/Kg **General Information** 

Inhalation LC50/2hr Rat: 510 mg/m3 LC50 (inhalation) 18 mg/m (guinea pig) LD50 (ingestion) 2140 mg/kg (rat)

TCLo (inhalation) 3 mg/m 24 weeks (human) Unreported Route LDLo (man) 135 mg/kg Eye Irritation (Rabbit) 5 mg/30S rinse (Severe) Highly corrosive. This product has the potential to cause serious adverse health effects. Use safe work practices to avoid eye or skin contact and inhalation. Over exposure may result in severe skin, eye and respiratory burns with permanent lung and tissue damage. Upon dilution, the potential for adverse health effects may be reduced. Strong inorganic acid mists containing sulphuric acid is classified as carcinogenic to humans (IARC Group 1). Occupational exposures to strong inorganic acid mists containing sulfuric acid are specifically associated with laryngeal and lung cancer in humans.

For the component Sulphuric acid: Repeated overexposure may lead to chronic conjunctivitis, lung damage and dental erosion. The International Agency for Research on Cancer (IARC) have concluded that occupational exposure to strong inorganic acid mists containing sulphuric acid is carcinogenic to humans, causing cancer of the larynx and to a lesser extent, the lung. No direct link has been established with sulphuric acid, itself, and cancer in humans. Exposure to any mist or aerosol during the use of this product should be avoided and exposure should not exceed the exposure standard.

Eyelrritant Corrosive to eyes and may injure the cornea. Contamination of eyes can result in permanent injury. Symptoms

include stinging, tearing, redness and swelling of eyes, lacrimation, corneal burns and possible permanent damage.

Ingestion Highly corrosive. Ingestion may result in burns to the mouth and throat, nausea, vomiting, abdominal pain and

diarrhoea. Ingestion of large quantities may result in ulceration, unconsciousness, convulsions and death.

Inhalation Highly corrosive - severe irritant. Over exposure may result in mucous membrane irritation of the respiratory tract,

coughing, bronchitis, ulceration, bloody nose, lung tissue damage, chemical pneumonitis, pulmonary oedema and

death.

SkinIrritant Corrosive to skin - may cause skin burns. May not produce an immediate burning sensation upon contact, delaying

the awareness that contact has occurred. Symptoms may include redness, burning, and swelling of skin, burns, and

other skin damage.

Carcinogen Category No Data Available

#### 12. ECOLOGICAL INFORMATION

Ecotoxicity Fish, fresh water: Gambusia affinis LC50/96hr: 42 mg/L

Crustacean: Pandalus montagui (Pink shrimp) EC50/48hr: 42.5 mg/l

Mosquito fish LC50 42 mg/L/96hr Shore Crab LC50 70 - 80 mg/L/48hr Hooknose fish LC50 80-90 mg/L/48hr Cockle LC50 200 - 500 mg/l/48hr

Persistence/Degradability Rapidly Degradable: YES

Mobility No information available on mobility for this product. Sulphuric acid is miscible with water and its dilution will increase

the velocity of downward movement

in the soil where it may dissolve the soil material.

Environmental Fate Do NOT allow product to enter waterways, drains or sewers. The product is highly acidic. If large spills occurred a

water pH drop could be responsible for an environmental effect on aquatic organisms. Sulphuric acid is harmful to aquatic life in very low concentrations. May cause corrosion and deterioration of many common materials found in

the environment (eg steel, limestone).

Bioaccumulation Potential Bioccumulative: NO
Environmental Impact No Data Available

#### 13. DISPOSAL CONSIDERATIONS

General Information

Dispose of in accordance with all local regulations. All empty packaging should be disposed of in accordance with Local Regulations or recycled/reconditioned at an approved facility. Reuse or

recycle if possible. Contaminated acid and concentrated acidic wash down water can be neutralised to pH 6-8 by the SLOW addition of the waste to a saturated basic solution containing up to 20% sodium carbonate (Soda Ash) or 10% sodium bicarbonate. The order of mixing is not critical for dilute or weak wash down water, i.e. the dilute acid can be added to the basic solution, or vice versa. contaminated soil and other sorbent materials can be neutralized with a solution of sodium carbonate or sodium bicarbonate, a dry liming material such as Hydrated Lime (calcium hydroxide), or any other suitable neutralizing agent. Collect neutralized waste for disposal. Containers pose a hazard when empty and must be treated as still containing acid until thoroughly rinsed. Empty containers can be

decontaminated with a 5% aqueous solution of Soda Ash or sodium bicarbonate, and then rinsing again with water. Waste disposal should only be undertaken in a well ventilated area. Contact Waste Management Authority for advice on whether waste needs to be treated as hazardous, or requires further dilution, neutralization or treatment, and

acceptable disposal methods.

Special Precautions for Land Fill

Contact a specialist disposal company or the local waste regulator for advice. The product is suitable for disposal by landfill through an approved agent. Incineration of the product is not recommended, as it is unlikely to adequately

burn

#### 14. TRANSPORT INFORMATION

# **Land Transport**

Proper Shipping Name SULPHURIC ACID with more than 51% acid

Class 8 Corrosive Substances
Subsidiary Risk(s) No Data Available

ERG 137 Substances - Water-Reactive - Corrosive

 UN Number
 1830

 Hazchem
 2P

 Pack Group
 II

Special Provision No Data Available

# **Sea Transport**

**IMDG** 

Proper Shipping Name SULPHURIC ACID with more than 51% acid

Class 8 Corrosive Substances
Subsidiary Risk(s) No Data Available

 UN Number
 1830

 Hazchem
 2P

 Pack Group
 II

Special Provision No Data Available

EMS FA,SB Marine Pollutant No

# Air Transport

IATA

Proper Shipping Name SULPHURIC ACID with more than 51% acid

Class 8 Corrosive Substances
Subsidiary Risk(s) No Data Available

 UN Number
 1830

 Hazchem
 2P

 Pack Group
 II

Special Provision No Data Available

#### 15. OTHER INFORMATION

Revision

Key/Legend < Less Than > Greater Than

atm Atmosphere

CAS Chemical Abstracts Service (Registry Number)

cm Square Centimetres CO2 Carbon Dioxide

**COD Chemical Oxygen Demand** 

Degrees Celcius Degrees Farenheit g Grams

g/cm Grams per Cubic Centimetre

g/I Grams per Litre

**HSNO Hazardous Substance and New Organism** 

IDLH Immediately Dangerous to Life and Health

immiscible Liquids are insoluable in each other.

inHg Inch of Mercury

inH2O Inch of Water

K Kelvin

kg Kilogram

kg/m Kilograms per Cubic Metre

**Ib Pound** 

LC50 LC stands for lethal concentration. LC50 is the concentration of a material in air which causes the death of 50% (one half) of a group of test animals. The material is inhaled over a set period of time, usually 1 or 4 hours. LD50 LD stands for Lethal Dose. LD50 is the amount of a material, given all at once, which causes the death of 50% (one half) of a group of test animals.

Itr or L Litre

m Cubic Metre

mbar Millibar

mg Milligram

mg/24H Milligrams per 24 Hours

mg/kg Milligrams per Kilogram

mg/m Milligrams per Cubic Metre

Misc or Miscible Liquids form one homogeneous liquid phase regardless of the amount of either component present.

mm Millimetre

mmH2O Millimetres of Water

mPa.s Millipascals per Second

N/A Not Applicable

NIOSH National Institute for Occupational Safety and Health

NOHSC National Occupational Heath and Safety Commission

**OECD Organisation for Economic Co-operation and Development** 

Oz Ounce

PEL Permissible Exposure Limit

Pa Pascal

ppb Parts per Billion

ppm Parts per Million

ppm/2h Parts per Million per 2 Hours

ppm/6h Parts per Million per 6 Hours

psi Pounds per Square Inch

R Rankine

**RCP Reciprocal Calculation Procedure** 

STEL Short Term Exposure Limit

**TLV Threshold Limit Value** 

tne Tonne

TWA Time Weighted Average

ug/24H Micrograms per 24 Hours

**UN United Nations** 

wt Weight