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RE-ISSUED by CHEMSUPP Infosafe No™ 3CH5V Issue Date: January 2019

PERCHLORIC ACID SOLUTION IN ACETIC ACID, 0.1M/0.1N Product Name:

Classified as hazardous

1. Identification

**GHS Product** 

PERCHLORIC ACID SOLUTION IN ACETIC ACID, 0.1M/0.1N

Identifier

CHEM-SUPPLY PTY LTD (ABN 19 008 264 211) **Company Name** 

38 - 50 Bedford Street GILLMAN **Address** 

> SA 5013 Australia Tel: (08) 8440-2000

Telephone/Fax Number

Fax: (08) 8440-2001 CHEMCALL 1800 127 406 (Australia) / +64-4-917-9888 (International)

**Emergency phone** 

number

Laboratory reagent, analytical chemistry, titrant in volumentric analysis.

Recommended use of the chemical and restrictions on use Other Names

**Product Code** Name

Perchloric acid in acetic acid, 0.1M/0.1N LR

Other Information

Chem-Supply 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 Chem-Supply 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 Chem-Supply 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.

PL360

#### 2. Hazard Identification

Flammable Liquids: Category 3 **GHS** classification

of the

Skin Corrosion/Irritation: Category 1A

H226 Flammable liquid and vapour.

substance/mixture

**DANGER** Signal Word (s)

**Hazard Statement** (s)

H314 Causes severe skin burns and eye damage.

Pictogram (s) Corrosion, Flame





**Precautionary** statement -

P210 Keep away from heat/sparks/open flames/hot surfaces. - No smoking.

P233 Keep container tightly closed.

Prevention

P241 Use explosion-proof electrical/ventilating/lighting/.../equipment.

P242 Use only non-sparking tools.

P243 Take precautionary measures against static discharge. P260 Do not breathe dust/fume/gas/mist/vapours/spray.

P280 Wear protective gloves/protective clothing/eye protection/face protection. P301+P330+P331 IF SWALLOWED: rinse mouth. Do NOT induce vomiting.

**Precautionary** statement -Response

P303+P361+P353 IF ON SKIN (or hair): Remove/Take off immediately all contaminated clothing. Rinse

skin with water/shower.

P363 Wash contaminated clothing before reuse.

P304+P340 IF INHALED: Remove victim to fresh air and keep at rest in a position comfortable for

breathing.

P310 Immediately call a POISON CENTER or doctor/physician.

P305+P351+P338 IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses.

if present and easy to do. Continue rinsing.

P370+P378 In case of fire: Use dry chemical, CO2, or water spray for extinction.

**Precautionary** 

P403+P235 Store in a well-ventilated place. Keep cool.

statement - Storage P405 Store locked up.



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**Precautionary** 

P501 Dispose of contents/container to an approved waste disposal plnt.

statement -Disposal

3. Composition/information on ingredients

Liquid

Characterization

Ingredients Name CAS **Proportion Hazard Symbol Risk Phrase** 

> Acetic acid 64-19-7 99 % Perchloric acid 7601-90-3 1 %

4. First-aid measures

Inhalation If inhaled, remove from contaminated area to fresh air immediately. If breathing is difficult, give oxygen.

Apply artificial respiration with a respiratory medical device if not breathing. Do not use mouth to mouth

resuscitation. Immediately medical attention is required.

Rinse mouth thoroughly with water immediately, repeat until all traces of product have been removed. Ingestion

DO NOT INDUCE VOMITING. Seek immediate medical advice.

Immediately remove contaminated clothing and wash affected area with water for at least 15 minutes. Skin

Ensure contaminated clothing is washed before re-use. Seek medical advice /attention depending on the

If contact with the eye(s) occurs, wash with copious amounts of water for approximately 15 minutes Eye contact

holding eyelid(s) open. Take care not to rinse contaminated water into the non-affected eye. Seek

medical attention.

Maintain eyewash fountain and safety shower in work area. **First Aid Facilities** 

**Advice to Doctor** Treat symptomatically based on judgement of doctor and individual reactions of the patient.

**Other Information** For advice, contact the National Poisons Information Centre (Phone Australia 13 11 26; New Zealand

0800 764 766) or a doctor.

5. Fire-fighting measures

Hazards from Combustion **Products** 

Vapours may form explosive mixtures with air creating an explosion hazard. May evolve irritating, poisonous/corrosive, toxic fumes in fire such as oxides of carbon and hydrogen chloride gas.

**Specific Methods** Small fire: Use foam, dry chemical, CO2, or water spray.

Large fire: Use foam, fog or water spray - Do not use water jets.

If safe to do so, move undamaged containers from fire area. Cool containers with flooding quantities of

water until well after fire is out. Avoid getting water inside containers.

Specific hazards arising from the chemical

May be ignited by heat, sparks or flame. Vapours may form explosive mixtures with air. Vapours may travel to source of ignition and flash back. Most vapours are heavier than air and will collect in low or confined areas (drains, basements, tanks). Many liquids are lighter than water. Containers may explode when heated. Vapours from runoff may create an explosion hazard. Fire will produce irritating, poisonous

and/or corrosive gases. May decompose exposively when heated or involed in a fire.

**Hazchem Code** 

Precautions in

Wear SCBA and fully-encapsulating, gas-tight suit when handling these substances. Structural

connection with Fire firefighter's uniform is NOT effective for these materials.

6. Accidental release measures

Spills & Disposal ELIMINATE all ignition sources (no smoking, flares, sparks or flame) within at least 25m. All equipment

used when handling the product must be earthed. Do NOT touch or walk through spilled material. Stop leak if safe to do so. Prevent entry into waterways, drains or confined areas. Vapour-suppressing foam may be used to control vapours. Water spray may be used to knock down or divert vapour clouds. Absorb with earth, sand or other non-combustible material. Use clean, non-sparking tools to collect absorbed material and place it into loosely-covered metal or plastic containers for later disposal.

SEEK EXPERT ADVICE ON HANDLING AND DISPOSAL.

Avoid contact with skin, eyes and clothing. Avoid inhalation and ingestion of product. **Personal** 

**Precautions** 

Personal Protection Wear protective clothing specified for normal operations (see Section 8)

**Small Spillages** 

Clean-up Methods - Absorb or contain liquid with sand, earth or spill control 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.



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**Environmental Precautions** 

Prevent from entering into drains, ditches, rivers or the sea.

7. Handling and storage

Handling

Precautions for Safe Extinguish any naked flames. Remove ignition sources. Avoid sparks. Do not smoke. Take precautionary measures against static discharges. Earth all equipment Ensure the appropriate personal protective equipment is used when handling this material. All electrical equipment must be flameproofed. Use in well ventilated areas away from all ignition sources. In case of insufficient

ventilation, wear suitable respiratory equipment.

Conditions for safe storage, including any

Do not smoke in storage areas. Keep container tightly closed and in a cool, well-ventilated place Keep away from heat and other sources of ignition.

incompatabilities

Corrosiveness Corrosive in presence of stainless steel. Slightly corrosive in presence of aluminium and copper.

Storage Regulations Refer Australian Standard AS 3780-1994 'The storage and handling of corrosive substances'. Refer Australian Standard AS 1940-2017 'The storage and handling of flammable and combustible liquids'.

Store at room temperature (15 to 25 °C recommended). Storage

**Temperatures** 

8. Exposure controls/personal protection

STEL Occupational **Name TWA** 

exposure limit values

> mg/m3 mg/m3 ppm <u>ppm</u> **Footnote**

Acetic acid 37 15 25 10

Other Exposure Information

A time weighted average (TWA) has been established for Acetic acid (Safe Work Australia) of 25 mg/m³, (10 ppm). The corresponding STEL level is 37 mg/m³, (15 ppm). The STEL (Short Term Exposure Limit) is an exposure value that should not be exceeded for more than 15 minutes and should not be repeated for more than 4 times per day. There should be at least 60 minutes between successive exposures at the STEL. 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.

**Appropriate** 

In industrial situations maintain the concentrations values below the TWA. This may be achieved by engineering controls process modification, use of local exhaust ventilation, capturing substances at the source, or other methods. These methods should be used in preference to personal protective equipment.

Respiratory **Protection** 

Where ventilation is not adequate, respiratory protection may be required. Avoid breathing dust, vapours or mists. Respiratory protection should comply with AS 1716 - Respiratory Protective Devices and be selected in accordance with AS 1715 - Selection, Use and Maintenance of Respiratory Protective Devices. Filter capacity and respirator type depends on exposure levels. In event of emergency or planned entry into unknown concentrations a positive pressure, full-facepiece SCBA should be used. If respiratory protection is required, institute a complete respiratory protection program including selection,

fit testing, training, maintenance and inspection.

The use of a face shield, chemical goggles or safety glasses with side shield protection as appropriate. Eye Protection

Must comply with Australian Standards AS 1337 and be selected and used in accordance with AS 1336. Wear gloves of impervious material conforming to AS/NZS 2161: Occupational protective gloves -

Selection, use and maintenance. Final choice of appropriate glove type will vary according to individual circumstances. This can include methods of handling, and engineering controls as determined by

appropriate risk assessments.

Recommendation: Excellent: polyvinyl chloride gloves. Poor: Nitrile, Neoprene, NR latex.

**Personal Protective Equipment** 

**Hand Protection** 

Personal protective equipment should not solely be relied upon to control risk and should only be used when all other reasonably practicable control measures do not eliminate or sufficiently minimise risk. Guidance in selecting personal protective equipment can be obtained from Australian, Australian/New

Zealand or other approved standards.

Safety boots in industrial situations is advisory, foot protection should comply with AS 2210, **Footwear** 

Occupational protective footwear - Guide to selection, care and use.

**Body Protection** Clean clothing or protective clothing should be worn, preferably with an apron. Clothing for protection

against chemicals should comply with AS 3765 Clothing for Protection Against Hazardous Chemicals. Always wash hands before smoking, eating or using the toilet. Wash contaminated clothing and other

**Hygiene Measures** protective equipment before storing or re-using.

#### 9. Physical and chemical properties



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Liquid **Form** 

**Appearance** Clear, colourless liquid. Of acetic acid; pungent. Odour 120°C at 1013 hPa **Boiling Point** 

Solubility in Water Soluble.

Solubility in Organic Soluble in diethyl ether, acetone, glycerol, alcohol, Benzene and carbon tetrachloride. Practically

insoluble in carbon disulfide. **Solvents** 

1.06 g/cm3 at 20°C **Specific Gravity** 

pH <1; pH 2.4 (1.0M solution). Hq

Partition Coefficient: log Kow = -0.17

n-octanol/water

**Flash Point** 40°C (closed cup). **Flammability** Flammable liquid.

**Auto-Ignition** 

454°C

**Temperature** 

Flammable Limits -3.3% (V)

Lower

Flammable Limits -

22% (V)

Upper

Other Information Taste: Burning taste.

Heat of combustion: 874.2 kJ/mol

#### 10. Stability and reactivity

**Chemical Stability** Stable under normal use conditons.

Releases heat and toxic, irritating vapors when mixed with water.

**Conditions to Avoid** Heat, sources of ignition. Incompatibles.

Incompatible **Materials** 

Oxidizing agents, chromates, soluble carbonates and phosphates, hydroxides, metals, hydrogen peroxide, peroxi compounds, permanganates (e.g. potassium permanganate), amines, alkalis, and

alcohols.

**Hazardous Decomposition Products** 

Vapours may form explosive mixtures with air creating an explosion hazard. May evolve irritating, poisonous/corrosive, toxic fumes in fire such as oxides of carbon and hydrogen chloride gas.

**Hazardous** 

Will not occur.

**Polymerization** 

#### 11. Toxicological Information

Acute Toxicity - Oral LD50 (rat): 3310 mg/kg (acetic acid) LD50 (rabbit): 1113 mg/k (acetic acid) **Acute Toxicity -**

**Dermal** 

Corrosive. Poisonous, may be fatal if swallowed. Causes severe burns and irritation to gastrointestinal Ingestion

> tract (mouth, oesophagus, stomach, abdomen). Symptoms include sore throat, coughing, ulceration, nausea, gastric spasms, bleeding, bloody vomiting, dyspnoea (breathing difficulty), asphyxia (suffocation), hematemesis and diarrhea which may lead to shock, pulmonary failure possible after aspiration of vomit, cardiovascular failture, acidosis, coma and death. Risk of perforation in the oesophagus and stomach! May also affect and damage the liver (impaired liver function), behaviour (convulsions, giddiness, muscular weakness), and the kidneys (Hematuria, Albuminuria, Nephrosis,

acute renal failure, acute tubular necrosis).

Inhalation Poisonous, may be fatal if inhaled. Material is extremely descructive to the tissue of the mucous

membranes and upper respiratory tract (nose, throat, lungs). Symptoms may include of burning sensation, coughing, wheezing, sneezing, chest pain, salivation, nausea, muscular weakness, shortness of breath/breathing difficulties and irritation. Inhalation may lead to inflammation and edema of the larynx and bronchi, spasms, pneumonitis and pulmonary edema. Neither odour nor degree of irritation

are adequate to indicate vapor concentration.

Poisonous, may be fatal if absorbed through the skin. Causes severe skin irritation and burns. Material is Skin

extremely desctructive to the tissue of the mucous membranes and the skin. Skin inflammation is

characterized by itching, scaling, reddening and blistering.



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Causes severe irritation and burns to eyes. Material is extremely destructive to the tissue of the mucous Eye

membranes and eyes. Inflammation of the eye is characterised by redness, watering, and itching. Risk

of blindness! Risk of corneal clouding!

Carcinogenicity Not listed in the IARC Monographs.

**Chronic Effects** Prolonged or repeated exposure may result in lung tissue damage (such as bronchial infection,

pulmonary edema) and corrosion (ulceration) of the mucous membranes. Swallowing can cause severe injury leading to cardiovascular failure, acidosis, liver and kidney damage, perforation of the oesophagus and stomach, and death. Prolonged or repeated exposure can result in blindness and or corneal clouding to the eye. Repeated or prolonged contact with spray mist may produce chronic eye irritation and severe skin irritation. Repeated or prolonged exposures may cause darkening of the skin, erosion of

exposed front teeth, and chronic inflammation of the nose and throat.

Mutagenicity No evidence of mutagenic properties.

12. Ecological information

**Ecotoxicity** Harmful effect due to pH shift. Persistence and BOD: 0.88 g/g (chief component).

degradability Mobility

log P(o/w): -0.31 (chief component).

**Bioaccumulative** 

**Potential** 

Low probability of bioaccumulation is expected (log P(o/w) < 1).

**Environmental** 

Do not allow product to enter drains, waterways or sewers.

**Protection** 

Acute Toxicity - Fish LC50 (L.macrochirus): 75 mg/l /96h (chief component). **Acute Toxicity -**LC50 (Daphnia magna): 47 mg/l /24h (chief component).

Daphnia

13. Disposal considerations

Disposal Whatever cannot be saved for recovery or recycling should be disposed of according to relevant local. state and federal government regulations. Considerations

14. Transport information

Dangerous Goods of Class 8 Corrosives are incompatible in a placard load with any of the following: -Transport Information

Class 1, Class 4.3, Class 5, Class 6, if the Class 6 dangerous goods are cyanides and the Class 8

dangerous goods are acids and Class 7.

**U.N. Number** 

UN proper shipping ACETIC ACID SOLUTION

name

**Transport hazard** 

class(es)

Sub.Risk

3

**Hazchem Code** •2P **Packing Group** Ш **EPG Number** 8B1 **IERG Number** 19

15. Regulatory information

Regulatory Information Not listed under WHS Regulation 2011, Schedule 10 - Prohibited carcinogens, restricted carcinogens

and restricted hazardous chemicals.

**Poisons Schedule** S<sub>6</sub>

16. Other Information

Literature Standard for the Uniform Scheduling of Medicines and Poisons .', Commonwealth of Australia. References

Lewis, Richard J. Sr. 'Hawley's Condensed Chemical Dictionary 13th. Ed.', Rev., John Wiley and Sons,

Inc., NY, 1997.

National Road Transport Commission, 'Australian Code for the Transport of Dangerous Goods by Road

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Standards Australia, 'SAA/SNZ HB 76:2010 Dangerous Goods - Initial Emergency Response Guide',

Standards Australia/Standards New Zealand, 2010.

Safe Work Australia, 'Approved Criteria for Classifying Hazardous Substances [NOHSC:1008 (2004)]'.

Safe Work Australia, 'Hazardous Chemical Information System, 2005'.

Safe Work Australia, 'National Code of Practice for the Labelling of Safe Work Hazardous Substances

(2011)

Safe Work Australia, 'National Exposure Standards for Atmospheric Contaminants in the Occupational

Environment [NOHSC:1003(1995) 3rd Edition]'.

Contact Person/Point Paul McCarthy Ph. (08) 8440 2000 DISCLAIMER STATEMENT:

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