

Callington Haven Pty Ltd

Chemwatch; 4512-19 Version No: 9.1.1.1

Safety Data Sheet according to WHS and ADG requirements

Chemwatch Hazard Alert Code: 4

tssue Date: 07/03/2020 Print Date: 20/07/2020 S.GHS.AUS.EN

SECTION 1 IDENTIFICATION OF THE SUBSTANCE / MIXTURE AND OF THE COMPANY / UNDERTAKING Product Identifier Product name Weldbrite weld scale remover stainless steel pickling paste Synonyms CORROSIVE LIQUID, TOXIC, N.O.S. (contains nitric acid and hydrofluoric acid) Proper shipping name Other means of identification Not Available Relevant identified uses of the substance or mixture and uses advised against Relevant identified uses Thickened stainless steel pickling paste. Details of the supplier of the safety data sheet Registered company name Callington Haven Pty Ltd Address 30 South Street Rydalmere NSW 2116 Australia +61 2 9898 2700 Telephone Fax +61 2 9475 0449 www.callingtonhaven.com Website Email customerservice@callington.com Emergency telephone number Association / Organisation CHEMWATCH EMERGENCY RESPONSE Chemwatch Emergency telephone +61 1800 951 288 +61 1800 951 288 numbers Other emergency telephone +61 2 9186 1132 +61 2 9186 1132 numbers Once connected and if the message is not in your prefered language then please dial 01 **SECTION 2 HAZARDS IDENTIFICATION** Classification of the substance or mixture Poisons Schedule Acute Toxicity (Oral) Category 3, Acute Toxicity (Dermal) Category 3, Acute Toxicity (Inhalation) Category 3, Skin Corrosion/Irritation Category Classification [1] 1A, Serious Eye Damage Category 1, Skin Sensitizer Category 1, Germ cell mutagenicity Category 2 1. Classified by Chemwatch: 2. Classification drawn from HGIS: 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI Legend: Label elements Hazard pictogram(s) SIGNAL WORD DANGER Hazard statement(s) H301 Toxic if swallowed. H311 Toxic in contact with skin. H331 Toxic if inhaled. H314 Causes severe skin burns and eye damage. H317 May cause an altergic skin reaction. H341 Suspected of causing genetic defects. Precautionary statement(s) Prevention

Obtain special instructions before use.

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P260	Do not breathe mist/vapours/spray.
P270	Do not eat, drink or smoke when using this product.
P271	Use only outdoors or in a well-ventilated area.
P280	Wear protective gloves/protective clothing/eye protection/face protection.
P281	Use personal protective equipment as required.
P272	Contaminated work clothing should not be allowed out of the workplace.

Precautionary statement(s) Response

P301+P310	IF SWALLOWED: Immediately call a POISON CENTER or doctor/physician.
P301+P330+P331	IF SWALLOWED: Rinse mouth. Do NOT induce vomiting.
P303+P361+P353	IF ON SKIN (or hair): Remove/Take off immediately all contaminated clothing. Rinse skin with water/shower.
P305+P351+P338	IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.
P308+P313	IF exposed or concerned: Get medical advice/attention.
P321	Specific treatment (see advice on this label).
P322	Specific measures (see advice on this label).
P363	Wash contaminated clothing before reuse.
P302+P352	IF ON SKIN: Wash with plenty of water.
P304+P340	IF INHALED: Remove victim to fresh air and keep at rest in a position comfortable for breathing.
P333+P313	If skin irritation or rash occurs: Get medical advice/attention.

Precautionary statement(s) Storage

P403+P233	Store in a well-ventilated place. Keep container tightly closed.
P405	Store locked up.
	er karakan kan beranggalan dan beranggalan beranggalan beranggalan kepadahan beranggalan beranggalan beranggal

Precautionary statement(s) Disposal

P501 Dispose of contents/container to authorised hazardous or special waste collection point in accordance with any local regulation.

SECTION 3 COMPOSITION / INFORMATION ON INGREDIENTS

Substances

See section below for composition of Mixtures

Mixtures

CAS No	%[weight]	Name
7697-37-2	10-30	nitric acid
7664-39-3	<10	hvdrofluoric acid
Not Available	10-30	performance additives
7732-18-5	>60	waler
Not Available	:	NOTE: Manufacturer has supplied full ingredient
Not Available		information to allow CHEMWATCH assessment.

SECTION 4 FIRST AID MEASURES

Description of first aid measur	es
Eye Contact	DO NOT delay. If this product or its vapours come in contact with the eyes, DO NOT DELAY: Immediately irrigate continuously by holding the cyclids apart and washing with fresh running water. Ensure complete irrigation of the eye by keeping cyclids apart and away from eye and moving the cyclids 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, eye clinic or eye specialist, ophthalmologist without delay.
Skin Contact	DO NOT delay. If there is evidence of severe skin irritation or skin burns: Avoid further contact. Immediately remove contaminated clothing, including footwear. Flush skin under running water for 15 minutes. Avoiding contamination of the hands, massage calcium gluconate gel into affected areas, pay particular attention to creases in skin. Contact the Poisons Information Centre. Continue gel application for at least 15 minutes after burning sensation ceases. If pain recurs, repeat application of calcium gluconate gel or apply every 20 minutes. If no gel is available, continue washing for at least 15 minutes, using soap if available. If patient is conscious, give six calcium gluconate or calcium carbonate tablets in water by mouth. Transport to hospital, or doctor, urgently.
	For massive exposures:

If dusts, vapours, aerosols, fumes or combustion products are inhaled, remove from contaminated area.
 Lay patient down.

Inhalation

- 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. If victim is conscious, give six calcium gluconate or calcium carbonate tablets in water by mouth.

Transport to hospital, or doctor, urgently.

DO NOT delay

Rinse mouth out with plenty of water.

Indestion

Transport to hospital or doctor and seek immediate medical attention,

DO NOT INDUCE vomiting.

If patient is conscious, give six calcium gluconate or calcium carbonate tablets dissolved in water, by mouth,

Indication of any immediate medical attention and special treatment needed

Following acute or short term repeated exposure to hydrofluoric acid:

- Subculaneous injections of Calcium Gluconate may be necessary around the burnt area. Continued application of Calcium Gluconate Get or subcutaneous Calcium Gluconate should then continue for 3-4 days at a frequency of 4-6 times per day. If a "burning" sensation recurs, apply more frequently.
- Systemic effects of extensive hydrofluoric acid burns include renal damage, hypocalcaemia and consequent cardiac arrhythmias. Monitor haematological, respiratory, renal, cardiac and electrolyte status at least daily. Tests should include FBE, blood gases, chest X-ray, creatinine and electrolytes, urine output, Ca ions, Mg ions and phosphate ions. Continuous ECG monitoring may be required.
- Where serum calcium is low, or clinical, or ECG signs of hypocalcaemia develop, infusions of calcium gluconate, or if less serious, oral Sandocal, should be given. Hydrocortisone 500 mg in a four to six hourly infusion may help.
- Antibiotics should not be given as a routine, but only when indicated.
- Eye contact pain may be excruciating and 2-3 drops of 0.05% pentocaine hydrochloride may be instilled, followed by further irrigation

BIOLOGICAL EXPOSURE INDEX - BEI

These represent the determinants observed in specimens collected from a healthy worker exposed at the Exposure Standard (ES or TLV);

Index

Sampling Time

Comments

1. Methaemoglobin in blood

1.5% of haemoglobin

During or end of shift

B, NS, SQ

B: Background levels occur in specimens collected from subjects NOT exposed.

NS: Non-specific determinant: Also seen after exposure to other materials

SQ: Semi-quantitative determinant - Interpretation may be ambiguous; should be used as a screening test or confirmatory test.

SECTION 5 FIREFIGHTING MEASURES

Extinguishing media

- Foam.
- Dry chemical powder.
- BCF (where regulations permit).
- Carbon dioxide
- Water spray or fog Large fires only.

Special hazards arising from the substrate or mixture

· Avoid any contamination of this material as it is very reactive and any contamination is potentially hazardous Avoid storage with glass, cement, concrete and other silicon materials; reaction produces toxic silicon tetrafluoride gas; which may pressurise and/or rupture containers. Avoid reaction with

organic materials / compounds

Fire Incompatibility powdered metals

reducing agents and

hydrogen sulfide (H2S)

as ignition may result

Reacts with mild steel, galvanised steel / zinc producing hydrogen gas which may form an explosive mixture with air.

Advice for firefighters

- · Alert Fire Brigade and tell them location and nature of hazard.
- Wear full body protective clothing with breathing apparatus. Consider evacuation (or protect in place).
- - Use water delivered as a fine spray to control fire and cool adjacent area.
 - DO NOT approach containers suspected to be hot.
 - Cool fire exposed containers with water spray from a protected location. If safe to do so, remove containers from path of fire.

 - Equipment should be thoroughly decontaminated after use.
 - Non combustible.
 - Not considered to be a significant fire risk.
 - Acids may react with metals to produce hydrogen, a highly flammable and explosive gas.
 - Heating may cause expansion or decomposition leading to violent rupture of containers.
 - Decomposes on heating and may produce toxic fumes of carbon monoxide (CO).

 May emit acrid smoke. May emit corrosive fumes. Other decomposition products include:

carbon dioxide (CO2) hydrogen fluoride

nitrogen oxides (NOx)

HAZCHEM

Fire/Explosion Hazard

Fire Fighting

SECTION 6 ACCIDENTAL RELEASE MEASURES

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Personal precautions, protective equipment and emergency procedures

See section 8

Environmental precautions

See section 12

Methods and material for containment and cleaning up

Minor Spills

Slippery when spilt. ▶ DO NOT touch the spill material

Clean up all spills immediately.

Avoid breathing vapours and contact with skin and eyes.

Wear impervious gloves and safety glasses. Use soda ash or slaked lime to neutralise.

Trowel up/scrape up.

Place spilled material in clean, dry, sealable, labelled container.

Flush spill area with water.

Slippery when spilt.

- Clear area of personnel and move upwind.
- Alert Fire Brigade and tell them location and nature of hazard.
- Wear full body protective clothing with breathing apparatus.
- Prevent, by any means available, spillage from entering drains or water course.
- Consider evacuation (or protect in place).
- Major Spills
- Stop leak if safe to do so. Contain spill with sand, earth or vermiculite.
- Collect recoverable product into labelled containers for recycling.
- Neutralise/decontaminate residue (see Section 13 for specific agent).
- Collect solid residues and seal in labelled drums for disposal.
- Wash area and prevent runoff into drains.
- After clean up operations, decontaminate and launder all protective clothing and equipment before storing and re-using.
- If contamination of drains or waterways occurs, advise emergency services.

Personal Protective Equipment advice is contained in Section 8 of the SDS.

SECTION 7 HANDLING AND STORAGE

Precautions for safe handling

DO NOT allow clothing wet with material to stay in contact with skin

Use good occupational work practice. Observe manufacturer's storage and handling recommendations contained within this SDS.

Atmosphere should be regularly checked against established exposure standards to ensure safe working conditions are maintained.

Avoid all personal contact, including inhalation

Wear protective clothing when risk of exposure occurs. Avoid contact with incompatible materials.

Safe handling

WARNING: To avoid violent reaction, ALWAYS add material to water and NEVER water to material.

Handle and open container with care
 When handling, DO NOT eat, drink or smoke.

Keep containers securely sealed when not in use

Avoid physical damage to containers.

Wash hands with soap and water after handling.

Work clothes should be laundered separately: NOT at home.

- Store in original containers.
 - Keep containers securely sealed.
 - Store in a cool, dry, well-ventilated area. Store away from incompatible materials and foodstuff containers.

 - Protect containers against physical damage and check regularly for leaks.
 - Observe manufacturer's storage and handling recommendations contained within this SDS.

Conditions for safe storage, including any incompatibilities

Suitable container

Other information

- Polyethylene or polypropylene container.
- Packing as recommended by manufacturer.
- Check all containers are clearly labelled and free from leaks.

Avoid storage with glass, cement, concrete and other silicon materials; reaction produces toxic silicon tetrafluoride gas; which may pressurise and/or rupture containers.

- DO NOT use unlined steel containers
- DO NOT use aluminium, galvanised or tin-plated containers

Storage incompatibility

Segregate from alkalies, oxidising agents and chemicals readily decomposed by acids, i.e. cyanides, sulfides, carbonates,

combustible materials

metal oxides

SECTION 8 EXPOSURE CONTROLS / PERSONAL PROTECTION

Control parameters

OCCUPATIONAL EXPOSURE LIMITS (OEL)

INGREDIENT DATA

Source Ingredient Material name TWA STEL Peak Notes Australia Exposure Standards nitric acid Nitric acid 2 ppm / 5.2 mg/m3 10 mg/m3 / 4 ppm Not Available Not Available

Australia Exposure Standards hydrofluoric acid Hydrogen fluoride (as F) Not Available Not Available : 3 ppm / 2.6 mg/m3 Not Available **EMERGENCY LIMITS** Ingredient Material name TEEL-1 TEEL-2 TEEL-3 nitric acid Nitric acid Not Available Not Available Not Available hydrofluoric acid Hydrogen fluoride; (Hydrofluoric acid) Not Available Not Available Not Available Ingredient Original IDLH Revised IDLH nitric acid 25 ppm Not Available

Exposure controls

hydrofluoric acid

water

Use in a well-ventilated area

Type of Contaminant:

30 ppm

Not Available

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:

Not Available

Not Available

Process controls which involve changing the way a job activity or process is done to reduce the risk.

Enclosure and/or isolation of emission source which keeps a selected hazard "physically" away from the worker and ventilation that strategically "adds" and "removes" air in the work environment. Ventilation can remove or dilute an air contaminant if designed properly. The design of a ventilation system must match the particular process and chemical or contaminant in use.

Employers may need to use multiple types of controls to prevent employee overexposure.

General exhaust is adequate under normal operating conditions. Local exhaust ventilation may be required in specific circumstances. If risk of overexposure exists, wear approved respirator. Correct fit is essential to obtain adequate protection. Provide adequate ventilation in warehouse or closed storage areas. Air contaminants generated in the workplace possess varying "escape" velocities which, in turn, determine the "capture velocities" of fresh circulating air required to effectively remove the contaminant.

Appropriate engineering controls

solvent, vapours, degreasing etc., evaporating from tank (in still air).

aerosols, furnes from pouring operations, intermittent container filling, low speed conveyer transfers, welding, spray drift, plating acid furnes, pickling (released at low velocity into zone of active generation)

direct spray, spray painting in shallow booths, drum filling, conveyer loading, crusher dusts, gas discharge (active generation into zone of rapid air motion)

grinding, abrasive blasting, tumbling, high speed wheel generated dusts (released at high initial velocity into zone of very high rapid air motion).

(50-100 f/min) 0.5-1 m/s (100-200 f/min.)

Air Speed: 0.25-0.5 m/s

1-2.5 m/s (200-500 f/min.)

2.5-10 m/s (500-2000 f/min.)

Within each range the appropriate value depends on:

Lower end of the range

1: Room air currents minimal or favourable to capture

1: Disturbing room air currents

2: Contaminants of low toxicity or of nuisance value only,

3: Intermittent, low production.

3: High production, heavy use

4: Large hood or large air mass in motion

4: Small hood-local control only

Simple theory shows that air velocity falls rapidly with distance away from the opening of a simple extraction pipe. Velocity generally decreases with the square of distance from the extraction point (in simple cases). Therefore the air speed at the extraction point should be adjusted, accordingly, after reference to distance from the contaminating source. The air velocity at the extraction fan, for example, should be a minimum of 1-2 m/s (200-400 f/min) for extraction of solvents generated in a tank 2 meters distant from the extraction point. Other mechanical considerations, producing performance deficits within the extraction apparatus, make it essential that theoretical air velocities are multiplied by factors of 10 or more when extraction systems are installed or used.

Personal protection

Eye and face protection











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- Chemical goggles.
- Full face shield may be required for supplementary but never for primary protection of eyes.
- Contact lenses may pose a special hazard; soft contact tenses may absorb and concentrate irritants. A written policy document, describing the wearing of lenses or restrictions on use, should be created for each workplace or task. This should include a review of lens absorption and adsorption for the class of chemicals in use and an account of injury experience. Medical and first-aid personnel should be trained in their removal and suitable equipment should be readily available. In the event of chemical exposure, begin eye irrigation immediately and remove contact lens as soon as practicable. Lens should be removed at the first signs of eye redness or irritation lens should be removed in a clean environment only after workers have washed hands thoroughly. [CDC NIOSH Current Intelligence Bulletin 59], [AS/NZS 1336 or national equivalent]

Skin protection

See Hand protection below

- ▶ Barrier cream
- Neoprene rubber gloves

Hands/feet protection

Nitrile rubber gloves (Note: Nitric acid penetrates nitrile gloves in a few minutes.)

- ▶ Elbow length PVC gloves
- Rubber boots
- ▶ PVC safety gumboots

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Weldbrite

Body protection

See Other protection below

- Overalls.
- ▶ PVC Apron.
- PVC protective suit may be required if exposure severe.
- Other protection
 - Evewash unit. Ensure there is ready access to a safety shower.

Always ensure that a supply, is on hand, of calcium gluconate get for treatment of burns and calcium carbonate tablets for accidental ingestion.

Recommended material(s)

GLOVE SELECTION INDEX

Glove selection is based on a modified presentation of the:

"Forsberg Clothing Performance Index".

The effect(s) of the following substance(s) are taken into account in the computergenerated selection: Weldbrite

Material	CPI
NEOPRENE	А
BUTYL	С
BUTYL/NEOPRENE	C
HYPALON	C
NAT+NEOPR+NITRILE	С
NATURAL RUBBER	С
NATURAL+NEOPRENE	С
NEOPRENE/NATURAL	С
NITRILE	С
NITRILE+PVC	C
PE	С
PE/EVAL/PE	С
PVA	С
PVC	С
SARANEX-23	С
VITON	C
VITON/NEOPRENE	С

^{*} CPI - Chemwatch Performance Index

C: Poor to Dangerous Choice for other than short term immersion NOTE: As a series of factors will influence the actual performance of the glove, a final selection must be based on detailed observation. -

Respiratory protection

Type BE-P Filter of sufficient capacity. (AS/NZS 1716 & 1715, EN 143:2000 & 149:2001, ANSI Z88 or national equivalent)

Where the concentration of gas/particulates in the breathing zone, approaches or exceeds the "Exposure Standard" (or ES), respiratory protection is required. Degree of protection varies with both face-piece and Class of filter; the nature of protection varies with Type of filter.

Required Minimum Protection Factor	Half-Face Respirator	Full-Face Respirator	Powered Air Respirator
up to 10 x ES	BE-AUS P2	•	BE-PAPR-AUS / Class 1 P2
up to 50 x ES	•	BE-AUS / Class 1 P2	
up to 100 x ES	-	BE-2 P2	BE-PAPR-2 P2 ^

^ - Full-face

A(All classes) = Organic vapours, B AUS or B1 = Acid gasses, B2 = Acid gas or hydrogen cyanide(HCN), B3 = Acid gas or hydrogen cyanide(HCN), E = Sulfur dioxide(SO2), G = Agricultural chemicals, K = Ammonia(NH3), Hg = Mercury, NO = Oxides of nitrogen, MB = Methyl bromide, AX = Low boiling point organic compounds(below 65 degC)

SECTION 9 PHYSICAL AND CHEMICAL PROPERTIES

Appearance	Opaque white acidic gel; mixes with w	ater. Pungent acidic odour,	
Physical state	Non Slump Paste	Relative density (Water = 1)	1.12-1.18
Odour	Not Available	Partition coefficient n-octanol / water	Not Available
Odour threshold	Not Available	Auto-ignition temperature (°C)	Not Available
pH (as supplied)	<1	Decomposition temperature	Not Available
Melting point / freezing point (°C)	Not Available	Viscosity (cSt)	Not Available
nitial boiling point and boiling range (°C)	Not Available	Molecular weight (g/mol)	Not Applicable
Flash point (°C)	Not Applicable	Taste	Not Available
Evaporation rate	Not Available	Explosive properties	Not Available
Flammability	Not Applicable	Oxidising properties	Not Available
Upper Explosive Limit (%)	Not Applicable	Surface Tension (dyn/cm or mN/m)	Not Available
Lower Explosive Limit (%)	Not Applicable	Volatile Component (%vol)	Not Available
Vapour pressure (kPa)	Not Available	Gas group	Not Available
Solubility in water	Miscible	pH as a solution (1%)	Not available.

A: Best Selection

B: Satisfactory; may degrade after 4 hours continuous immersion

^{*} Where the glove is to be used on a short term, casual or infrequent basis, factors such as "feet" or convenience (e.g. disposability), may dictate a choice of gloves which might otherwise be unsuitable following long-term or frequent use. A qualified practitioner should be consulted.

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Vapour density (Air = 1) Not available.

VOC g/L Not Available

SECTION 10 STABILITY AND REACTIVITY

Reactivity

See section 7

- Unstable in the presence of incompatible materials. Chemical stability
 - Product is considered stable.
 - Hazardous polymerisation will not occur.

Possibility of hazardous reactions

See section 7

Conditions to avoid Incompatible materials

See section 7 See section 7

Hazardous decomposition products

See section 5

SECTION 11 TOXICOLOGICAL INFORMATION

Inhaled

Information on toxicological effects

There is some evidence to suggest that the material can cause respiratory irritation in some persons. The body's response to such irritation can cause further lung damage.

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 hazard is increased at higher temperatures.

Acute effects of fluoride inhalation include irritation of nose and throat, coughing and chest discomfort. A single acute over-exposure may even

cause nose bleed. Acute inhalation of hydrogen fluoride (hydrofluoric acid) vapours causes severe irritation of the eye, nose and throat, delayed fever, bluing of the

extremities and water in the lungs, and may cause death. The above imitation occurs even with fairly low concentrations of hydrogen fluoride. Hydrogen fluoride has a strong irritating odour, that can be detected at concentrations of about 0.04 parts per million. Higher fevels cause corrosion of the throat, nose and lungs, leading to severe inflammation and water buildup in the lungs (which may occur with 1 hour of exposure). A vapour concentration of 10 parts per million is regarded as intolerable, but a vapour concentration of 30 parts per million is considered as immediately dangerous to life and health.

It is estimated that the lowest tethal concentration for a 5-minute human exposure to hydrogen fluoride is in the range of 50 to 250 parts per million. Exposure by either skin contact or inhalation may lead to low levels of calcium and magnesium in the blood, which may result in heart rhythm disturbances. Animal testing suggests that repeated exposure produces liver and kidney damage.

Inhalation of nitric acid mist or furnes may produce respiratory symptoms. Depending on the concentration and duration of exposure, cough, gagging, chest pain, low body oxygen, lung imitation and damage may occur. Deaths have occurred and may be delayed for several days. inhalation of aerosols (mists, furnes), generated by the material during the course of normal handling, may produce severely toxic effects. Relatively small amounts absorbed from the lungs may prove fatal.

Severely toxic effects may result from the accidental ingestion of the material; animal experiments indicate that ingestion of less than 5 gram may be fatal or may produce serious damage to the health of the individual.

Ingestion of acidic corrosives may produce burns around and in the mouth, the throat and desophagus. Immediate pain and difficulties in swallowing and speaking may also be evident.

Ingestion

Fluoride causes severe loss of calcium in the blood, with symptoms appearing several hours later including painful and rigid muscle contractions of the limbs. Cardiovascular collapse can occur and may cause death with increased heart rate and other heart rhythm irregularities. Exposure to nitric acid causes burning pain, severe corrosion and scaring of the digestive tract with adhesions, narrowing and obstruction and even anaemia. There may be vomiting, aspiration, lung inflammation and shock. Death may be delayed 12 hours to 14 days or several months from these complications. Survivors may have strictures of the stornach fining and subsequent pernicious anaemia. fatal if swallowed unless immediate treatment is applied

Skin contact with the material may produce severely toxic effects; systemic effects may result following absorption and these may be fatat. Skin contact with acidic corrosives may result in pain and bums; these may be deep with distinct edges and may heal slowly with the formation of

Fluorides are easily absorbed through the skin and cause death of soft tissue and erode bone. Healing is delayed and death of tissue may continue to spread beneath skin.

Open cuts, abraded or irritated skin should not be exposed to this material

Contact of the skin with liquid hydrofluoric acid (hydrogen fluoride) may cause severe burns, erythema, and swelling, vesiculation, and serious crusting. With more serious burns, ulceration, blue-gray discoloration, and necrosis may occur. Solutions of hydrofluoric acid, as dilute as 2%, may cause severe skin burns.

Dermat burns may not be readily noticed or painful, unlike the warning properties of other acids. Skin contact with HF concentrations in the 20% to 50% range may not produce symptoms for one to eight hours. With concentrations less than 20%, the latency period may be up to twenty-four hours. A solution of only 1-2% HF exposed to greater than 10% of the body is fatal without medical attention; however dermal burns are not

Skin Contact

Fluoride ions form insoluble salts with calcium and magnesium in bodily tissue. Soluble salts can form with other cations, which dissociate rapidly causing further disruption and damage to tissue. The severe, throbbing pain associated with HF burns is thought to result from nerve imitation due to potassium cations entering the extracellular space to compensate for reduced calcium ion concentrations. Fluoride poisoning is associated with hypocalcemia (low calcium levels), hyperkalemia (high potassium levels), hypomagnesemia (low magnesium levels), and sudden

Systemic hypocalcemia should be considered a risk whenever the body surface area of skin burns from concentrated hydrofluoric acid exceed 160 cm2 (25 sq.in).), or about the size of the palm of your hand. Concentrated hydrogen fluoride burns can be fatal if only 2% of the body surface area is exposed.

Entry into the blood-stream, through, for example, cuts, abrasions or lesions, may produce systemic injury with harmful effects. Examine the skin prior to the use of the material and ensure that any external damage is suitably protected.

Skin contact with nitric acid may cause corrosion, skin thickening, yellow discolouration of the skin, blisters and scars depending on the concentration exposed.

Direct eye contact with acid corrosives may produce pain, tears, sensitivity to light and burns. Mild burns of the epithelia generally recover rapidly

Animal testing showed that a 20% solution of hydrofluoric acid (hydrogen fluoride) in water caused immediate damage in the form of total clouding of the lens and ischaemia of the conjunctiva. Swelling of the stroma of the cornea occurred within 1 hour, followed by tissue death (necrosis) of structures of the front of the eye.

Eye contact with bolh diluted and concentrated nitric acid may result in burns causing pain, adhesions, corneal damage, blindness or permanent eye damage. Pain may be absent after contact with concentrated nitric acid.

The vapour may produce pronounced discomfort of the eyes when present at higher concentrations and this generally gives warning of excessive exposure and the need for control measures to ensure safe working conditions

Chronic

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Repeated or prolonged exposure to acids may result in the erosion of teeth, swelling and/or ulceration of mouth lining. Imitation of airways to lung, with cough, and inflammation of lung tissue often occurs.

Long-term exposure to respiratory irritants may result in airways disease, involving difficulty breathing and related whole-body problems. Extended exposure to inorganic fluorides causes fluorosis, which includes signs of joint pain and stiffness, tooth discolouration, nausea and vomiting, loss of appetite, diarrhoea or constipation, weight loss, anaemia, weakness and general unwellness. There may also be frequent urination and thirst.

Prolonged or repeated overexposure to low concentrations of nitric acid vapour may cause chronic airway inflammation, corrosion of teeth and chemical lung inflammation.

Hydrogen fluoride easily penetrates the skin and causes destruction and corrosion of the bone and underlying tissue. Ingestion causes severe pains and burns in the mouth and throat and blood calcium levels are dangerously reduced.

The state of the second se	pains and burns in the mouth and throat and blood calcit	um fevels are dangerously reduced.
	тохісіту	IRRITATION
Weldbrite	Not Available	Not Available
	тохісіту	IRRITATION
nitric acid	50500 mg/kg ^[2]	Eye: adverse effect observed (imitating)[1]
	Inhalation (rat) LC50: 0.13 mg/l/4h ^[2]	Skin: adverse effect observed (corrosive)[1]
	τοχισιτγ	IRRITATION
hydrofluoric acid	Inhalation (rat) LC50: 0.275 mg/l/60M ^[2]	Eye (human): 50 mg - SEVERE
water	TOXICITY	RRITATION
*****	Oral (rat) LD50: >90000 mg/kg ^[2]	Not Available
Legend:	1 Value obtained from Europe ECHA Registered Substa specified data extracted from RTECS - Register of Toxic	nces - Acute toxicity 2.* Value obtained from manufacturer's SDS. Unless otherwise Effect of chemical Substances
NITRIC ACID	not been examined in this respect. Mucous secretion may protects the stomach lining from the hydrochloric acid sec	to genetic damage when the pH falls to about 6.5. Cells from the respiratory tract have y protect the cells of the airway from direct exposure to inhaled acidic mists (which also creted there). ged or repeated exposure and may produce on contact skin redness, swelling, the

The material may cause severe 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. Repeated exposures may produce severe ulceration.

(liver and kidney damage) [Manufacturer] for hydrogen fluoride (as vapour)

Laboratory (in vitro) and animal studies show, exposure to the material may result in a possible risk of irreversible effects, with the possibility of producing mutation.

The following information refers to contact allergens as a group and may not be specific to this product.

HYDROFLUORIC ACID

NITRIC ACID & HYDROFLUORIC ACID

Contact allergies quickly manifest themselves as contact eczema, more rarely as unicaria or Quincke's oedema. The palhogenesis of contact eczema involves a cell-mediated (T lymphocytes) immune reaction of the delayed type. Other allergic skin reactions, e.g. contact urticaria, involve antibody-mediated immune reactions. The significance of the contact allergen is not simply determined by its sensitisation potential; the distribution of the substance and the opportunities for contact with it are equally important. A weakly sensitising substance which is widely distributed can be a more important allergen than one with stronger sensitising potential with which few individuals come into contact. From a clinical point of view, substances are noteworthy if they produce an allergic test reaction in more than 1% of the persons tested.

Asthma-like symptoms may continue for months or even years after exposure to the material ends. This may be due to a non-allergic condition known as reactive airways dysfunction syndrome (RADS) which can occur after exposure to high levels of highly irritating compound. Main criteria for diagnosing RADS include the absence of previous airways disease in a non-atopic individual, with sudden onset of persistent asthma-like symptoms within minutes to hours of a documented exposure to the irritant. Other criteria for diagnosis of RADS include a reversible airflow pattern on lung function tests, moderate to severe bronchial hyperreactivity on methacholine challenge testing, and the lack of minimal lymphocytic inflammation, without eosinophilia. 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. On the other hand, industrial bronchitis is a disorder that occurs as a result of exposure due to high concentrations of irritating substance (often particles) and is completely reversible after exposure ceases. The disorder is characterized by difficulty breathing, cough and mucus production.

The material may produce severe irritation to the eye causing pronounced inflammation. Repeated or prolonged exposure to irritants may produce conjunctivitis.

The material may produce respiratory tract irritation, and result in damage to the lung including reduced lung function.

HYDROFLUORIC ACID & WATER

No significant acute toxicological data identified in literature search.

Acute Toxicity	₩.	Carcinogenicity	×
Skin Irritation/Corrosion	₩.	Reproductivity	×
Serious Eye Damage/Irritation	<^	STOT - Single Exposure	X
Respiratory or Skin sensitisation	49	STOT - Repeated Exposure	×
Mutagenicity	V	Aspiration Hazard	×

Legend: 💢 - Data either not available or does not fill the critena for classification

Data available to make classification

SECTION 12 ECOLOGICAL INFORMATION

Toxicity					
	ENDPOINT	TEST DURATION (HR)	SPECIES	VALUE	SOURCE
Weldbrite	Not Available	Not Available	Not Available	Not Available	Not Available

	ENDPOINT	TEST DURATION (HR)	SPECIES	VALUE	SOURCE
	LC50	96	Fish		
nîtrîc acid	1	free contract the second contract to	$\Phi_{ij} = \frac{1}{2} \left(\frac{1}{2} \right) \right) \right) \right) \right)}{1} \right) \right)}{1} \right) \right)} \right) \right)} \right) \right)} \right) \right)} \right) \right) \right) \right)}$	1-354mg/L	2
	EC50	48	Crustacea	490mg/L	2
	NOEC	720	Fish	58mg/L	. 2
		1	·	4	:
	ENDPOINT	TEST DURATION (HR)	SPECIES	VALUE	SOURCE
	LC50	96	Fish	51mg/L	2
hydrofluoric acid	EC50	48	Crustacea	97mg/L	2
	EC50	96	Algae or other aquatic plants	43mg/L	2
	NOEC	504	Crustacea	3.7mg/L	2
		*		:	
	ENDPOINT	TEST DURATION (HR)	SPECIES	VALUE	SOURCE
	LC50	96	Fish	897.520mg/L	3
water		in Table and the second and the second and the second			
	EC50	96	Algae or other aquatic plants	8768.874mg/L	3
Legend: DO NOT discharge into sewer or wa	EC50 Extracted from V3.12 (QSAR) - Data 6. NITE (J		Algae or other aquatic plants istered Substances - Ecotoxicological Informa . Ecotox database - Aquatic Toxicity Data 5. E	ition - Aquatic Toxicity 3. I	EPIWIN Sun
Legend:	EC50 Extracted from V3.12 (QSAR) - Data 6. NITE (J	96 1 IUCLID Toxicity Data 2. Europe ECHA Reg. Aquatic Toxicity Data (Estimated) 4. US EPA	Algae or other aquatic plants istered Substances - Ecotoxicological Informa . Ecotox database - Aquatic Toxicity Data 5. E	ition - Aquatic Toxicity 3. I	EPIWIN Suit
Legend: Legend: DO NOT discharge into sewer or wa	EC50 Extracted from V3.12 (QSAR) - Data 6. NITE (J	96 1 IUCLID Toxicity Data 2. Europe ECHA Reg. Aquatic Toxicity Data (Estimated) 4. US EPA apan) - Bioconcentration Data 7. METI (Japan	Algae or other aquatic plants istered Substances - Ecotoxicological Informa . Ecotox database - Aquatic Toxicity Data 5. E	ition - Aquatic Toxicity 3. I	EPIWIN Suit
Legend: DO NOT discharge into sewer or we ersistence and degradability	EC50 Extracted from V3.12 (OSAR) - Data 6. NITE (J	96 1 IUCLID Toxicity Data 2. Europe ECHA Reg. Aquatic Toxicity Data (Estimated) 4. US EPA apan) - Bioconcentration Data 7. METI (Japan	Algae or other aquatic plants istered Substances - Ecotoxicological Informa , Ecotox database - Aquatic Toxicity Data 5. E a) - Bioconcentiation Data 8. Vendor Data	ition - Aquatic Toxicity 3. I	EPIWIN Suit
Legend: DO NOT discharge into sewer or was ersistence and degradability Ingredient	EC50 Extracted from V3.12 (QSAR) - Data 6. MITE (Jatorways. Persistence: W	96 1 IUCLID Toxicity Data 2. Europe ECHA Reg. Aquatic Toxicity Data (Estimated) 4. US EPA apan) - Bioconcentration Data 7. METI (Japan	Algae or other aquatic plants istered Substances - Ecotoxicological Informa, Ecotox database - Aquatic Toxicity Data 5. E a) - Bioconcentiation Data 8. Vendor Data Persistence: Air	ition - Aquatic Toxicity 3. I	EPIWIN Suit
Legend: DO NOT discharge into sewer or was ersistence and degradability Ingredient water	EC50 Extracted from V3.12 (QSAR) - Data 6. MITE (Jatorways. Persistence: W	96 1 IUCLID Toxicity Data 2. Europe ECHA Reg. Aquatic Toxicity Data (Estimated) 4. US EPA (apan) - Bioconcentration Data 7. METI (Japan fater/Soil	Algae or other aquatic plants istered Substances - Ecotoxicological Informa, Ecotox database - Aquatic Toxicity Data 5. E a) - Bioconcentiation Data 8. Vendor Data Persistence: Air	ition - Aquatic Toxicity 3. I	EPIWIN Suit
Legend: DO NOT discharge into sewer or water Ingredient water ioaccumulative potential	EC50 Extracted from V3.12 (QSAR) - Data 6. NITE (Jaterways. Persistence: W	96 1 IUCLID Toxicity Data 2. Europe ECHA Reg. Aquatic Toxicity Data (Estimated) 4. US EPA (apan) - Bioconcentration Data 7. METI (Japan fater/Soil	Algae or other aquatic plants istered Substances - Ecotoxicological Informa, Ecotox database - Aquatic Toxicity Data 5. E a) - Bioconcentiation Data 8. Vendor Data Persistence: Air	ition - Aquatic Toxicity 3. I	EPIWIN Sun
Legend: DO NOT discharge into sewer or water water ioaccumulative potential	EC50 Extracted from V3.12 (OSAR) - Data 6. NITE (Jatonways. Persistence: W. LOW.	96 1 IUCLID Toxicity Data 2. Europe ECHA Reg. Aquatic Toxicity Data (Estimated) 4. US EPA (apan) - Bioconcentration Data 7. METI (Japan fater/Soil	Algae or other aquatic plants istered Substances - Ecotoxicological Informa, Ecotox database - Aquatic Toxicity Data 5. E a) - Bioconcentiation Data 8. Vendor Data Persistence: Air	ition - Aquatic Toxicity 3. I	EPIWIN Suit
Legend: DO NOT discharge into sewer or water state water ioaccumulative potential ingredient water	EC50 Extracted from V3.12 (OSAR) - Data 6. NITE (Jatonways. Persistence: W. LOW.	96 1 IUCLID Toxicity Data 2. Europe ECHA Reg. Aquatic Toxicity Data (Estimated) 4. US EPA (apan) - Bioconcentration Data 7. METI (Japan fater/Soil	Algae or other aquatic plants istered Substances - Ecotoxicological Informa, Ecotox database - Aquatic Toxicity Data 5. E a) - Bioconcentiation Data 8. Vendor Data Persistence: Air	ition - Aquatic Toxicity 3. I	EPIWIN Suit

SECTION 13 DISPOSAL CONSIDERATIONS

Waste treatment methods

- Recycle wherever possible or consult manufacturer for recycling options.
- Consult State Land Waste Management Authority for disposal.
 Treat and neutralise at an effluent treatment plant.
- Product / Packaging disposal
 - Use soda ash or slaked lime to neutralise.
 - Recycle containers, otherwise dispose of in an authorised landfill.

SECTION 14 TRANSPORT INFORMATION

Labels Required



Marine Pollutant

NO 2X

HAZCHEM

Land transport (ADG)

UN number	2922		
UN proper shipping name	CORROSIVE LIQUID, TOXIC, N.O.S. (contains nitric acid and hydrofluoric acid)		
Transport hazard class(es)	Class 8		
	Subrisk 6,1		

Packing group Environmental hazard

Not Applicable

Special precautions for user

Special provisions 274
Limited quantity 1 L

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Air transport (ICAO-IATA / DGR)

UN number	2922		
UN proper shipping name	Corrosive liquid, toxic, n.o.s. * (contains nitric acid and hydrofluoric acid)		
Transport hazard class(es)	ICAO/IATA Class 8 ICAO / IATA Subrisk 6.1 ERG Code 8P		
Packing group	п		
Environmental hazard	Not Applicable		
	Special provisions	A3 A803	
	Cargo Only Packing Instructions	855	
	Cargo Only Maximum Qty / Pack	30 L	
Special precautions for user	Passenger and Cargo Packing Instructions	851	
	Passenger and Cargo Maximum Qty / Pack	1 L	
	Passenger and Cargo Limited Quantity Packing Instructions	Y840	
	Passenger and Cargo Limited Maximum Qty / Pack	0,5 L	

Sea transport (IMDG-Code / GGVSee)

UN number	2922
UN proper shipping name	CORROSIVE LIQUID, TOXIC, N.O.S. (contains nitric acid and hydrofluoric acid)
Transport hazard class(es)	IMDG Class 8 IMDG Subrisk 6.1
Packing group	
Environmental hazard	Not Applicable
Special precautions for user	EMS Number F-A , S-B Special provisions 274 Limited Quantities 1 L

Transport in bulk according to Annex II of MARPOL and the IBC code Not Applicable

SECTION 15 REGULATORY INFORMATION

Safety, health and environmental regulations / legislation specific for the substance or mixture

\S NITRIC ACID IS FOUND ON THE FOLLOWING REGULATORY LISTS

Australia Hazardous Chemical Information System (HCIS) - Hazardous Chemicals Australia Inventory of Chemical Substances (AICS)

Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) - Schedule 5

HYDROFLUORIC ACID IS FOUND ON THE FOLLOWING REGULATORY LISTS

Australia Hazardous Chemical Information System (HCIS) - Hazardous Chemicals Australia Inventory of Chemical Substances (AICS)

Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) - Schedule 2

Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) - Schedule $\bf 3$

Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) - Schedule 4

Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) - Schedule 5 $\,$

Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) - Schedule 6

Australia Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) - Schedule 7

International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Monographs

WATER IS FOUND ON THE FOLLOWING REGULATORY LISTS

Australia Inventory of Chemical Substances (AICS)

ECHA SUMMARY

Ingredient	CAS number	Index No I	CHA Dossier	
nitric acid	7697-37-2	007-004-00-1	1-2119487297-23-XXXX 01-2119987992-14-XXXX	01-2120763162-60-XXXX
Harmonisation (C&L Inventory)	Hazard Class and Category Code(s)		Pictograms Signal Word Code(s)	Hazard Statement Code(s)
1	Ox. Sol. 2; Skin Corr. 1A		GHS03; GHS05; Dgr	H272; H314
Harmonisation Code 1 = The	most provalent classificatio	n Harmonisation Cods 2 =	The most severe classification	
Ingredient	CAS number	Index No	ECHA Dossier	
hydrofluoric acid	7664-39-3	009-002-00-6 009-00	3-00-1 01-2119458860-33-XXXXI0	11-2120762785-41-XXXX

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Harmonisation (C&L Inventory)	Hazard Class and Category Code(s)		Pictograms Signal Word Code(s) Hazard Statement Code(s)
1	Acute Tox, 2; Acute Tox, 1; Skin Corr, 1A; A	cute Tox. 2	GHS05; GHS06; Dgr	H300; H310; H314; H330
1	Acute Tox, 2; Acute Tox, 1; Skin Corr, 1A; Acute Tox, 2		GHS05; GHS06; Dgr	H300; H310; H314; H330
1	Acute Tox. 2; Acute Tox. 1; Skin Corr. 1A; Acute Tox. 2		GHS05; GHS06; Dgr	H300; H310; H314; H330
Harmonisation Code 1 = The mos	t prevalent classification. Harmonisation Code 2	? = The most severe (dassification.	
ngredient	CAS number	Index No	ECH	A Dossier
vater	7732-18-5	Not Available	Not /	Available
farmonisation (C&L nventory)	Hazard Class and Category Code(s)	Pictogra	ns Signal Word Code(s)	Hazard Statement Code(s)
the second control of the second seco	Not Classified	Not Avail	able	Not Available
	t prevalent classification. Harmonisation Code 2	t = The most severe c	lassification.	
itional Inventory Status	t prevalent classification. Harmonisation Code 2 Status	? = The most severe o	lassification.	
tional Inventory Status lational Inventory		= The most severe a	lassification.	
tional Inventory Status lational Inventory ustralia - AICS	Status	= The most severe o	lassification.	
tional Inventory Status lational Inventory ustralia - AICS canada - DSL	Status Yes	P = The most severe of	lassification.	
tional Inventory Status lational Inventory lustralia - AICS lanada - DSL lanada - NDSL	Status Yes Yes	P = The most severe of	lassification.	
tional Inventory Status lational Inventory kustralia - AICS Canada - DSL Canada - NDSL China - IECSC	Status Yes Yes No (nitric acid; hydrofluoric acid; water)	= The most severe s	lassification.	
tional Inventory Status lational Inventory ustralia - AICS lanada - DSL lanada - NDSL hina - IECSC urope - EINEC / ELINCS / NLP	Status Yes Yes No (nitric acid; hydrofiuoric acid; water) Yes	P = The most severe of	lassification.	
tional Inventory Status lational Inventory ustralia - AICS lanada - DSL lanada - NDSL lhina - IECSC urope - EINEC / ELINCS / NLP	Status Yes Yes No (nitric acid; hydrofluoric acid; water) Yes Yes	P = The most severe of	lassification.	
tional Inventory Status lational Inventory lational Inventory lational Inventory lational - AICS lanada - DSL lanada - NDSL lanada - NDSL lanada - NECSC lurope - EINEC / ELINCS / NLP lapan - ENCS lorea - KECI	Status Yes Yes No (nitric acid; hydrofiuoric acid; water) Yes Yes	P = The most severe s	lassification.	
Ational Inventory Status National Inventory Australia - AICS Canada - DSL Canada - NDSL China - IECSC Europe - EINEC / ELINCS / NLP Iapan - ENCS Korea - KECI Iew Zealand - NZIoC	Status Yes Yes No (nitric acid; hydrofluoric acid; water) Yes Yes Yes	P = The most severe s	lassification.	
Attional Inventory Status National Inventory Status National Inventory Australia - AICS Canada - DSL Canada - NDSL China - IECSC Europe - EINEC / ELINCS / NLP Iapan - ENCS Korea - KECI New Zealand - NZIoC Phillippines - PICCS USA - TSCA	Status Yes Yes No (nitric acid; hydrofluoric acid; water) Yes Yes Yes Yes Yes	P= The most severe s	lassification.	

SECTION 16 OTHER INFORMATION

Yes

Yes

Yes

Yes = All CAS declared ingredients are on the inventory

	y		
Revision Date	07/03/2020		
Initial Date	30/12/2002		

No = One or more of the CAS listed ingredients are not on the inventory and are not exempt from listing(see specific ingredients in brackets)

SDS Version Summary

Mexico - INSQ

Vietnam - NCI

Russia - ARIPS

Leaend:

Version	Issue Date	Sections Updated	
8.1.1.1	15/01/2020	Acute Health (inhaled)	
9.1.1.1	07/03/2020	Classification change due to full database hazard calculation/update.	

Other information

Classification of the preparation and its individual components has drawn on official and authoritative sources as well as independent review by the Chemwatch Classification committee using available literature references.

The SDS is a Hazard Communication tool and should be used to assist in the Risk Assessment. Many factors determine whether the reported Hazards are Risks in the workplace or other settings. Risks may be determined by reference to Exposures Scenarios. Scale of use, frequency of use and current or available engineering controls must be considered.

Definitions and abbreviations

PC-TWA: Permissible Concentration-Time Weighted Average

PC-STEL: Permissible Concentration-Short Term Exposure Limit

IARC: International Agency for Research on Cancer

ACGIH: American Conference of Governmental Industrial Hygienists

STEL: Short Term Exposure Limit
TEEL: Temporary Emergency Exposure Limit.

IDLH: Immediately Dangerous to Life or Health Concentrations

OSF: Odour Safety Factor

NOAEL :No Observed Adverse Effect Level

LOAEL: Lowest Observed Adverse Effect Level

TLV: Threshold Limit Value

LOD: Limit Of Detection

OTV: Odour Threshold Value

BCF: BioConcentration Factors

BEI: Biological Exposure Index

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