

High-Purity Standards

Catalogue number: CWW-TM-C

Version No: 1.1 Safety Data Sheet according to OSHA HazCom Standard (2012) requirements

SECTION 1 IDENTIFICATION

Product Identifier

Product name	CWW-TM-C - Certified Waste Water Trace Metals Solution C
Synonyms	CWW-TM-C
Proper shipping name	Corrosive liquid, acidic, inorganic, n.o.s.
Other means of identification	CWW-TM-C

Recommended use of the chemical and restrictions on use

Relevant identified uses Use according to manufacturer's directions.

Name, address, and telephone number of the chemical manufacturer, importer, or other responsible party

Registered company name	High-Purity Standards
Address	PO Box 41727 SC 29423 United States
Telephone	843-767-7900
Fax	843-767-7906
Website	highpuritystandards.com
Email	Not Available

Emergency phone number

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Association / Organisation	INFOTRAC
Emergency telephone numbers	1-800-535-5053
Other emergency telephone numbers	1-352-323-3500

SECTION 2 HAZARD(S) IDENTIFICATION

H314

Causes severe skin burns and eye damage.

Classification of the subst	Serious Eye Damage Category 1, Acute Toxicity (Oral) Category 4, Acute Toxicity (Dermal) Category 4, Metal Corrosion Category 1, Skin Corrosion/Irritation Category 1A
Label elements	
Hazard pictogram(s)	
SIGNAL WORD	DANGER
Hazard statement(s)	
H302	Harmful if swallowed.
H312	Harmful in contact with skin.
H290	May be corrosive to metals.

Chemwatch Hazard Alert Code: 3

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Hazard(s) not otherwise specified

Not Applicable

Precautionary statement(s) Prevention

P260	Do not breathe dust/fume/gas/mist/vapours/spray.	
Precautionary statement(s)	Response	
P301+P330+P331 IF SWALLOWED: Rinse mouth. Do NOT induce vomiting.		
Precautionary statement(s) P405	Storage Store locked up.	
Precautionary statement(s)	Disposal	
P501	Dispose of contents/container in accordance with local regulations.	

SECTION 3 COMPOSITION / INFORMATION ON INGREDIENTS

Substances

See section below for composition of Mixtures

Mixtures

CAS No	%[weight]	Name
7429-90-5	0.005	aluminium
7440-36-0	0.0015	antimony
7440-38-2	0.0015	arsenic
7440-39-3	0.005	barium
7440-41-7	0.0015	beryllium
7440-42-8	0.005	boron
7440-43-9	0.0015	cadmium
7440-47-3	0.005	<u>chromium</u>
7440-48-4	0.005	cobalt
7440-50-8	0.005	copper
7439-89-6	0.005	iron
7439-92-1	0.005	lead
6156-78-1	0.005	manganese(II) acetate tetrahydrate
7439-97-6	0.0001	mercury (elemental)
7439-98-7	0.005	molybdenum
7440-02-0	0.005	nickel
7782-49-2	0.0015	selenium
7440-22-4	0.0015	silver
7440-24-6	0.005	strontium
7440-28-0	0.0015	thallium
7803-55-6	0.005	ammonium metavanadate
7440-66-6	0.005	zinc
7697-37-2	10	nitric acid
7664-39-3	0-0.49	hydrofluoric acid
7732-18-5	balance	water

SECTION 4 FIRST-AID MEASURES

Description of first aid measures

Eye Contact	 If this product comes in contact with the eyes: Immediately hold eyelids apart and flush the eye continuously with running water. Ensure complete irrigation of the eye by keeping eyelids apart and away from eye and moving the eyelids by occasionally lifting the upper and lower lids. Continue flushing until advised to stop by the Poisons Information Centre or a doctor, or for at least 15 minutes. Transport to hospital or doctor without delay. Removal of contact lenses after an eye injury should only be undertaken by skilled personnel.
Skin Contact	 For thermal burns: Decontaminate area around burn. Consider the use of cold packs and topical antibiotics. For first-degree burns (affecting top layer of skin) Hold burned skin under cool (not cold) running water or immerse in cool water until pain subsides. Use compresses if running water is not available.

	 Cover with sterile non-adhesive bandage or clean cloth. Do NOT apply butter or ointments; this may cause infection. Give over the counter pain increases or swelling, redness, fever occur. For second-degree burns (affecting top two layers of skin) Cool the burn by immerse in cold running water for 10-15 minutes. Use compresses if running water is not available. Do NOT apply ice as this may lower body temperature and cause further damage. Do NOT paphy ice as this may lower body temperature and cause further damage. Do NOT paphy ice as this may lower body temperature and cause further damage. Do NOT paphy ice as this temile, nonstick bandage and secure in place with gauze or tape. To prevent shock: (unless the person has a head, neck, or leg injury, or it would cause discomfort): Lay the person flat. Elevate feet about 12 inches. Elevate feet about 12 inches. Elevate heat level, if possible. Cover the person with coat or blanket. Seek medical assistance. For third-degree burns Seek medical cor emergency assistance. In the mean time: Protect burn area acover loosely with sterile, nonstick bandage or, for large areas, a sheet or other material that will not leave lint in wound. Separate burne or apply other or apply other or spins in may cause infection. For an airway burn, do not place pillow under the person's head when the person is tying down. This can dose the airway. Have a person with a facial burn sit up. Check pulse and breathing to monitor for shock until emergency help arrives. If there is evidence of severe skin iritation or skin curns: Avoid impredive framework containsted clothing, including footwear. Flues is widence of severe skin iritation or skin curns. Avoid impredive for to tab, many asses calcium gluconate gel into affected areas,
Inhalation	 If fumes or combustion products are inhaled remove from contaminated area. Lay patient down. Keep warm and rested. Prostheses such as false teeth, which may block airway, should be removed, where possible, prior to initiating first aid procedures. Apply artificial respiration if not breathing, preferably with a demand valve resuscitator, bag-valve mask device, or pocket mask as trained. Perform CPR if necessary. Transport to hospital, or doctor, without delay. Inhalation of vapours or aerosols (mists, fumes) may cause lung oedema. Corrosive substances may cause lung damage (e.g. lung oedema, fluid in the lungs). As this reaction may be delayed up to 24 hours after exposure, affected individuals need complete rest (preferably in semi-recumbent posture) and must be kept under medical observation even if no symptoms are (yet) manifested. Before any such manifestation, the administration of a spray containing a dexamethasone derivative or beclomethasone derivative may be considered. This must definitely be left to a doctor or person authorised by him/her. (ICSC13719) For massive exposures: If dusts, vapours, aerosols, fumes or combustion products are inhaled, remove from contaminated area. Lay patient down. Keep warm and rested. Prostheses such as false teeth, which may block airway, should be removed, where possible, prior to initiating first aid procedures. Apply artificial respiration if not breathing, preferably with a demand valve resuscitator, bag-valve mask device, or pocket mask as trained. Perform CPR if necessary. If victim is conscious, give six calcium gluconate or calcium carbonate tablets in water by mouth. Transport to hospital, or doctor, urgenty.
Ingestion	 For advice, contact a Poisons Information Centre or a doctor at once. Urgent hospital treatment is likely to be needed. If swallowed do NOT induce vomiting. If vomiting occurs, lean patient forward or place on left side (head-down position, if possible) to maintain open airway and prevent aspiration. Observe the patient carefully. Never give liquid to a person showing signs of being sleepy or with reduced awareness; i.e. becoming unconscious. Give water to rinse out mouth, then provide liquid slowly and as much as casualty can comfortably drink. Transport to hospital or doctor without delay.

Most important symptoms and effects, both acute and delayed

See Section 11

Indication of any immediate medical attention and special treatment needed

Following acute or short term repeated exposure to hydrofluoric acid:

- Subcutaneous injections of Calcium Gluconate may be necessary around the burnt area. Continued application of Calcium Gluconate Gel 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

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

Determinant	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.

For acute or short term repeated exposures to fluorides:

- Fluoride absorption from gastro-intestinal tract may be retarded by calcium salts, milk or antacids.
- Fluoride particulates or fume may be absorbed through the respiratory tract with 20-30% deposited at alveolar level.
- ▶ Peak serum levels are reached 30 mins. post-exposure; 50% appears in the urine within 24 hours.
- For acute poisoning (endotracheal intubation if inadequate tidal volume), monitor breathing and evaluate/monitor blood pressure and pulse frequently since shock may supervene with little warning. Monitor ECG immediately; watch for arrhythmias and evidence of Q-T prolongation or T-wave changes. Maintain monitor. Treat shock vigorously with isotonic saline (in 5% glucose) to restore blood volume and enhance renal excretion.
- > Where evidence of hypocalcaemic or normocalcaemic tetany exists, calcium gluconate (10 ml of a 10% solution) is injected to avoid tachycardia.

BIOLOGICAL EXPOSURE INDEX - BEI

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

Determinant Fluorides in urine	Index 3 mg/gm creatinine 10mg/gm creatinine	Sampling Time Prior to shift End of shift	Comments B, NS B, NS
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B: Background levels occur in specimens collected from subjects **NOT** exposed

NS: Non-specific determinant; also observed after exposure to other exposures.

SECTION 5 FIRE-FIGHTING MEASURES

Extinguishing media

- Jets of water.
- Water spray or fog.
- Foam.
- Dry chemical powder.
- BCF (where regulations permit).
- Carbon dioxide.

Special hazards arising from the substrate or mixture

Fire Incompatibility	None known.
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Special protective equipment and precautions for fire-fighters

Fire Fighting	
Fire/Explosion Hazard	 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. May emit corrosive, poisonous fumes. May emit acrid smoke. When aluminium oxide dust is dispersed in air, firefighters should wear protection against inhalation of dust particles, which can also contain hazardous substances from the fire absorbed on the alumina particles.

SECTION 6 ACCIDENTAL RELEASE MEASURES

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	 Drains for storage or use areas should have retention basins for pH adjustments and dilution of spills before discharge or disposal of material. Check regularly for spills and leaks. Clean up all spills immediately. Avoid breathing vapours and contact with skin and eyes. Control personal contact with the substance, by using protective equipment. Contain and absorb spill with sand, earth, inert material or vermiculite. Wipe up. Place in a suitable, labelled container for waste disposal.
Major Spills	 Clear area of personnel and move upwind. Alert Fire Brigade and tell them location and nature of hazard. Wear breathing apparatus plus protective gloves. Prevent, by any means available, spillage from entering drains or water course. Stop leak if safe to do so. Contain spill with sand, earth or vermiculite. Collect recoverable product into labelled containers for recycling.

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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 Avoid all personal contact, including inhalation. Wear protective clothing when risk of exposure occurs. Use in a well-ventilated area. Avoid contact with moisture. Avoid contact with incompatible materials. When handling, DO NOT eat, drink or smoke. Safe handling Keep containers securely sealed when not in use. Avoid physical damage to containers. Always wash hands with soap and water after handling. Work clothes should be laundered separately. Launder contaminated clothing before re-use. 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. Store in original containers. ۲ Keep containers securely sealed. Store in a cool, dry, well-ventilated area. Other information 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

Conditions for safe storage	ge, including any incompatibilities
Suitable container	 DO NOT use aluminium or galvanised containers Check regularly for spills and leaks Lined metal can, lined metal pail/ can. Plastic pail. Polyliner drum. Packing as recommended by manufacturer. Check all containers are clearly labelled and free from leaks. For low viscosity materials Drums and jerricans must be of the non-removable head type. Where a can is to be used as an inner package, the can must have a screwed enclosure. For materials with a viscosity of at least 2680 cSt. (23 deg. C) and solids (between 15 C deg. and 40 deg C.): Removable head packaging; Cans with friction closures and low pressure tubes and cartridges may be used. - Where combination packages are used, and the inner packages are of glass, porcelain or stoneware, there must be sufficient inert cushioning material in contact with inner and outer packages unless the outer packaging is a close fitting moulded plastic box and the substances are not incompatible with the plastic. Material is corrosive to most metals, glass and other siliceous materials.
Storage incompatibility	 For aluminas (aluminium oxide): Incompatible with hot chlorinated rubber. In the presence of chlorine trifluoride may react violently and ignite. May initiate explosive polymerisation of olefin oxides including ethylene oxide. -Produces exothermic reaction above 200 C with halocarbons and an exothermic reaction at ambient temperatures with halocarbons in the presence of other metals. -Produces exothermic reaction with oxygen difluoride. -May initiate explosive polymerisation of olefin oxides including ethylene oxide. -Produces exothermic reaction with oxygen difluoride. -May form explosive mixtures with sodium nitrate. -Reacts vigorously with vinyl acetate. Aluminium oxide is an amphoteric substance, meaning it can react with both acids and bases, such as hydrofluoric acid and sodium hydroxide, acting as an acid with a base and a base with an acid, neutralising the other and producing a salt. Nitric acid: is a strong acid and oxidiser reacts violently with water when added as the concentrated acid with generation of heat (always add acid to water to dilute) reacts violently with water when added as the concentrated acid with generation of heat (always add acid to water to dilute) reacts violently with reducing agents, bases, combustible materials, finely dispersed or powdered metals and proguence diamine, all of horide, armonia, aniline, anioric exchange resins, 1.4-benzoquinone dimine, a.2-bis(trimethylsily)/hydrazine, bromine pentafluoride, cresol, crotonaldehyde, currene, cyanides, diethyl ether, 1,2-dimethyl-2-timethyl-2-timethyl-1-timethyl

Continued...

 presence of light with formation of nitrogen dioxide (which cause discolouration) attacks most metals and some plastics, rubber and coatings
Inorganic acids are generally soluble in water with the release of hydrogen ions. The resulting solutions have pH's of less than 7.0.
Inorganic acids neutralise chemical bases (for example: amines and inorganic hydroxides) to form salts - neutralisation can generate dangerously large amounts of heat in small spaces.
The dissolution of inorganic acids in water or the dilution of their concentrated solutions with additional water may generate significant heat.
The addition of water to inorganic acids often generates sufficient heat in the small region of mixing to cause some of the water to boil explosively. The resulting "bumping" can spatter the acid.
Inorganic acids react with active metals, including such structural metals as aluminum and iron, to release hydrogen, a flammable gas.
▶ Inorganic acids can initiate the polymerisation of certain classes of organic compounds.
Inorganic acids react with cyanide compounds to release gaseous hydrogen cyanide.
Inorganic acids generate flammable and/or toxic gases in contact with dithiocarbamates, isocyanates, mercaptans, nitrides, nitriles, sulfides, and strong reducing agents. Additional gas-generating reactions occur with sulfites, nitrites, thiosulfates (to give H2S and SO3), dithionites (SO2), and even
carbonates.
Acids often catalyse (increase the rate of) chemical reactions.
 WARNING: Avoid or control reaction with peroxides. All transition metal peroxides should be considered as potentially explosive. For example transition metal
complexes of alkyl hydroperoxides may decompose explosively.
 The pi-complexes formed between chromium(0), variatium(0) and other transition metals (haloarene-metal complexes) and mono-or poly-fluorobenzene
show extreme sensitivity to heat and are explosive.
Avoid reaction with borohydrides or cyanoborohydrides
Salts of inorganic fluoride:
 react with water forming acidic solutions.
+ are violent reactive with boron, bromine pentafluoride, bromine trifluoride, calcium disilicide, calcium hydride, oxygen difluoride, platinum, potassium.
 in aqueous solutions are incompatible with sulfuric acid, alkalis, ammonia, aliphatic amines, alkanolamines, alkylene oxides, amides, epichlorohydrin,
isocyanates, nitromethane, organic anhydrides, vinyl acetate.
corrode metals in presence of moisture
▶ may be incompatible with glass and porcelain
Reacts with mild steel, galvanised steel / zinc producing hydrogen gas which may form an explosive mixture with air.
Hydrogen fluoride:
 reacts violently with strong oxidisers, acetic anhydride, alkalis, 2-aminoethanol, arsenic trioxide (with generation of heat), bismuthic acid, calcium oxide,
chlorosulfonic acid, cyanogen fluoride, ethylenediamine, ethyleneimine, fluorine (fluorine gas reacts vigorously with a 50% hydrofluoric acid solution and may
burst into flame), nitrogen trifluoride, N-phenylazopiperidine, oleum, oxygen difluoride, phosphorus pentoxide, potassium permanganate, potassium
tetrafluorosilicate(2-), beta-propiolactone, propylene oxide, sodium, sodium tetrafluorosilicate, sulfuric acid, vinyl acetate
Freats (possibly violently) with aliphatic amines, alcohols, alkanolamines, alkylene oxides, aromatic amines, amides, ammonia, ammonium hydroxide,
epichlorohydrin, isocyanates, metal acetylides, metal silicides, methanesulfonic acid, nitrogen compounds, organic anhydrides, oxides, silicon compounds, vinvlidene fluoride
attacks glass and siliceous materials, concrete, ceramics, metals (flammable hydrogen gas may be produced), metal alloys, some plastics, rubber coatings, leather, and most other metarials with the execution of load platinum potential and up at the execution
leather, and most other materials with the exception of lead, platinum, polyethylene, wax.
Segregate from alkalies, oxidising agents and chemicals readily decomposed by acids, i.e. cyanides, sulfides, carbonates.

SECTION 8 EXPOSURE CONTROLS / PERSONAL PROTECTION

Control parameters

OCCUPATIONAL EXPOSURE LIMITS (OEL)

INGREDIENT DATA

Source	Ingredient	Material name	TWA	STEL	Peak	Notes
US OSHA Permissible Exposure Levels (PELs) - Table Z1	aluminium	Aluminum, metal	15 mg/m3	Not Available	Not Available	Total dust; (as Al)
US OSHA Permissible Exposure Levels (PELs) - Table Z1	aluminium	Aluminum, metal- Respirable fraction	5 mg/m3	Not Available	Not Available	(as Al)
US NIOSH Recommended Exposure Limits (RELs)	aluminium	Aluminium, Aluminum metal, Aluminum powder, Elemental aluminum	10 (total), 5 (resp) mg/m3	Not Available	Not Available	Not Available
US NIOSH Recommended Exposure Limits (RELs)	antimony	Antimony metal, Antimony powder, Stibium	0.5 mg/m3	Not Available	Not Available	[*Note: The REL also applies to other antimony compounds (as Sb).]
US NIOSH Recommended Exposure Limits (RELs)	arsenic	Arsenic metal: Arsenia	Not Available	Not Available	0.002 mg/m3	Ca See Appendix A
US NIOSH Recommended Exposure Limits (RELs)	beryllium	Beryllium metal: Beryllium	Not Available	Not Available	0.0005 mg/m3	Ca See Appendix A
US OSHA Permissible Exposure Levels (PELs) - Table Z1	cadmium	Cadmium	0.005 mg/m3	Not Available	Not Available	see 1910.1027;(as Cd)
US NIOSH Recommended Exposure Limits (RELs)	cadmium	Cadmium metal: Cadmium	0.01 mg/m3	Not Available	Not Available	Ca See Appendix A [*Note: The REL applies to all Cadmium compounds (as Cd).]
US ACGIH Threshold Limit Values (TLV)	cadmium	Cadmium	Not Available	Not Available	Not Available	TLV® Basis: Kidney dam; BEI
US NIOSH Recommended Exposure Limits (RELs)	chromium	Chrome, Chromium	0.5 mg/m3	Not Available	Not Available	Not Available
US OSHA Permissible Exposure Levels (PELs) - Table Z1	cobalt	Cobalt metal, dust, and fume	0.1 mg/m3	Not Available	Not Available	(as Co)
US NIOSH Recommended Exposure Limits (RELs)	cobalt	Cobalt metal dust, Cobalt metal fume	0.05 mg/m3	Not Available	Not Available	TLV® Basis: Pneumonitis
US ACGIH Threshold Limit Values (TLV)	cobalt	Hard metals containing Cobalt and Tungsten carbide, as Co	0.005 mg/m3	Not Available	Not Available	Not Available

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US NIOSH Recommended Exposure Limits (RELs)	copper	Copper metal dusts, Copper metal fumes	1 mg/m3	Not Available	Not Available	[*Note: The REL also applies to other copper compounds (as Cu) except Copper fume.]	
US ACGIH Threshold Limit Values (TLV)	copper	Copper - Fume, as Cu	0.2 mg/m3	Not Available	Not Available	TLV® Basis: Irr; GI; metal fume fever; BEI	
US ACGIH Threshold Limit Values (TLV)	copper	Copper - Dusts and mists, as Cu	1 mg/m3	Not Available	Not Available	TLV® Basis: Irr; GI; metal fume fever BEI	
US NIOSH Recommended Exposure Limits (RELs)	lead	Lead metal, Plumbum	0.050 mg/m3	Not Available	Not Available	See Appendix C [*Note: The REL also applies to other lead compounds (as Pb see Appendix C.]	
US OSHA Permissible Exposure Levels (PELs) - Table Z1	mercury (elemental)	Mercury (vapor)	Hg Vapor: 0.05 mg/m3	Not Available	0.1 mg/m3	See Table Z-2;(as Hg)	
US OSHA Permissible Exposure Levels (PELs) - Table Z2	mercury (elemental)	Mercury	Not Available	Not Available	Other:0.1 mg/m3	(Z37.8–1971)	
US NIOSH Recommended Exposure Limits (RELs)	mercury (elemental)	Mercury metal: Colloidal mercury, Metallic mercury, Quicksilver	Not Available	Not Available	Not Available	Not Available	
US NIOSH Recommended Exposure Limits (RELs)	molybdenum	Molybdenum metal	0.5 mg/m3	Not Available	Not Available	See Appendix D	
US ACGIH Threshold Limit Values (TLV)	molybdenum	Molybdenum, as Mo	Not Available	Not Available	Not Available	TLV® Basis: LRT irr	
US NIOSH Recommended Exposure Limits (RELs)	nickel	Nickel metal: Elemental nickel, Nickel catalyst	0.015 mg/m3	Not Available	Not Available	Ca See Appendix A [*Note: The REL does not apply to Nickel carbonyl.]	
US ACGIH Threshold Limit Values (TLV)	nickel	Nickel and inorganic compounds including Nickel subsulfide, as Ni - Elemental	1.5 mg/m3	Not Available	Not Available	TLV® Basis: Dermatitis; pneumoconiosis	
US NIOSH Recommended Exposure Limits (RELs)	selenium	Elemental selenium, Selenium alloy	0.2 mg/m3	Not Available	Not Available	[*Note: The REL also applies to other selenium compounds (as Se) except Selenium hexafluoride.]	
US NIOSH Recommended Exposure Limits (RELs)	silver	Silver metal: Argentum	0.01 mg/m3	Not Available	Not Available	Not Available	
US OSHA Permissible Exposure Levels (PELs) - Table Z1	nitric acid	Nitric acid	5 mg/m3 / 2 ppm	10 mg/m3 / 4 ppm	Not Available	TLV® Basis: URT & eye irr; dental erosion	
US NIOSH Recommended Exposure Limits (RELs)	nitric acid	Aqua fortis, Engravers acid, Hydrogen nitrate, Red fuming nitric acid (RFNA), White fuming nitric acid (WFNA)	5 mg/m3 / 2 ppm	4 ppm	Not Available	Not Available	
US ACGIH Threshold Limit Values (TLV)	nitric acid	Nitric acid	2 ppm	Not Available	Not Available	Not Available	
US OSHA Permissible Exposure Levels (PELs) - Table Z1	hydrofluoric acid	Hydrogen fluoride	2.5 mg/m3 / 3 ppm	Not Available	5 mg/m3 / 6 ppm	See Table Z-2;(as F)	
JS OSHA Permissible Exposure Levels (PELs) - Table Z2	hydrofluoric acid	Hydrogen fluoride	3 ppm	Not Available	2 ppm	(Z37.28–1969)	
US NIOSH Recommended Exposure Limits (RELs)	hydrofluoric acid	Anhydrous hydrogen fluoride; Aqueous hydrogen fluoride (i.e., Hydrofluoric acid); HF-A	0.5 ppm	Not Available	Not Available	[15-minute]	
US ACGIH Threshold Limit Values (TLV)	hydrofluoric acid	Hydrogen fluoride, as F	Not Available	Not Available	Not Available	TLV® Basis: URT, LRT, skin, & eye irr; fluorosis; BEI	

EMERGENCY LIMITS

Ingredient	Material name	TEEL-1	TEEL-2	TEEL-3
antimony	Antimony	1.5 mg/m3	13 mg/m3	80 mg/m3
barium	Barium	1.5 mg/m3	180 mg/m3	1,100 mg/m3
beryllium	Beryllium	0.0023 mg/m3	Not Available	Not Available
boron	Boron	1.9 mg/m3	21 mg/m3	130 mg/m3
cadmium	Cadmium	Not Available	Not Available	Not Available
chromium	Chromium	1.5 mg/m3	17 mg/m3	99 mg/m3
cobalt	Cobalt	0.18 mg/m3	2 mg/m3	20 mg/m3
copper	Copper	3 mg/m3	33 mg/m3	200 mg/m3
iron	Iron	3.2 mg/m3	35 mg/m3	150 mg/m3
lead	Lead	0.15 mg/m3	120 mg/m3	700 mg/m3
manganese(II) acetate tetrahydrate	Acetic acid, manganese(2+) salt, tetrahydrate	13 mg/m3	22 mg/m3	740 mg/m3
manganese(II) acetate tetrahydrate	Acetic acid, manganese(II) salt (2:1)	9.4 mg/m3	16 mg/m3	96 mg/m3
mercury (elemental)	Mercury vapor	0.15 mg/m3	Not Available	Not Available
molybdenum	Molybdenum	30 mg/m3	330 mg/m3	2,000 mg/m3

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nickel	Nickel		4.5 mg/m3	50 mg/m3	99 mg/m3
selenium	Selenium		0.6 mg/m3	6.6 mg/m3	40 mg/m3
silver	Silver		0.3 mg/m3	170 mg/m3	990 mg/m3
strontium	Strontium		30 mg/m3	330 mg/m3	2,000 mg/m3
thallium	Thallium		0.06 mg/m3	13 mg/m3	20 mg/m3
ammonium metavanadate	Ammonium vanadate; (Ammonium vanadium oxide; Ammonium metavanadate)		0.01 mg/m3	0.11 mg/m3	80 mg/m3
zinc	Zinc		6 mg/m3	21 mg/m3	120 mg/m3
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	Revis	sed IDLH		
aluminium	Not Available	Not A	vailable		
antimony	80 mg/m3	50 mg	j/m3		
arsenic	100 mg/m3	5 mg/	m3		
barium	1,100 mg/m3	50 mg	j/m3		
beryllium	10 mg/m3	4 mg/	4 mg/m3		
boron	Not Available	Not A	Not Available		
cadmium	50 mg/m3 / 9 mg/m3	9 mg/	m3 / 9 [Unch] mg/m3		
chromium	N.E. / N.E.	250 m	250 mg/m3		
cobalt	20 mg/m3	20 [Ui	20 [Unch] mg/m3		
copper	N.E. / N.E.	100 m	100 mg/m3		
iron	Not Available	Not A	Not Available		
lead	700 mg/m3	100 m	ng/m3		
manganese(II) acetate tetrahydrate	N.E. / N.E.	500 m	500 mg/m3		
mercury (elemental)	10 mg/m3 / 28 mg/m3	2 mg/	m3 / 10 mg/m3		
molybdenum	N.E. / N.E.	5,000	mg/m3		
nickel	N.E. / N.E.	10 mg	ŋ/m3		
selenium	Unknown mg/m3 / Unknown ppm	1 mg/	m3		
silver	N.E. / N.E.	10 mg	ŋ/m3		
strontium	Not Available	Not A	vailable		
thallium	Not Available	Not A	vailable		
ammonium metavanadate	Not Available	Not A	vailable		
zinc	Not Available	Not A	vailable		
nitric acid	100 ppm	25 pp	m		
hydrofluoric acid	30 ppm	30 [Ui	nch] ppm		
water	Not Available	Not A	Not Available		

Exposure controls

	Engineering controls are used to remove a hazard or place a barrier between the worker and the hazard. Well-designed engineering controls can be highly effective in protecting workers and will typically be independent of worker interactions to provide this high level of protection. The basic types of engineering controls are: Process controls which involve changing the way a job activity or process is done to reduce the risk. Enclosure and/or isolation of emission source which keeps a selected hazard "physically" away from the worker and ventilation that strategically "adds" and "removes" air in the work environment. 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. Local exhaust ventilation usually required. If risk of overexposure exists, wear approved respirator. Correct fit is essential to obtain adequate protection. Supplied-air type respirator may be required in special circumstances. Correct fit is essential to ensure adequate protection. An approved self contained breathing apparatus (SCBA) may be required in some situations. Provide adequate ventilation in warehouse or closed storage area. 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	Type of Contaminant:		Air Speed:		
controls	solvent, vapours, degreasing etc., evaporating from tank (in still air).		0.25-0.5 m/s (50-100 f/min.)		
	aerosols, fumes from pouring operations, intermittent container filling, low speed conveyer transfe acid fumes, pickling (released at low velocity into zone of active generation)	rs, welding, spray drift, plating	0.5-1 m/s (100-200 f/min.)		
	direct spray, spray painting in shallow booths, drum filling, conveyer loading, crusher dusts, gas discharge (active generation into zone of rapid air motion)		1-2.5 m/s (200-500 f/min.)		
	grinding, abrasive blasting, tumbling, high speed wheel generated dusts (released at high initial ve air motion).	elocity into zone of very high rapid	2.5-10 m/s (500-2000 f/min.)		
	Within each range the appropriate value depends on:				
	Lower end of the range	Upper 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.	2: Contaminants of high toxicity				
	3: Intermittent, low production.	3: High production, heavy use				
	4: Large hood or large air mass in motion	4: Small hood-local control only				
	of distance from the extraction point (in simple cases). Therefore the air speed at the distance from the contaminating source. The air velocity at the extraction fan, for exam solvents generated in a tank 2 meters distant from the extraction point. Other mechan	air velocity falls rapidly with distance away from the opening of a simple extraction pipe. Velocity generally decreases with the square action point (in simple cases). Therefore the air speed at the extraction point should be adjusted, accordingly, after reference to inating 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 ank 2 meters distant from the extraction point. Other mechanical considerations, producing performance deficits within the extraction tial 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	 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 lenses 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 Intelling the source shullen the source shull be sources and the store shull be sources and the store source shullen the source source sources and the store source sources are stored as 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 Intelling the sources are shullen to the store of the source source of the source source of the of the source of the source of the source of the source source of the source of					
	Current Intelligence Bulletin 59], [AS/NZS 1336 or national equivalent]	nvironment only after workers have washed hands thoroughly. [CDC NIOSF				
Skin protection	See Hand protection below	nvironment only after workers have washed hands thoroughly. [CDC NIOSF				
Skin protection Hands/feet protection						
· · ·	See Hand protection below Wear chemical protective gloves, e.g. PVC. Wear safety footwear or safety gumboots, e.g. Rubber					
Hands/feet protection	See Hand protection below Wear chemical protective gloves, e.g. PVC. Wear safety footwear or safety gumboots, e.g. Rubber When handling corrosive liquids, wear trousers or overalls outside of boots, to a					

Respiratory protection

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

SECTION 9 PHYSICAL AND CHEMICAL PROPERTIES

Information on basic physical and chemical properties

Appearance	Colourless		
Physical state	Liquid	Relative density (Water = 1)	Not Available
Odour	Not Available	Partition coefficient n-octanol / water	Not Available
Odour threshold	Not Available	Auto-ignition temperature (°C)	Not Available
pH (as supplied)	Not Available	Decomposition temperature	Not Available
Melting point / freezing point (°C)	Not Available	Viscosity (cSt)	Not Available
Initial boiling point and boiling range (°C)	Not Available	Molecular weight (g/mol)	Not Available
Flash point (°C)	Not Available	Taste	Not Available
Evaporation rate	Not Available	Explosive properties	Not Available
Flammability	Not Available	Oxidising properties	Not Available
Upper Explosive Limit (%)	Not Available	Surface Tension (dyn/cm or mN/m)	Not Available
Lower Explosive Limit (%)	Not Available	Volatile Component (%vol)	Not Available
Vapour pressure (kPa)	Not Available	Gas group	Not Available
Solubility in water (g/L)	Miscible	pH as a solution (1%)	Not Available
Vapour density (Air = 1)	Not Available	VOC g/L	Not Available

SECTION 10 STABILITY AND REACTIVITY

Reactivity	See section 7
Chemical stability	Contact with alkaline material liberates heat
Possibility of hazardous reactions	See section 7

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Conditions to avoid	See section 7
Incompatible materials	See section 7
Hazardous decomposition products	See section 5
SECTION 11 TOXICOLOG	SICAL INFORMATION
Information on toxicologic	cal effects
Inhaled	Inhalation of vapours or aerosols (mists, fumes), generated by the material during the course of normal handling, may be harmful. 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. Bronchial and alveolar exudate are apparent in animals exposed to molybdenum by inhalation. Molybdenum fume may produce bronchial irritation and moderate fatty changes in liver and kidney. Inhalation of nitric acid mist or fumes may produce respiratory symptoms. Depending on the concentration and duration of exposure, cough, gagging, chest pain, low body oxygen, lung irritation and damage may occur. Deaths have occurred and may be delayed for several days. 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 irritation 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 levels 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 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
Ingestion	Accidental ingestion of the material may be harmful; animal experiments indicate that ingestion of less than 150 gram may be fatal or may produce serious damage to the health of the individual. The material can produce chemical burns within the oral cavity and gastrointestinal tract following ingestion. Molybdenum, an essential trace element, can in large doses hamper growth and cause loss of appetite, listlessness and diarrhoea. Anaemia also occurs, and other symptoms include greying of hair, shrinking of the testicles, reduced fertility and milk production, shortness of breath, inccordination and irritation of the mucous membranes. Ingestion of acidic corrosives may produce burns around and in the mouth, the throat and oesophagus. Immediate pain and difficulties in swallowing and speaking may also be evident. 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 stomach lining and subsequent pemicious anaemia. 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.
Skin Contact	Skin contact with the material may be harmful; systemic effects may result following absorption. The material can produce chemical burns following direct contact with the skin. Though considered non-harmful, slight irritation may result from contact because of the abrasive nature of the aluminium oxide particles. Thus it may cause itching and skin reaction and inflammation. 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. Skin contact with acidic corrosives may result in pain and burns; these may be deep with distinct edges and may heal slowly with the formation of scar tissue.

Skin contact with nitric acid may cause corrosion, skin thickening, yellow discolouration of the skin, blisters and scars depending on the concentration exposed. 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 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. The material can produce chemical burns to the eye following direct contact. Vapours or mists may be extremely irritating. If applied to the eyes, this material causes severe eye damage

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

Eye Eye contact with both 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.

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

Long-term exposure to respiratory irritants may result in airways disease, involving difficulty breathing and related whole-body problems.

Substance accumulation, in the human body, may occur and may cause some concern following repeated or long-term occupational exposure.

Animal testing shows long term exposure to aluminium oxides may cause lung disease and cancer, depending on the size of the particle. The smaller the size, the greater the tendencies of causing harm.

High levels of molybdenum can cause joint problems in the hands and feet with pain and lameness. Molybdenum compounds can also cause liver changes with elevated levels of enzymes and cause over-activity of the thyroid gland.

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

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

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. Hydrogen fluoride easily penetrates the skin and causes destruction and corrosion of the bone and underlying tissue. Ingestion causes severe pains and burns

cww-Waste W

TM-C - Certified ater Trace Metals	TOXICITY	IRRITATION
Solution C	Not Available	Not Available

in the mouth and throat and blood calcium levels are dangerously reduced.

	ΤΟΧΙΟΙΤΥ			IRR	ITATION
aluminium	Oral (rat) LD50: >2000 mg/kg ^[1] No				Available
	ΤΟΧΙΟΙΤΥ				IRRITATION
antimony	Dermal (rabbit) LD50: >8300 mg/kg ^[1]				Not Available
	Oral (rat) LD50: 100 mg/kg ^[2]				
	ΤΟΧΙΟΙΤΥ			IRRIT	ATION
arsenic	Oral (rat) LD50: 763 mg/kg ^[2]			Not Av	
	ΤΟΧΙΟΙΤΥ		IRRITATION		
barium	Not Available		Not Available		
			1		
	ΤΟΧΙΟΙΤΥ			IRR	ITATION
beryllium	Oral (rat) LD50: >2000 mg/kg ^[1]			Not	Available
	ΤΟΧΙΟΙΤΥ			IRRIT/	ATION
boron	Oral (rat) LD50: 650 mg/kg ^[2]			Not Av	vailable
	ΤΟΧΙΟΙΤΥ				IRRITATION
cadmium	Oral (rat) LD50: >63<259 mg/kg> ^[1]				Not Available
	ΤΟΧΙΟΙΤΥ		IRRITATION		
chromium	Not Available		Not Available		
	ΤΟΧΙΟΙΤΥ			IF	RITATION
cobalt	dermal (rat) LD50: >2000 mg/kg ^[1]				lot Available
	Oral (rat) LD50: 6170 mg/kgd ^[2]				
	ΤΟΧΙΟΙΤΥ				IRRITATION
	dermal (rat) LD50: >2000 mg/kg ^[1]				Not Available
	Inhalation (rat) LC50: 0.733 mg/l/4hr ^[1]				
copper	Inhalation (rat) LC50: 1.03 mg//4hr ^[1]				
	Inhalation (rat) LC50: 1.67 mg//4hr ^[1]				
	Oral (rat) LD50: 300-500 mg/kg ^[1]				
					1
	ΤΟΧΙΟΙΤΥ			IRR	RITATION
iron	Oral (rat) LD50: 98600 mg/kg] ^[2]			Not	Available
	ΤΟΧΙΟΙΤΥ				IRRITATION
	dermal (rat) LD50: >2000 mg/kg ^[1]				Not Available
lead	Inhalation (rat) LC50: >5.05 mg/l/4hr ^[1]				
	Oral (rat) LD50: >2000 mg/kg ^[1]				
	ΤΟΧΙΟΙΤΥ			IPP	ITATION
manganese(II) acetate tetrahydrate	Oral (rat) LD50: 3730 mg/kga ^[2]				Available
	ΤΟΧΙΟΙΤΥ			IPPIT	TATION
mercury (elemental)					ration
	Oral (rat) LD50: >9.2 mg/kg ^[1]			NULA	

	L						
	TOXICITY			IRRITATION			
molybdenum	dermal (rat) LD50: >2000 mg/kg ^[1]		Not Available				
	Oral (rat) LD50: >2000 mg/kg ^[1]						
	ΤΟΧΙΟΙΤΥ			IRRITATION			
nickel	Oral (rat) LD50: 5000 mg/kg ^[2]			Not Available			
	TOXICITY			IRRITATION			
selenium	Oral (rat) LD50: 6700 mg/kgd ^[2]			Not Available			
	ΤΟΧΙΟΙΤΥ			IRRITATION			
silver	Oral (rat) LD50: >2000 mg/kg ^[1]			Not Available			
strontium	TOXICITY	IRF	RITATION				
strontium	Not Available	Not	t Available				
	ΤΟΧΙΟΙΤΥ	IRE	RITATION				
thallium	Not Available	Not Available					
	TOXICITY			IRRITATION			
ammonium metavanadate	dermal (rat) LD50: 2102 mg/kg ^[2]	dermal (rat) LD50: 2102 mg/kg ^[2]					
	Oral (rat) LD50: 160 mg/kgd ^[2]						
	ΤΟΧΙΟΙΤΥ			IRRITATION			
zinc	Dermal (rabbit) LD50: 1130 mg/kg ^[2]			Not Available			
	Oral (rat) LD50: >2000 mg/kg ^[1]						
nitric acid	ΤΟΧΙΟΙΤΥ			IRRITATION			
	Inhalation (rat) LC50: 625 ppm/1h*t ^[2]			Not Available			
	ΤΟΧΙΟΙΤΥ		IRRITATION				
hydrofluoric acid	Inhalation (rat) LC50: 1276 ppm/4hr ^[2]		Eye (human): 50 mg	- SEVERE			
·	Inhalation (rat) LC50: 319 ppm/1hr ^[2]						
	ΤΟΧΙΟΙΤΥ						
water		RITATION t Available					
	Not Available		t Avallable				
Legend:	1. Value obtained from Europe ECHA Registered Substances extracted from RTECS - Register of Toxic Effect of chemical S		lue obtained from manufa	acturer's SDS. Unless otherwise specified data			
ARSENIC	Arsenic compounds are classified by the European Union as to Tumorigenic - Carcinogenic by RTECS criteria.	oxic by inhalation and	ingestion and toxic to aq	uatic life and long lasting in the environment.			
	33nix&11b						
	WARNING: Beryllium and compounds are classified by IARC	as					

BERYLLIUM	33nix&11b WARNING: Beryllium and compounds are classified by IARC as Group 1- CARCINOGENIC TO HUMANS Beryllium oxide fume is very toxic to the respiratory tract, lungs and skin and is quick acting. Mutation DNA damage Human Tumorigenic - neoplastic by RTECS criteria.
BORON	Elemental boron produces lower foetal body weight in rats.
CHROMIUM	On skin and inhalation exposure, chromium and its compounds (except hexavalent) can be a potent sensitiser, as particulates. Gastrointestinal tumours, lymphoma, musculoskeletal tumours and tumours at site of application recorded.
COBALT	Allergic reactions involving the respiratory tract are usually due to interactions between IgE antibodies and allergens and occur rapidly. Attention should be paid to atopic diathesis, characterised by increased susceptibility to nasal inflammation, asthma and eczema. Exogenous allergic alveolitis is induced essentially by allergen specific immune-complexes of the IgG type; cell-mediated reactions (T lymphocytes) may be involved.

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COPPER	for copper and its compounds (typically copper chloride): Acute toxicity: There are no reliable acute oral toxicity results available. WARNING: Inhalation of high concentrations of copper fume may cause "metal fume fever", a like respiratory tract irritation with fever.	n acute industrial disease of short duration. tiredness, influenza						
LEAD	WARNING: Lead is a cumulative poison and has the potential to cause abortion and intellectua	al impairment to unborn children of pregnant workers.						
MERCURY (ELEMENTAL)	Animal studies have shown that mercury may be a reproductive effector.							
NICKEL	enth Annual Report on Carcinogens: Substance anticipated to be Carcinogen National Toxicology Program: U.S. Dep. Oral (rat) TDLo: 500 mg/kg/5D-I Inhalation (rat) TCLo: 0.1 mg/m3/24H/17W-C							
THALLIUM	Structural changes in nerves and sheath, changes in extraocular muscles, hair loss recorded							
ZINC	The material may cause skin irritation after prolonged or repeated exposure and may produce scaling and thickening of the skin.	he material may cause skin irritation after prolonged or repeated exposure and may produce on contact skin redness, swelling, the production of vesicles, caling and thickening of the skin.						
NITRIC ACID	For acid mists, aerosols, vapours Test results suggest that eukaryotic cells are susceptible to genetic damage when the pH falls to about 6.5. 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. Oral (?) LD50: 50-500 mg/kg * [Various Manufacturers]							
HYDROFLUORIC ACID	(liver and kidney damage) [Manufacturer] for hydrogen fluoride (as vapour)							
ALUMINIUM & BARIUM & CHROMIUM & MOLYBDENUM & STRONTIUM & HYDROFLUORIC ACID & WATER	No significant acute toxicological data identified in literature search.							
ARSENIC & BERYLLIUM	WARNING: This substance has been classified by the IARC as Group 1: CARCINOGENIC TO HUMANS.							
BARIUM & BORON & MANGANESE(II) ACETATE TETRAHYDRATE & MERCURY (ELEMENTAL) & AMMONIUM METAVANADATE & NITRIC ACID & HYDROFLUORIC ACID	Asthma-like symptoms may continue for months or even years after exposure to the material en	ds.						
BERYLLIUM & COBALT & NICKEL	The following information refers to contact allergens as a group and may not be specific to this product.							
BERYLLIUM & CHROMIUM	Tenth Annual Report on Carcinogens: Substance known to be Carcinogenic [National Toxicology Program: U.S. Dep.							
CHROMIUM & SELENIUM	The substance is classified by IARC as Group 3: NOT classifiable as to its carcinogenicity to humans.							
COBALT & NICKEL	WARNING: This substance has been classified by the IARC as Group 2B: Possibly Carcino	genic to Humans.						
NITRIC ACID & HYDROFLUORIC ACID	The material may produce severe irritation to the eye causing pronounced inflammation.							
NITRIC ACID & HYDROFLUORIC ACID	The material may produce respiratory tract irritation, and result in damage to the lung includin	g reduced lung function.						
Acute Toxicity	✓ Carcinogenicit	у 🛇						
Skin Irritation/Corrosion	✓ Reproductivit							
Serious Eye Damage/Irritation	✓ STOT - Single Exposur							
Respiratory or Skin sensitisation	STOT - Repeated Exposur	• 🛇						
Mutagenicity	S Aspiration Hazar	a 🛇						
	Legend:	 Data available but does not fill the criteria for classification Data available to make classification 						

- Data available to make classification
- S Data Not Available to make classification

SECTION 12 ECOLOGICAL INFORMATION

CWW-TM-C - Certified	ENDPOINT	TEST DURATION (HR)		SPECIES		E	SOURCE	
Waste Water Trace Metals Solution C	Not Applicable	able Not Applicable		Not Applicable	Not Ap	plicable	Not Applicable	
	ENDPOINT	TEST DURATION (HR)	SPECIE	SPECIES		VALUE		SOURCE
	LC50	96	Fish	Fish				2
aluminium	EC50	48	Crustac	Crustacea				2
	EC50	96	Algae o	Algae or other aquatic plants				2
	BCF	360	Algae o	Algae or other aquatic plants				4

	EC50	120		Fis	sh		0.000)051mg/L	5
	NOEC	72			gae or other aqua	atic plants		004mg/L	2
	NOLO	12			gae of other aqua		0.	oo-ing/L	2
	ENDPOINT	TES	T DURATION (HR)		SPECIES		١	/ALUE	SOURCE
	LC50	96			Fish).93mg/L	2
	EC50	48			Crustacea			mg/L	2
antimony	EC50	72				auatic plants			2
	EC50				Algae or other ad	qualic plants		>2.4mg/L	2
		96			Crustacea).5mg/L	
	NOEC	720			Fish		>	>0.0075mg/L	2
	ENDPOINT	TES	ST DURATION (HR)		SPECIES			VALUE	SOURCE
	LC50	96			Fish			9.9mg/L	4
arsenic	EC50	336			Algae or other	aquatic plants		0.63mg/L	4
	NOEC	336			Algae or other			<0.75mg/L	4
	NOLC	550			Aigae of other a	aqualle plants		<0.7 Shig/L	4
	ENDPOINT	TEO			SPECIES		V	ALUE	SOURCE
			T DURATION (HR)						
	LC50	96			Fish			500mg/L	4
barium	EC50	96			Algae or other aq	uatic plants		Smg/L	4
	BCF	24			Crustacea			000002mg/L	4
	EC50	240			Algae or other aq	uatic plants		10306mg/L	4
	NOEC	48		(Crustacea		68	3mg/L	4
	ENDPOINT		TEST DURATION (HR)			SPECIES	VALU		SOURCE
beryllium	EC50	96			Fish		0.19mg/L		5
	NOEC		48			Crustacea	0.25n	ng/L	4
					1				
	ENDPOINT		ST DURATION (HR)		SPECIES			VALUE	SOURCE
	LC50	96			Fish			74mg/L	2
	EC50	48			Crustacea			230mg/L	5
boron	EC50	72			Algae or other	aquatic plants		54mg/L	2
	BCF	336			Algae or other	aquatic plants		8.5mg/L	4
	EC50	336			Algae or other	aquatic plants		8.5mg/L	4
	NOEC	576			Fish			0.001mg/L	5
	ENDPOINT	TES	T DURATION (HR)	SI	PECIES		VAL	UE	SOURCE
	LC50	96		Fi	ish		0.00	1mg/L	4
	EC50	48		С	rustacea		0.00	33mg/L	5
cadmium	EC50	72		AI	lgae or other aqu	atic plants	0.01	8mg/L	2
	BCF	960		Fi	ish		500r	ng/L	4
	EC50	336		C	rustacea		0.00	065mg/L	5
	NOEC	168		Fi	ish		0.00	001821mg/L	4
	ENDPOINT	TES	T DURATION (HR)		SPECIES		1	VALUE	SOURCE
	LC50	96			Fish			13.9mg/L	4
	EC50	48			Crustacea		().0225mg/L	5
chromium	EC50	72			Algae or other aquatic plants		().104mg/L	4
	BCF	1440)		Algae or other a	quatic plants	().0495mg/L	4
	EC50	48			Crustacea		().0245mg/L	5
	NOEC	672			Fish		(0.00019mg/L	4
	ENDPOINT	TES	T DURATION (HR)		SPECIES			VALUE	SOURCE
	LC50	96			Fish			1.406mg/L	2
		48			Crustacea			>0.89mg/L	2
cobalt	EC50	40			Crustacea Algae or other aquatic plants				
cobalt	EC50 EC50	40 72			Algae or other a	equatic plants		0.144mg/L	2
cobalt			4		Algae or other a Fish	aquatic plants		0.144mg/L 0.99mg/L	2 4

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Control Sign Pain Control Con		ENDPOINT	TE	ST DURATION (HR)	SPEC	FS		VALUE	SOURCE
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EC50 48 Crustacea 0.013mg/L 5 EC50 72 Algae or other aquatic plants 0.0407mg/L 2 BCF 1440 Algae or other aquatic plants 0.47mg/L 4 EC50 720 Crustacea 0.0062mg/L 2 NOEC 72 Algae or other aquatic plants 0.0035mg/L 2 NOEC 72 Algae or other aquatic plants 0.0035mg/L 2 NOEC 72 Algae or other aquatic plants 0.0035mg/L 2 VENE VENE VENE 2 2 NOEC 72 Algae or other aquatic plants 0.0035mg/L 2 VENE VENE VENE SOURCE 2 LC50 96 Fish >0.0262mg/L 2 EC50 48 Crustacea >0.1603mg/L 2 EC50 72 Algae or other aquatic plants >0.00173mg/L 2				DURATION (MR)					
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BCF 1440 Algae or other aquatic plants 0.47mg/L 4 EC50 720 Crustacea 0.0062mg/L 2 NOEC 72 Algae or other aquatic plants 0.0035mg/L 2 VALUE VALUE SOURCE LC50 96 Fish >0.0262mg/L 2 LC50 96 Fish >0.0262mg/L 2 EC50 48 Crustacea >0.1603mg/L 2 EC50 72 Algae or other aquatic plants >0.00173mg/L 2			48		Crusta	Crustacea		.013mg/L	5
EC50 720 Crustacea 0.0062mg/L 2 NOEC 72 Algae or other aquatic plants 0.0035mg/L 2 selenium ENDPOINT TEST DURATION (HR) SPECIES VALUE SOURCE LC50 96 Fish >0.0262mg/L 2 EC50 48 Crustacea >0.1603mg/L 2 EC50 72 Algae or other aquatic plants >0.00262mg/L 2		EC50				Algae or other aquatic plants			-
NOEC 72 Algae or other aquatic plants 0.0035mg/L 2 Image: selenium ENDPOINT TEST DURATION (HR) SPECIES VALUE SOURCE LC50 96 Fish >0.0262mg/L 2 EC50 48 Crustacea >0.1603mg/L 2 EC50 72 Algae or other aquatic plants >0.00173mg/L 2	nickel	EC50 EC50	72		Algae			-	
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keenium LC50 96 Fish >0.0262mg/L 2 EC50 48 Crustacea >0.1603mg/L 2 EC50 72 Algae or other aquatic plants >0.00173mg/L 2	nickel	EC50 EC50 BCF	72 144 720		Algae o	or other aquatic plants	0	.47mg/L	4 2
kelenium LC50 96 Fish >0.0262mg/L 2 EC50 48 Crustacea >0.1603mg/L 2 EC50 72 Algae or other aquatic plants >0.00173mg/L 2	nickel	EC50 EC50 BCF EC50	72 144 720		Algae o Algae o Crusta	or other aquatic plants cea	0	.47mg/L .0062mg/L	4 2
selenium EC50 48 Crustacea >0.1603mg/L 2 EC50 72 Algae or other aquatic plants >0.00173mg/L 2	nickel	EC50 EC50 BCF EC50 NOEC	72 144 720 72		Algae o Algae o Crusta Algae o	or other aquatic plants cea or other aquatic plants	0	.47mg/L .0062mg/L .0035mg/L	4 2 2
EC50 72 Algae or other aquatic plants >0.00173mg/L 2	nickel	EC50 EC50 BCF EC50 NOEC	72 144 720 72 TES		Algae o Algae o Crusta Algae o SPEC	or other aquatic plants cea or other aquatic plants	000000000000000000000000000000000000000	.47mg/L .0062mg/L .0035mg/L VALUE	4 2 2
		EC50 EC50 BCF EC50 NOEC ENDPOINT LC50	72 144 720 72 72 72 72		Algae of Algae of Algae of Crustal Algae of SPEC	or other aquatic plants cea or other aquatic plants ES		.47mg/L .0062mg/L .0035mg/L VALUE >0.0262mg/L	4 2 2 2 SOURCE 2
		EC50 EC50 BCF EC50 NOEC ENDPOINT LC50 EC50	72 144 720 72 72 72 72 72 72 72 72 72 72		Algae of Algae of Crustal Algae of SPEC	or other aquatic plants cea or other aquatic plants ES cea		47mg/L .0062mg/L .0035mg/L VALUE >0.0262mg/L >0.1603mg/L	4 2 2 2 SOURCE 2 2 2

	EC50	96			or other aqua			0.355mg/L		
	NOEC	72		Algae	or other aquat	ic plants		0.000547n	ng/L	2
	ENDPOINT	TES	T DURATION (HR)	SPECIE	S			VALUE		SOURCE
	LC50	96		Fish				0.00148mg/L		2
	EC50	48		Crustace	ea			0.00024mg/L		4
silver	EC50	96		Algae or	other aquatic	plants		0.001628837r	ng/L	4
	BCF	336		Crustace	ea			0.02mg/L		4
	EC50	48		Crustace	ea			0.00024mg/L		4
	NOEC	480		Crustace	ea			0.00031mg/L		2
	ENDPOINT		TEST DURATION (HR)		SPECIES		VALU	E	SO	URCE
strontium	Not Applicable		Not Applicable		Not Applica	ble	_	oplicable		t Applicable
			riot / ppilocolo		- tot, pp.ice		1.0074	spiloubio		
	ENDPOINT	TES	T DURATION (HR)	SPEC	IES			VALUE		SOURCE
	LC50	96	\ /	Fish				21mg/L		4
thallium	EC50	96			or other aqua	tic plants		0.13mg/L		4
	EC50	240		-	or other aqua	-		0.040876	ma/l	4
	NOEC	720		Fish	or ourier aqua			0.04mg/L	11g/L	5
	NOLO	120		11311				0.04Mg/L		3
	ENDPOINT	TES	ST DURATION (HR)	SPE	CIES			VALUE		SOURCE
	LC50	96		Fish				0.693m	0.693mg/L	
	EC50	48		Crus	tacea			2.387m	ng/L	2
monium metavanadate	EC50	72		Alga	e or other aqu	atic plants		0.9894	mg/L	2
	EC50	72			e or other aqu			1.162m	-	2
	NOEC	72			e or other aqu			0.0168	-	2
	ENDPOINT	TES	ST DURATION (HR)	SPEC	CIES			VALUE		SOURCE
	LC50	96		Fish				0.00272	mg/L	4
	EC50	48		Crust	acea			0.04mg/	L	5
zinc	EC50	72		Algae	or other aqua	atic plants		0.106mg	ı/L	4
	BCF	360		Algae	or other aqua	atic plants		9mg/L		4
	EC50	120		Fish				0.00033	mg/L	5
	NOEC	336		Algae	or other aqua	atic plants		0.00075	mg/L	4
nitric acid	ENDPOINT		TEST DURATION (HR)			SPECIES		VALUE		SOURCE
	NOEC		16			Crustacea		107mg/L	2	4
	ENDPOINT		TEST DURATION (HR)		SF	ECIES	,	VALUE		SOURCE
	LC50		96		Fis			51mg/L		2
hydrofluoric acid	EC50		48			ustacea		=270mg/L		1
nyuronuone aciu										
	EC50 NOEC		96 504		Fis	ustacea		26-48mg/L 4mg/L		2 2
	NOEU		JUT			211	·	±ing/∟		۷
	ENDPOINT		TEST DURATION (HR)		SPECIES		VALU	E	SO	URCE
water	Not Applicable		Not Applicable		Not Applica	ble	Not A	oplicable	Not	t Applicable
					I INUL AUUIUd		I NUL A	pillane		L APPIICADIE

(QSAR) - Aquatic Toxicity Data (Estimated) 4. US EPA, Ecotox database - Aquatic Toxicity Data 5. ECETOC Aquatic Hazard Assessment Data 6. NITE (Japan) - Bioconcentration Data 7. METI (Japan) - Bioconcentration Data 8. Vendor Data

For Molybdenum:

Environmental Fate: Molybdenum is an essential micronutrient in plants and animals. It is commonly used in the manufacture of steel alloys. Based on the high concentration of molybdenum in all analyzed waste types, the exposure of the environment to molybdenum is regarded as significant. The limited amount of data regarding its toxicity makes it impossible to evaluate the potential for adverse environmental and health effects from molybdenum exposure. Molybdenum is generally found in two oxidation states in nature, Mo(IV) and Mo(VI). In oxidizing environments, Mo(VI) dominates and it is commonly present as molybdenum contains seven isotopes. Molybdenum oxidizes at elevated temperatures.

Atmospheric Fate: Molybdenum can be deposited via dry/wet deposition; however, atmospheric exposure has been identified as a minor source to terrestrial and aquatic habitats. Terrestrial Fate: Molybdenum is a naturally occurring substance in soil. Soil molybdenum is a potentially toxic element, but no cases have been reported of molybdenum toxicity to animals from consumption of forage grown on sludge-amended soils. Microbes are expected to transform the substance.

Aquatic Fate: Molybdenum disulfide is sparingly soluble in water but oxidizes to more soluble molybdates, which are stable in water. At pH 3-5, molybdate frequently shifts to hydrogen molybdate.

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Low pH molybdenum is usually adsorbed to sediment composed of clay, or other minerals that are prone to weathering. Molybdenum in the water is expected to be taken up by aquatic organisms. Concentrations of the substance in sediments are by site-specific factors like flow rate, and other factors, (e.g. organic content, pH)

Ecotoxicology: Molybdenum cause adverse effects in ruminant animals. Livestock have been injured by forage grown on soils containing the element. The substance stoxicological properties in mammals are governed, to a large extent, by its interaction with copper and sulfur; residues of molybdenum alone are not sufficient to diagnose poisoning by the substance. Domestic ruminants, especially cattle, are especially sensitive to molybdenum poisoning, when copper and inorganic sulfate are deficient. The resistance of small laboratory animals, and wildlife, is at least 10X that of cattle. Mule deer are not adversely affected by the substance. The substance may have a negative impact on reproduction in domestic birds and there is inadequate data on its effects on waterfowl and most mammals.

For Vanadium Compounds:

Environmental Fate: Vanadium is travels through the environment via long-range transportation in the atmosphere, water, and land by natural and man-made sources, wet and dry deposition, adsorption and complexing. From natural sources, vanadium is probably in the form of less soluble trivalent mineral particles.

Atmospheric Fate: Vanadium generally enters the atmosphere as an aerosol. Natural and man-made sources of vanadium tend to release large particles that are more likely to settle near the source. Smaller particles, such as those emitted from oil-fueled power plants, have a longer residence time in the atmosphere and are more likely to be transported farther away from the site of release.

Terrestrial Fate: Soil - Transport and partitioning of vanadium in soil is influenced by pH and reduction potential. Ferric hydroxides and solid bitumens (organic) are the main carriers of vanadium in the sedimentation process. Iron acts as a carrier for trivalent vanadium and is responsible for its diffusion through molten rocks where it becomes trapped during crystallization. Vanadium is fairly mobile in neutral or alkaline soils, but its mobility decreases in acidic soils. Under oxidizing, unsaturated conditions, some mobility is observed, but under reducing, saturated conditions, vanadium is immobile. Plants - Vanadium levels in terrestrial plants are dependent upon the amount of water-soluble vanadium available in the soil as well as pH and growing conditions. The uptake of vanadium into the above-ground parts of many plants is low, although root concentrations have shown some correlation with levels in the soil. Certain legumes have been shown to be vanadium accumulators and the root nodules of these plants may contain vanadium levels three times greater than those of the surrounding soil. Fly agaric (Amanita muscaria) mushrooms are known to actively accumulate vanadium.

Aquatic Fate: Vanadium is eventually adsorbed to hydroxides or associated with organic compounds and is deposited on the sea bed. Vanadium is transported in water by solution (13%) or suspension (87%). Upon entering the oceans, vanadium is deposited to the sea bed. Only about 0.001% of vanadium entering the oceans is estimated to persist in soluble form. Sorption and biochemical processes are thought to contribute to the extraction of vanadium from sea water. Adsorption to organic matter as well as to manganese oxide and ferric hydroxide results in the precipitation of dissolved vanadium. Biochemical processes are also of importance in the partitioning from sea water to sediment.

Ecotoxicity: Some marine organisms, in particular the sea squirts, bioconcentrate vanadium very efficiently, attaining body concentrations approximately 10,000 times greater than the ambient sea water. Upon the death of the organism, the body burden adds to the accumulation of vanadium in silt. In general, marine plants and invertebrates contain higher levels of vanadium than terrestrial plants and animals. In the terrestrial environment, bioconcentration is more commonly observed amongst the lower plant phyla than in the higher, seed-producing phyla. Vanadium appears to be present in all terrestrial animals; however tissue concentrations in vertebrates are often so low that detection is difficult. The highest levels of vanadium in terrestrial mammals are generally found in the liver and skeletal tissues. No data are available regarding biomagnification of vanadium within the food chain, but human studies suggest that it is unlikely. Bioaccumulation appears to be unlikely.

Ecotoxicity:

The tolerance of water organisms towards pH margin and variation is diverse. Recommended pH values for test species listed in OECD guidelines are between 6.0 and almost 9. Acute testing with fish showed 96h-LC50 at about pH 3.5

For Fluorides: Small amounts of fluoride have beneficial effects however; excessive intake over long periods may cause dental and/or skeletal fluorosis. Fluorides are absorbed by humans following inhalation of workplace and ambient air that has been contaminated, ingestion of drinking water and foods and dermal contact. Populations living in areas with high fluoride levels in groundwater may be exposed to higher levels of fluorides in their drinking water or in beverages prepared with the water. Among these populations, outdoor labourers, people living in hot climates, and people with excessive thirst will generally have the greatest daily intake of fluorides because they consume greater amounts of water.

Atmospheric Fate: Both hydrogen fluoride and particulate fluorides will be transported in the atmosphere and deposited on land or water by wet and dry deposition. Non-volatile inorganic fluoride particulates are removed from the atmosphere via condensation or nucleation processes. Fluorides adsorbed on particulate matter in the atmosphere are generally stable and are not readily hydrolyzed, although they may be degraded by radiation if they persist in the atmosphere. Fluorine and the silicon fluorides (fluosilicates, silicofluorides) are hydrolyzed in the atmosphere to form hydrogen fluoride. Hydrogen fluoride may combine with water vapour to produce an aerosol or fog of aqueous hydrofluoric acid. Inorganic fluoride compounds, with the exception of sulfur hexafluoride, are not expected to remain in the troposphere for long periods or to migrate to the stratosphere. Estimates of the residence time of sulfur hexafluoride in aerosols can be transported over large distances by wind or as a result of atmospheric turbulence. Fluorosilicic acid and hydrofluoric acid in high aquatic concentrations such as may be found in industrial waste ponds may volatilize, releasing silicon tetrafluoride and hydrogen fluoride into the atmosphere. Soluble inorganic fluorides may also form aerosols at the air/water interface or vaporize into the atmosphere whereas undissolved species generally undergo sedimentation.

Terrestrial Fate: Soils - Atmospheric fluorides may be transported to soils and surface waters through both wet and dry deposition processes where they may form complexes and bind strongly to soil and sediment. Solubilisation of inorganic fluorides from minerals may also be enhanced by the presence of bentonite clays and humic acid. Factors that influence the mobility of inorganic fluorides in soil are pH and the formation of aluminium and calcium complexes. In more acidic soils, concentrations of inorganic fluoride were considerably higher in the deeper horizons. The low affinity of fluorides for organic material results in leaching from the more acidic surface horizon and increased retention by clay minerals and silts in the more alkaline, deeper horizons. The maximum adsorption of fluoride to soil was reported to occur at pH 5.5. In acidic soils with pH below 6, most of the fluoride is in complexes with either aluminium or iron. Fluoride in alkaline soils at pH 6.5 and above is almost completely fixed in soils as calcium fluoride, if sufficient calcium carbonate is available. Fluoride is extremely immobile in soil.

Aquatic Fate: Fresh Water: - In water, the transport and transformation of inorganic fluorides are influenced by pH, water hardness and the presence of ion-exchange materials such as clays. In natural water, fluoride forms strong complexes with aluminium in water, and fluorine chemistry in water is largely regulated by aluminium concentration and pH. Below pH 5, fluoride is almost entirely complexed with aluminium and consequently, the concentration of free F- is low. Once dissolved, inorganic fluorides remain in solution under conditions of low pH and hardness and in the presence of ion-exchange material. Sea Water - Fluoride forms stable complexes with calcium and magnesium, which are present in sea water. Calcium carbonate precipitation dominates the removal of dissolved fluoride from sea water. The residence time for fluoride in ocean sediment is calculated to be 2-3 million years.

Ecotoxicity: Fluorides have been shown to accumulate in animals that consume fluoride-containing foliage. However, accumulation is primarily in skeletal tissue and therefore, it is unlikely that fluoride will biomagnify up the food chain.

Prevent, by any means available, spillage from entering drains or water courses. **DO NOT** discharge into sewer or waterways.

Persistence and degradability

Ingredient	Persistence: Water/Soil	Persistence: Air
ammonium metavanadate	HIGH	HIGH
water	LOW	LOW

Bioaccumulative potential

Ingredient	Bioaccumulation
ammonium metavanadate	LOW (LogKOW = 2.229)
water	LOW (LogKOW = -1.38)

Mobility in soil

Ingredient	Mobility
ammonium metavanadate	LOW (KOC = 35.04)
water	LOW (KOC = 14.3)

	 Containers may still present a chemical hazard/ danger when empty.
	Return to supplier for reuse/ recycling if possible.
	Otherwise:
	If container can not be cleaned sufficiently well to ensure that residuals do not remain or if the container cannot be used to store the same product, then
	puncture containers, to prevent re-use, and bury at an authorised landfill.
	Where possible retain label warnings and SDS and observe all notices pertaining to the product.
Product / Packaging	Recycle wherever possible.
disposal	 Consult manufacturer for recycling options or consult local or regional waste management authority for disposal if no suitable treatment or disposal facility can be identified.
	 Treat and neutralise at an approved treatment plant. Treatment should involve: Neutralisation with soda-ash or soda-lime followed by: burial in a land-fill specifically licensed to accept chemical and / or pharmaceutical wastes or Incineration in a licensed apparatus (after admixture with suitable combustible material).
	 Decontaminate empty containers with 5% aqueous sodium hydroxide or soda ash, followed by water. Observe all label safeguards until containers are cleaned and destroyed.

SECTION 14 TRANSPORT INFORMATION

Labels Required





Land transport (DOT)

UN number	3264
UN proper shipping name	Corrosive liquid, acidic, inorganic, n.o.s.
Transport hazard class(es)	Class8SubriskNot Applicable
Packing group	II
Environmental hazard	Not Applicable
Special precautions for user	Hazard Label8Special provisions386, B2, IB2, T11, TP2, TP27

Air transport (ICAO-IATA / DGR)

UN number	3264			
UN proper shipping name	CORROSIVE LIQUID, ACIDIC, INORGANIC, N.O.S.			
Transport hazard class(es)	ICAO/IATA Class8ICAO / IATA SubriskNot ApplicableERG Code8L			
Packing group	II			
Environmental hazard	Not Applicable			
Special precautions for user	Special provisionsA3A803Cargo Only Packing Instructions855Cargo Only Maximum Qty / Pack30 LPassenger and Cargo Packing Instructions851Passenger and Cargo Maximum Qty / Pack1 LPassenger and Cargo Limited Quantity Packing InstructionsY840Passenger and Cargo Limited Maximum Qty / Pack0.5 L			

Sea transport (IMDG-Code / GGVSee)

UN number	3264		
UN proper shipping name	Corrosive liquid, acidic, inorganic, n.o.s. *		
Transport hazard class(es)	IMDG Class8IMDG SubriskNot Applicable		
Packing group	II.		

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

Source	Product name	Pollution Category	Ship Type
IMO MARPOL (Annex II) - List of Noxious Liquid Substances Carried in Bulk	Nitric acid (70% and over) Nitric acid (less than 70%)	Y; Y	2 2

SECTION 15 REGULATORY INFORMATION

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

ALUMINIUM(7429-90-5) IS FOUND ON THE FOLLOWING REGULATORY LISTS

- US Alaska Limits for Air Contaminants
- US Vermont Permissible Exposure Limits Table Z-1-A Transitional Limits for Air US - California Permissible Exposure Limits for Chemical Contaminants Contaminants US - Washington Permissible exposure limits of air contaminants US - Hawaii Air Contaminant Limits US - Wyoming Toxic and Hazardous Substances Table Z1 Limits for Air Contaminants US - Massachusetts - Right To Know Listed Chemicals US ACGIH Threshold Limit Values (TLV) US - Michigan Exposure Limits for Air Contaminants US - Minnesota Permissible Exposure Limits (PELs) US ACGIH Threshold Limit Values (TLV) - Carcinogens US ATSDR Minimal Risk Levels for Hazardous Substances (MRLs) US - Oregon Permissible Exposure Limits (Z-1) US EPCRA Section 313 Chemical List US - Pennsylvania - Hazardous Substance List US NIOSH Recommended Exposure Limits (RELs) US - Rhode Island Hazardous Substance List US OSHA Permissible Exposure Levels (PELs) - Table Z1 US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants US - Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminants

ANTIMONY(7440-36-0) IS FOUND ON THE FOLLOWING REGULATORY LISTS

- US Alaska Limits for Air Contaminants
- US California Permissible Exposure Limits for Chemical Contaminants
- US Hawaii Air Contaminant Limits
- US Idaho Limits for Air Contaminants
- US Massachusetts Right To Know Listed Chemicals
- US Michigan Exposure Limits for Air Contaminants
- US Minnesota Permissible Exposure Limits (PELs)
- US Oregon Permissible Exposure Limits (Z-1)
- US Pennsylvania Hazardous Substance List
- US Rhode Island Hazardous Substance List
- US Tennessee Occupational Exposure Limits Limits For Air Contaminants
- US Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminants

ARSENIC(7440-38-2) IS FOUND ON THE FOLLOWING REGULATORY LISTS

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

Monographs

- US Alaska Limits for Air Contaminants
- US California OEHHA/ARB Acute Reference Exposure Levels and Target Organs (RELs)
- US California OEHHA/ARB Chronic Reference Exposure Levels and Target Organs (CRELs)
- US California Permissible Exposure Limits for Chemical Contaminants
- US Hawaii Air Contaminant Limits
- US Idaho Limits for Air Contaminants
- US Massachusetts Right To Know Listed Chemicals
- US Minnesota Permissible Exposure Limits (PELs)
- US New Jersey Right to Know Special Health Hazard Substance List (SHHSL):
- Carcinogens
- US Pennsylvania Hazardous Substance List
- US Tennessee Occupational Exposure Limits Limits For Air Contaminants
- US Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminants
- US Vermont Permissible Exposure Limits Table Z-1-A Transitional Limits for Air Contaminants

BARIUM(7440-39-3) IS FOUND ON THE FOLLOWING REGULATORY LISTS

- US Alaska Limits for Air Contaminants
- US Hawaii Air Contaminant Limits
- US Idaho Limits for Air Contaminants
- US Massachusetts Right To Know Listed Chemicals
- US Minnesota Permissible Exposure Limits (PELs)
- US Pennsylvania Hazardous Substance List
- US Rhode Island Hazardous Substance List
- US Tennessee Occupational Exposure Limits Limits For Air Contaminants
- US Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminants

US - Vermont Permissible Exposure Limits Table Z-1-A Transitional Limits for Air Contaminants

- US Washington Permissible exposure limits of air contaminants
- US ACGIH Threshold Limit Values (TLV)
- US ACGIH Threshold Limit Values (TLV) Carcinogens US EPA Carcinogens Listing
- US EPCRA Section 313 Chemical List
- US OSHA Permissible Exposure Levels (PELs) Table Z1
- US Toxic Substances Control Act (TSCA) Chemical Substance Inventory

- US Toxic Substances Control Act (TSCA) Chemical Substance Inventory US - Vermont Permissible Exposure Limits Table Z-1-A Transitional Limits for Air Contaminants US - Washington Permissible exposure limits of air contaminants US - Wyoming Toxic and Hazardous Substances Table Z1 Limits for Air Contaminants US ACGIH Threshold Limit Values (TLV)
 - US Clean Air Act Hazardous Air Pollutants
 - US CWA (Clean Water Act) Priority Pollutants
 - US CWA (Clean Water Act) Toxic Pollutants

 - US EPCRA Section 313 Chemical List

US ACGIH Threshold Limit Values (TLV)

US Clean Air Act - Hazardous Air Pollutants

US CWA (Clean Water Act) - Priority Pollutants

US NIOSH Recommended Exposure Limits (RELs)

US OSHA Permissible Exposure Levels (PELs) - Table Z1

US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory

US CWA (Clean Water Act) - Toxic Pollutants

US EPCRA Section 313 Chemical List

US NIOSH Recommended Exposure Limits (RELs) US OSHA Permissible Exposure Levels (PELs) - Table Z1

US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory

US - Washington Toxic air pollutants and their ASIL, SQER and de minimis emission values

US National Toxicology Program (NTP) 14th Report Part A Known to be Human Carcinogens

US - Washington Permissible exposure limits of air contaminants

US ATSDR Minimal Risk Levels for Hazardous Substances (MRLs)

US ACGIH Threshold Limit Values (TLV) - Carcinogens

Version No: 1.1

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Catalogue number: CWW-TM-C Print Date: 05/31/2017 CWW-TM-C - Certified Waste Water Trace Metals Solution C International Agency for Research on Cancer (IARC) - Agents Classified by the IARC US - Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminants Monographs US - Vermont Permissible Exposure Limits Table Z-1-A Transitional Limits for Air US - Alaska Limits for Air Contaminants Contaminants US - Washington Permissible exposure limits of air contaminants US - California - Proposition 65 - Priority List for the Development of MADLs for Chemicals Causing Reproductive Toxicity US - Washington Toxic air pollutants and their ASIL, SQER and de minimis emission values US - California OEHHA/ARB - Chronic Reference Exposure Levels and Target Organs US - Wyoming Toxic and Hazardous Substances Table Z1 Limits for Air Contaminants (CRELs) US - Wyoming Toxic and Hazardous Substances Table Z-2 Acceptable ceiling concentration, US - California Permissible Exposure Limits for Chemical Contaminants Acceptable maximum peak above the acceptable ceiling concentration for an 8-hr shift US - California Proposition 65 - Carcinogens US ACGIH Threshold Limit Values (TLV) US - California Proposition 65 - No Significant Risk Levels (NSRLs) for Carcinogens US ACGIH Threshold Limit Values (TLV) - Carcinogens US - Hawaii Air Contaminant Limits US ATSDR Minimal Risk Levels for Hazardous Substances (MRLs) US - Idaho - Acceptable Maximum Peak Concentrations US Clean Air Act - Hazardous Air Pollutants US - Idaho - Limits for Air Contaminants US CWA (Clean Water Act) - Priority Pollutants US - Massachusetts - Right To Know Listed Chemicals US CWA (Clean Water Act) - Toxic Pollutants US - Michigan Exposure Limits for Air Contaminants US EPA Carcinogens Listing US - Minnesota Permissible Exposure Limits (PELs) US EPCRA Section 313 Chemical List US - New Jersey Right to Know - Special Health Hazard Substance List (SHHSL): US National Toxicology Program (NTP) 14th Report Part A Known to be Human Carcinogen Carcinogens US NIOSH Recommended Exposure Limits (RELs) US - Oregon Permissible Exposure Limits (Z-1) US OSHA Permissible Exposure Levels (PELs) - 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US - New Jersey Right to Know - Special Health Hazard Substance List (SHHSL): Carcinogens

- US Oregon Permissible Exposure Limits (Z-1)
- US Oregon Permissible Exposure Limits (Z-2)
- US Pennsylvania Hazardous Substance List
- US Rhode Island Hazardous Substance List
- US Tennessee Occupational Exposure Limits Limits For Air Contaminants

CHROMIUM(7440-47-3) IS FOUND ON THE FOLLOWING REGULATORY LISTS

- International Agency for Research on Cancer (IARC) Agents Classified by the IARC
- Monographs
- US Alaska Limits for Air Contaminants
- US California Permissible Exposure Limits for Chemical Contaminants
- US Hawaii Air Contaminant Limits
- US Idaho Limits for Air Contaminants
- US Massachusetts Right To Know Listed Chemicals
- US Michigan Exposure Limits for Air Contaminants
- US Oregon Permissible Exposure Limits (Z-1)
- US Pennsylvania Hazardous Substance List
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- US Tennessee Occupational Exposure Limits Limits For Air Contaminants
- US Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminants

COBALT(7440-48-4) IS FOUND ON THE FOLLOWING REGULATORY LISTS

US - Vermont Permissible Exposure Limits Table Z-1-A Transitional Limits for Air Contaminants

US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory

US - Washington Permissible exposure limits of air contaminants

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- US Clean Air Act Hazardous Air Pollutants
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- US EPCRA Section 313 Chemical List
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Carcinogens US NIOSH Recommended Exposure Limits (RELs) US - Oregon Permissible Exposure Limits (Z-1) US OSHA Permissible Exposure Levels (PELs) - Table Z1 US - Pennsylvania - Hazardous Substance List US Priority List for the Development of Proposition 65 Safe Harbor Levels - No Significant Risk US - Rhode Island Hazardous Substance List Levels (NSRLs) for Carcinogens and Maximum Allowable Dose Levels (MADLs) for Chemicals Causing Reproductive Toxicity US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory US - Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminants COPPER(7440-50-8) IS FOUND ON THE FOLLOWING REGULATORY LISTS US - Alaska Limits for Air Contaminants US - Vermont Permissible Exposure Limits Table Z-1-A Transitional Limits for Air US - California OEHHA/ARB - Acute Reference Exposure Levels and Target Organs (RELs) Contaminants US - Washington Permissible exposure limits of air contaminants US - California Permissible Exposure Limits for Chemical Contaminants US - Washington Toxic air pollutants and their ASIL, SQER and de minimis emission values US - Hawaii Air Contaminant Limits US - Idaho - Limits for Air Contaminants US - Wyoming Toxic and Hazardous Substances Table Z1 Limits for Air Contaminants US ACGIH Threshold Limit Values (TLV) US - Massachusetts - Right To Know Listed Chemicals US - Michigan Exposure Limits for Air Contaminants US ATSDR Minimal Risk Levels for Hazardous Substances (MRLs) US - Minnesota Permissible Exposure Limits (PELs) US CWA (Clean Water Act) - Priority Pollutants US CWA (Clean Water Act) - Toxic Pollutants US - Oregon Permissible Exposure Limits (Z-1) US EPA Carcinogens Listing US - Pennsylvania - Hazardous Substance List US EPCRA Section 313 Chemical List US - Rhode Island Hazardous Substance List US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants US NIOSH Recommended Exposure Limits (RELs) US OSHA Permissible Exposure Levels (PELs) - Table Z1 US - Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminants US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory IRON(7439-89-6) IS FOUND ON THE FOLLOWING REGULATORY LISTS International Agency for Research on Cancer (IARC) - Agents Classified by the IARC US - Oregon Permissible Exposure Limits (Z-1) Monographs US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants US - California OEHHA/ARB - Chronic Reference Exposure Levels and Target Organs US - Washington Permissible exposure limits of air contaminants (CRELs) US - Wyoming Toxic and Hazardous Substances Table Z1 Limits for Air Contaminants US - California Permissible Exposure Limits for Chemical Contaminants US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory US - Hawaii Air Contaminant Limits US - Michigan Exposure Limits for Air Contaminants LEAD(7439-92-1) IS FOUND ON THE FOLLOWING REGULATORY LISTS International Agency for Research on Cancer (IARC) - Agents Classified by the IARC US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants Monographs US - Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminants US - Alaska Limits for Air Contaminants US - Vermont Permissible Exposure Limits Table Z-1-A Transitional Limits for Air US - California - Proposition 65 - Priority List for the Development of MADLs for Chemicals Contaminants Causing Reproductive Toxicity US - Washington Permissible exposure limits of air contaminants US - California Permissible Exposure Limits for Chemical Contaminants US - Washington Toxic air pollutants and their ASIL. 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- US Hawaii Air Contaminant Limits
- US Idaho Limits for Air Contaminants
- US Michigan Exposure Limits for Air Contaminants
- US Minnesota Permissible Exposure Limits (PELs)
- US Oregon Permissible Exposure Limits (Z-1)
- US Tennessee Occupational Exposure Limits Limits For Air Contaminants

- US Clean Air Act Hazardous Air Pollutants
- US EPCRA Section 313 Chemical List
- US OSHA Permissible Exposure Levels (PELs) Table Z1

US - Washington Permissible exposure limits of air contaminants

US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory

US - Washington Toxic air pollutants and their ASIL, SQER and de minimis emission values

US - Wyoming Toxic and Hazardous Substances Table Z1 Limits for Air Contaminants

MERCURY (ELEMENTAL)(7439-97-6) IS FOUND ON THE FOLLOWING REGULATORY LISTS

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

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- US California Permissible Exposure Limits for Chemical Contaminants
- US California Proposition 65 Reproductive Toxicity
- US Hawaii Air Contaminant Limits
- US Idaho Acceptable Maximum Peak Concentrations
- US Idaho Limits for Air Contaminants
- US Massachusetts Right To Know Listed Chemicals
- US Michigan Exposure Limits for Air Contaminants
- US Minnesota Permissible Exposure Limits (PELs)
- US Oregon Permissible Exposure Limits (Z-2)
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- MOLYBDENUM(7439-98-7) IS FOUND ON THE FOLLOWING REGULATORY LISTS
- US Alaska Limits for Air Contaminants
- US Hawaii Air Contaminant Limits

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- US Idaho Limits for Air Contaminants
- US Massachusetts Right To Know Listed Chemicals
- US Minnesota Permissible Exposure Limits (PELs)
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- US Tennessee Occupational Exposure Limits Limits For Air Contaminants

NICKEL(7440-02-0) IS FOUND ON THE FOLLOWING REGULATORY LISTS

US - Alaska Limits for Air Contaminants

- US California OEHHA/ARB Acute Reference Exposure Levels and Target Organs (RELs)
- US California OEHHA/ARB Chronic Reference Exposure Levels and Target Organs
- (CRELs)
- US California Permissible Exposure Limits for Chemical Contaminants
- US California Proposition 65 Carcinogens
- US Hawaii Air Contaminant Limits
- US Idaho Limits for Air Contaminants
- US Massachusetts Right To Know Listed Chemicals
- US Michigan Exposure Limits for Air Contaminants
- US Minnesota Permissible Exposure Limits (PELs)
- US New Jersey Right to Know Special Health Hazard Substance List (SHHSL):
- Carcinogens
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- US Pennsylvania Hazardous Substance List
- US Rhode Island Hazardous Substance List
- US Tennessee Occupational Exposure Limits Limits For Air Contaminants
- US Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminants

SELENIUM(7782-49-2) IS FOUND ON THE FOLLOWING REGULATORY LISTS

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

- US Alaska Limits for Air Contaminants
- US California OEHHA/ARB Acute Reference Exposure Levels and Target Organs (RELs) US - California OEHHA/ARB - Chronic Reference Exposure Levels and Target Organs
- (CRELs)
- US Hawaii Air Contaminant Limits
- US Idaho Limits for Air Contaminants
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- US Minnesota Permissible Exposure Limits (PELs)
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SILVER(7440-22-4) IS FOUND ON THE FOLLOWING REGULATORY LISTS

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US - Washington Toxic air pollutants and their ASIL, SQER and de minimis emission values

- US ATSDR Minimal Risk Levels for Hazardous Substances (MRLs)

US - Washington Permissible exposure limits of air contaminants

- US Clean Air Act Hazardous Air Pollutants
- US CWA (Clean Water Act) Priority Pollutants US CWA (Clean Water Act) - Toxic Pollutants
- US EPA Carcinogens Listing
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US Priority List for the Development of Proposition 65 Safe Harbor Levels - No Significant Risk Levels (NSRLs) for Carcinogens and Maximum Allowable Dose Levels (MADLs) for

- Chemicals Causing Reproductive Toxicity

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- US NIOSH Recommended Exposure Limits (RELs)
- US OSHA Permissible Exposure Levels (PELs) Table Z1

US Priority List for the Development of Proposition 65 Safe Harbor Levels - No Significant Risk

Levels (NSRLs) for Carcinogens and Maximum Allowable Dose Levels (MADLs) for Chemicals Causing Reproductive Toxicity

US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory

- US Washington Permissible exposure limits of air contaminants
- US Washington Toxic air pollutants and their ASIL, SQER and de minimis emission values

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- US ACGIH Threshold Limit Values (TLV)
- US ATSDR Minimal Risk Levels for Hazardous Substances (MRLs)
- US Clean Air Act Hazardous Air Pollutants
- US CWA (Clean Water Act) Priority Pollutants US CWA (Clean Water Act) - Toxic Pollutants

US NIOSH Recommended Exposure Limits (RELs)

US OSHA Permissible Exposure Levels (PELs) - Table Z1

US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory

US EPCRA Section 313 Chemical List

US EPA Carcinogens Listing

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US - Pennsylvania - Hazardous Substance List	US EPCRA Section 313 Chemical List
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STRONTIUM(7440-24-6) IS FOUND ON THE FOLLOWING REGULATORY LISTS	
US ATSDR Minimal Risk Levels for Hazardous Substances (MRLs)	US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory
THALLIUM(7440-28-0) IS FOUND ON THE FOLLOWING REGULATORY LISTS	
US - Massachusetts - Right To Know Listed Chemicals	US CWA (Clean Water Act) - Priority Pollutants
US - Minnesota Permissible Exposure Limits (PELs)	US CWA (Clean Water Act) - Toxic Pollutants
US - Pennsylvania - Hazardous Substance List	US EPCRA Section 313 Chemical List
US - Rhode Island Hazardous Substance List	US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory
US ACGIH Threshold Limit Values (TLV)	
AMMONIUM METAVANADATE(7803-55-6) IS FOUND ON THE FOLLOWING REGULATOR	Y LISTS
US - California OEHHA/ARB - Acute Reference Exposure Levels and Target Organs (RELs)	US EPCRA Section 313 Chemical List
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US - Pennsylvania - Hazardous Substance List	
ZINC(7440-66-6) IS FOUND ON THE FOLLOWING REGULATORY LISTS	
International Agency for Research on Cancer (IARC) - Agents Classified by the IARC	US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants
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US - California Permissible Exposure Limits for Chemical Contaminants	US CWA (Clean Water Act) - Priority Pollutants
US - Hawaii Air Contaminant Limits	US CWA (Clean Water Act) - Toxic Pollutants
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US - Michigan Exposure Limits for Air Contaminants	US EPCRA Section 313 Chemical List
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US - Pennsylvania - Hazardous Substance List	
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US - Pennsylvania - Hazardous Substance List

US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory

Federal Regulations

Superfund Amendments and Reauthorization Act of 1986 (SARA)

SECTION 311/312 HAZARD CATEGORIES

Immediate (acute) health hazard	Yes
Delayed (chronic) health hazard	No
Fire hazard	No
Pressure hazard	No
Reactivity hazard	No

US. EPA CERCLA HAZARDOUS SUBSTANCES AND REPORTABLE QUANTITIES (40 CFR 302.4)

Name	Reportable Quantity in Pounds (Ib)	Reportable Quantity in kg
Antimony	5000	2270
Arsenic	1	0.454
Beryllium	10	4.54
Cadmium	10	4.54
Chromium	5000	2270
Copper	5000	2270
Lead	10	4.54
Mercury	1	0.454
Nickel	100	45.4
Selenium	100	45.4
Silver	1000	454
Thallium	1000	454
Ammonium vanadate	1000	454
Zinc	1000	454
Nitric acid	1000	454
Hydrofluoric acid	100	45.4

State Regulations

US. CALIFORNIA PROPOSITION 65

WARNING: This product contains a chemical known to the State of California to cause cancer and birth defects or other reproductive harm

US - CALIFORNIA PREPOSITION 65 - CARCINOGENS & REPRODUCTIVE TOXICITY (CRT): LISTED SUBSTANCE

Beryllium and beryllium compounds: Beryllium, Cadmium and cadmium compounds: Cadmium, Cobalt metal powder, Lead and lead compounds: Lead, Mercury and mercury compounds, Nickel (Metallic) Listed

National Inventory	Status		
Australia - AICS	Y		
Canada - DSL	Y		
Canada - NDSL	N (strontium; thallium; lead; zinc; ammonium metavanadate; copper; boron; water; antimony; barium; selenium; aluminium; mercury (elemental); molybdenum; arsenic; cobalt; nickel; manganese(II) acetate tetrahydrate; iron; beryllium; chromium; silver; hydrofluoric acid; cadmium; nitric acid)		
China - IECSC	Y		
Europe - EINEC / ELINCS / NLP	Y		
Japan - ENCS	N (strontium; thallium; zinc; copper; boron; water; antimony; barium; selenium; aluminium; mercury (elemental); molybdenum; arsenic; cobalt; nickel; manganese(II) acetate tetrahydrate; iron; beryllium; chromium; silver; hydrofluoric acid; cadmium; nitric acid)		
Korea - KECI	Υ		
New Zealand - NZIoC	Υ		
Philippines - PICCS	Υ		
USA - TSCA	Υ		
Legend:	Y = All ingredients are on the inventory N = Not determined or one or more ingredients are not on the inventory and are not exempt from listing(see specific ingredients in brackets)		

SECTION 16 OTHER INFORMATION

Other information

Ingredients with multiple cas numbers

Name	CAS No
aluminium	7429-90-5, 91728-14-2
copper	7440-50-8, 133353-46-5, 133353-47-6, 195161-80-9, 65555-90-0, 72514-83-1

hydrofluoric acid

Catalogue number: CWW-TM-C

CWW-TM-C - Certified Waste Water Trace Metals Solution C

Version No: 1.1

7664-39-3, 790596-14-4

Classification of the preparation and its individual components has drawn on official and authoritative sources as well as independent review by the Chernwatch 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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