

QCS-21

High-Purity Standards

Catalogue number: QCS-21

Version No: 3.3 Safety Data Sheet according to OSHA HazCom Standard (2012) requirements

Chemwatch Hazard Alert Code: 3 Issue Date: 06/03/2017

Print Date: 06/03/2017 S.GHS.USA.EN

SECTION 1 IDENTIFICATION

Product Identifier

Product name	QCS-21
Synonyms 100µg/mL Antimony, Arsenic, Beryllium, Calcium, Cadmium, Chromium, Cobalt, Copper, Iron, Lead, Lithium, Magnesium, Manganese, Moly Selenium, Strontium, Thallium, Titanium, Vanadium, Zinc in 4% HNO3 + Tr HF	
Proper shipping name	Corrosive liquid, acidic, inorganic, n.o.s. (contains nitric acid and hydrofluoric acid)
Other means of identification	QCS-21

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

Association / Organisatio	INFOTRAC
Emergency telephon number	e 1-800-535-5053
Other emergency telephon number	P 1-352-323-3500

SECTION 2 HAZARD(S) IDENTIFICATION

Classification of the substance or mixture

Classification	Serious Eye Damage Category 1, Acute Toxicity (Oral) Category 4, Acute Toxicity (Dermal) Category 3, Metal Corrosion Category 1, Skin Corrosion/Irritation Category 1A

Label elements

Hazard pictogram(s)	
SIGNAL WORD	DANGER
SIGNAL WORD	DANGER

Hazard statement(s)

H302	Harmful if swallowed.	
H311 Toxic in contact with skin.		
H290	May be corrosive to metals.	
H314	Causes severe skin burns and eye damage.	

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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
7440-36-0	0.01	antimony
7440-38-2	0.01	arsenic
471-34-1	0.01 (as Ca)	calcium carbonate
543-81-7	0.01 (as Be)	beryllium acetate
7440-43-9	0.01	cadmium
7440-47-3	0.01	chromium
7440-48-4	0.01	cobalt
7440-50-8	0.01	copper
7439-89-6	0.01	iron
7439-92-1	0.01	lead
554-13-2	0.01 (as Li)	lithium carbonate
7439-95-4	0.01	magnesium
7439-98-7	0.01	molybdenum
7440-02-0	0.01	nickel
7782-49-2	0.01	selenium
10042-76-9	0.01 (as Sr)	strontium nitrate
7440-28-0	0.01	thallium
7440-32-6	0.01	titanium
7803-55-6	0.01 (as V)	ammonium metavanadate
7440-66-6	0.01	zinc
7697-37-2	4	nitric acid
7664-39-3	0-0.49	hydrofluoric acid
7732-18-5	balance	water
6156-78-1	0.01 (as Mn)	manganese(II) acetate tetrahydrate

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.		 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.

- Use compresses if running water is not available.
- Do NOT apply ice as this may lower body temperature and cause further damage.
- Do NOT break blisters or apply butter or ointments; this may cause infection.
 - Protect burn by cover loosely with sterile, 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 burn area above heart level, if possible.
 Cover the person with coat or blanket.
- Seek medical assistance.
- For third-degree burns
- Seek immediate medical or emergency assistance.
- In the mean time:
- Protect burn area cover loosely with sterile, nonstick bandage or, for large areas, a sheet or other material that will not leave lint in wound.
- Separate burned toes and fingers with dry, sterile dressings.
- Do not soak burn in water or apply ointments or butter; this may cause infection.
- To prevent shock see above.
- + For an airway burn, do not place pillow under the person's head when the person is lying down. This can close 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 irritation or skin burns: Avoid further contact. Immediately remove contaminated clothing, including footwear. Flush skin under running water for 15 minutes. Avoiding contamination of the hands, massage calcium gluconate gel into affected areas, pay particular attention to creases in skin. Contact the Poisons Information Centre. Continue gel application for at least 15 minutes after burning sensation ceases. ▶ If pain recurs, repeat application of calcium gluconate gel or apply every 20 minutes. F If no gel is available, continue washing for at least 15 minutes, using soap if available. If patient is conscious, give six calcium gluconate or calcium carbonate tablets in water by mouth. Transport to hospital, or doctor, urgently. 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. Inhalation 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. Lav 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, urgently. ▶ 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 vo If vomiting occurs, lean patient forward or place on left side (head-down position, if possible) to maintain open airway and prevent aspiration. Ingestion 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 casuality 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

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

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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 colle	cted from subjects NOT exposed.		
NS: Non-specific determinant; Also seen after	exposure to other materials		
SQ: Semi-quantitative determinant - Interpreta	tion 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 Index Fluorides in urine 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

- There is no restriction on the type of extinguisher which may be used.
- Use extinguishing media suitable for surrounding area.

Special hazards arising from the substrate or mixture

Fire Incompatibility None known.

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. May emit corrosive fumes.

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. 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.

SECTION 7 HANDLING AND STORAGE

Precautions for safe handling

Safe handling	 Avoid all personal contact, including inhalation. Wear protective clothing when risk of exposure occurs. Use in a well-ventilated area. WARNING: To avoid violent reaction, ALWAYS add material to water and NEVER water to material. Avoid smoking, naked lights or ignition sources. Avoid contact with incompatible materials. When handling, DO NOT eat, drink or smoke. 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.
Other information	 Store in original containers. Keep containers securely sealed. Store in a cool, dry, well-ventilated area. Store away from incompatible materials and foodstuff containers. Protect containers against physical damage and check regularly for leaks. Observe manufacturer's storage and handling recommendations contained within this SDS.

Conditions for safe storage, including any incompatibilities

Suitable container	 DO NOT use aluminium or galvanised containers 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 Iow 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 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	 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 heart 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 fore generates sufficient heat in the small region of mixing to cause some of the water to bol explosively. The resulting fourping can acids react with active metals, including such structural metals as aluminum and iron, to release hydrogen, a flammable gas. Inorganic acids react with active metals, including such structural metals as aluminum and iron, to release hydrogen, a flammable gas. Inorganic acids react with cyanide compounds to release gaseous hydrogen cyanide. Inorganic acids (an eitablyse (increase the rate of) chemical reactions. WARNING: Avoid or control reaction with peroxides. All <i>transition metal</i> peroxides should be considered as potentially explosive. For example transition metal complexes of alky hydroperoxides may decompose explosively. The pi-complexes formed between chromium(0), vanadium(0) and other transition metals (haloarene-metal complexes) and mono-or poly-fluorobenzene show externe sensitivity to heat and are explosive. Avoid forcation with boron, bromine pertafluoride, romine trifluoride, calcium hydride, oxygen difluoride, platinum, potassium. i react with water forming acidic solutions. are violent reactive with boron, bromine pertafluoride hornine trifluoride, calcium metals (haloarene-metal complexes) and mono-or poly-fluorobenzene show externe sensitivity to heat and are explosive. corrode metals in presence of motisture react with water forming acidic so

Control parameters

OCCUPATIONAL EXPOSURE LIMITS (OEL)

INGREDIENT DATA

INGREDIENT DATA						
Source	Ingredient	Material name	TWA	STEL	Peak	Notes
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)	calcium carbonate	Calcium salt of carbonic acid [Note: Occurs in nature as as limestone, chalk, marble, dolomite, aragonite, calcite and oyster shells.]	10 (total), 5 (resp) mg/m3	Not Available	Not Available	Total dust
US OSHA Permissible Exposure Levels (PELs) - Table Z1	calcium carbonate	Calcium carbonate	15 mg/m3	Not Available	Not Available	Not Available
US OSHA Permissible Exposure Levels (PELs) - Table Z1	calcium carbonate	Calcium carbonate - Respirable fraction	5 mg/m3	Not Available	Not Available	Not Available
US NIOSH Recommended Exposure Limits (RELs)	calcium carbonate	Calcium carbonate, Natural calcium carbonate [Note: Calcite & aragonite are commercially important natural calcium carbonates.]	10 (total), 5 (resp) mg/m3	Not Available	Not Available	Not Available
US NIOSH Recommended Exposure Limits (RELs)	calcium carbonate	Calcium carbonate, Natural calcium carbonate [Note: Marble is a metamorphic form of calcium carbonate.]	10 (total), 5 (resp) mg/m3	Not Available	Not Available	Not Available
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
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 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 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

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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)
US 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
calcium carbonate	Limestone; (Calcium carbonate; Dolomite)	45 mg/m3	500 mg/m3	3,000 mg/m3
calcium carbonate	Carbonic acid, calcium salt	45 mg/m3	210 mg/m3	1,300 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
lithium carbonate	Lithium carbonate	0.44 mg/m3	4.8 mg/m3	100 mg/m3
magnesium	Magnesium	18 mg/m3	200 mg/m3	1,200 mg/m3
nolybdenum	Molybdenum	30 mg/m3	330 mg/m3	2,000 mg/m3
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
strontium nitrate	Strontium nitrate	5.7 mg/m3	62 mg/m3	370 mg/m3
thallium	Thallium	0.06 mg/m3	13 mg/m3	20 mg/m3
litanium	Titanium	30 mg/m3	330 mg/m3	2,000 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
nydrofluoric acid	Hydrogen fluoride; (Hydrofluoric acid)	Not Available	Not Available	Not Available
nanganese(II) acetate etrahydrate	Acetic acid, manganese(2+) salt, tetrahydrate	13 mg/m3	22 mg/m3	740 mg/m3
nanganese(II) acetate etrahydrate	Acetic acid, manganese(II) salt (2:1)	9.4 mg/m3	16 mg/m3	96 mg/m3

Ingredient	Original IDLH	Revised IDLH
antimony	80 mg/m3	50 mg/m3
arsenic	100 mg/m3	5 mg/m3
calcium carbonate	Not Available	Not Available
beryllium acetate	10 mg/m3	4 mg/m3
cadmium	50 mg/m3 / 9 mg/m3	9 mg/m3 / 9 [Unch] mg/m3
chromium	N.E. / N.E.	250 mg/m3
cobalt	20 mg/m3	20 [Unch] mg/m3
copper	N.E. / N.E.	100 mg/m3
iron	Not Available	Not Available
lead	700 mg/m3	100 mg/m3
lithium carbonate	Not Available	Not Available
magnesium	Not Available	Not Available
molybdenum	N.E. / N.E.	5,000 mg/m3
nickel	N.E. / N.E.	10 mg/m3
selenium	Unknown mg/m3 / Unknown ppm	1 mg/m3
strontium nitrate	Not Available	Not Available
thallium	Not Available	Not Available
titanium	Not Available	Not Available
ammonium metavanadate	Not Available	Not Available
zinc	Not Available	Not Available
nitric acid	100 ppm	25 ppm
hydrofluoric acid	30 ppm	30 [Unch] ppm

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water	Not Available	Not Available
waler	Not Available	NUL AVAIIADIE
manganese(II) acetate tetrahydrate	N.E. / N.E.	500 mg/m3

Exposure controls

Appropriate enclosed Engineering controls on used to stroke a batter's towater and the band Weid-degree angenetic controls are the height be enclosed and the phage being controls are the height being controls are theight being being are the height being controls are		_			
Appropriate engineering 029-05 ms (80-100 mins.) accosh, turnes from pouring operations, intermittent container filling, tox speed conveyer transfers, welding, spany drit, plating 0.5 ms (100.200 mins.) accosh, turnes from pouring operations, intermittent container filling, tox speed conveyer transfers, welding, spany drit, plating 0.5 ms (100.200 mins.) accosh, turnes from pouring operations, intermittent container filling, tox speed conveyer transfers, welding, spany drit, plating 0.5 ms (100.200 mins.) accosh, turnes from pouring operations, intermittent container filling, tox speed conveyer transfers, welding, spany drit, plating 0.5 ms (100.200 mins.) accosh, turnes from pouring operations, intermittent container filling, tox speed conveyer transfers, welding, spany drit, plating 0.5 ms (100.200 mins.) With each range the appropriate value depends or: Upper end of the range Upper end of the range 1. Room air currents minimal of favourable to capture 1. Disturbing room air currents 2. Contaminutes of trigh postcolin, heavy use 1. Room air currents minimal of favourable to capture 1. Disturbing room air currents with a velocity fails rapidly with distance away from the operating of a simple most on order 3. Simple theory shows that air velocity fails rapidly with distance away from the operating of a simple most on order I. Room air current minimal of favourable to capture 1. Simult hoodocal current on order 1. Simult hoodocal current on order		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. Type of Contaminant: Air Speed: 0.25-0.5 m/s (50-100)			
Appropriate engineering temps temps temps Appropriate engineering accident to the temps temps temps accident to the temps temps temps temps temps accident to the temps temps temps temps temps temps accident to the temps temps temps temps temps temps temps accident to the temps temps <t< th=""><th></th><th>Type of Contaminant:</th><th></th><th>Air Speed:</th></t<>		Type of Contaminant:		Air Speed:	
Appropriate engineering control India tume, picking (released at two velocity vito zone of achie generation) Itims, indicessing (achieve generation) Itims, indicessing (achieve generation) <th< th=""><th></th><th colspan="3">solvent, vapours, degreasing etc., evaporating from tank (in still air).</th></th<>		solvent, vapours, degreasing etc., evaporating from tank (in still air).			
dreed singly gives parting in statew boots, durin limiting, ordreyer boarding, during to data give discription into interval 12: 5 mg 200-001 (min), interval grinding, discription blasting, turbling, high speed wheel generated dusts (released at high initial velocity into zone of vary high rapid 2:5-10 m/s (500-2000 (min)). 2: 5-10 m/s (500-2000 (min)). Within each range the appropriate value depends or: Lower and of the range Upper and of the range Contaminants of hold bookdy or of nulances value only. 2: Contaminants of hold bookdy or of nulances value only. 2: Contaminants of hold bookdy or of nulances value only. 3: High production, heavy use 4: Large hood or large at mass in motion 4: Small hood-local control only Single heavy shows that at velocity fails rapidly with distance avery from the contraction pice. Welchog generally decreases with the square of distance home the contraction pice (min) of the oblastical according to the structure on the contraction pice. Welchog generally decreases with the square of distance home the contraction pice. The ary velocity at the contraction pice (min) of the oblastical according to the structure on the contraction pice. Welchog generally decreases with the square of distance home the distance in the contraction pice. The ary velocity at the contraction pice (min) of the oblastical according to the structure on the contraction pice. Welchog generally decreases are multiplied by factors of 10 or more when estruction pice (min) for structure on a distance from the contraction pice. The minister distance more the arge and the square distance from the contraction pice. The minister distance more the arge and the structure distance more distance more when estruction pice (min) for structure			s, welding, spray drift, plating		
I air motion). (fmin.) White such range the appropriate value depends on: Upper end of the range (Ipper end of the range 1. Room air currents minimal or favourable to capture 1. Disturbing room air currents 2. Contaminants of high toxicity 2. Contaminants of high toxicity 3. High production, heavy use 4. Small hood/coal control only 4. Large fixed of large air mass in motion 4. Small hood/coal control only Simple theory shows that air velocities approximation on the contraining of a simple extractor, price (maining cases). Therefore the air speed at the extractoric price (maining), alter reference to a distance from the extractor price (maining cases). Therefore the air speed at the extractoric price (maining), alter reference to a distance from the extractoric price (maining cases). Therefore the air speed at the extractoric price (maining), alter reference to a distance from the extractoric price (maining cases). Therefore the air speed at the extractoric price (maining), alter reference to a distance from the extractoric price (maining cases). Therefore the air speed at the extractoric price (maining cases). Therefore the air speed at the extractoric price (maining cases). Therefore the air speed at the extractoric price (maining cases). Therefore the air speed at the extractoric price (maining cases). Therefore the air speed at the extractoric price (maining cases). Therefore the air speed at the extractoric price (maining cases). Therefore the air speed at the extractoric price (maining cases). Therefore the air speed at the extractoric price (maining cases). Therefore the air speed at the extractoric price (maining cases). Therefore the air speed at the extractoric price	controis		scharge (active generation into		
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3: Intermittent, low production. 3: High production, heavy use 4: Large hood or large air mass in motion 4: Small hood-local control only Single theory shows that air velocity falls rapidly with distance away from the opening of a simple extraction pipe. Velocity generally decreases with the square of distance from the extraction point (in simple cases). Therefore the air speed at the extraction point should be adjusted, accordingly, after reference to distance from the extraction point (in simple cases). Therefore the air speed at the extraction point of the adjusted, accordingly, after reference to observe the extraction point. Other morehand case animum of 12: Ne (20-06) (fimplif or extraction of solvenes) generated in a tank 2 meter distant from the extraction point. Other morehane extraction systems are installed or used. Personal protection Safety glasses with upperformed side shelds may be used where continuous eye protection is desirable, as in laboratories; spectacles are not sufficient where complete eye protection is needed such as when handing bulk-quantities, where there is a danger of splashing, or if the material may be under pressure. • Determine ball (20 or, 8) in minimum may be required for supplementary but new for primary protection of eyes; these afford face protection. • Aternatively a gas mask may replace splash togogles and face shields. • Contract incess may pose a special hazar, so to contract incess may absorb and concentrate infrants. A written policy document, describing the wearing of lenses or netricinal exposure, begin barger, so that environ of a wearing of lenses or netricinal exposure. Begin eye and tace or installed or used. • Safety glasses with upperformed side shie		1: Room air currents minimal or favourable to capture	1: Disturbing room air currents		
4: Large hood or large air mass in motion 4: Small hood-local control only Simple theory shows that air velocity fails rapidly with distance away from the opening of a simple extraction pipe. Velocity generally decreases with the square of distance from the extraction point (in simple cases). Therefore the air speed at the extraction point, or example, should be a minutom of 12 m/s (200400 fmin) for extraction of a solvents generated in a tank 2 meters distant from the extraction point. Other methenical considerations, producing performance deficits within the extraction apparatus, make it essential that theoretical air velocites are multiplied by factors of 10 or more when extraction systems are installed or used. Personal protection View View View View View View View View		2: Contaminants of low toxicity or of nuisance value only.	f nuisance value only. 2: Contaminants of high toxicity		
Simple theory shows that air velocity falls rapidly with distance away from the opening of a simple extraction pior. (Velocity generally decreases with the square distance from the extraction point (in simple cases). Therefore the air speed at the extraction point should be adjusted, accordingly, after reference to distance from the extraction point (in simple cases). Therefore the air speed at the extraction, point should be adjusted, accordingly, after reference to distance from the extraction point (in simple cases). The air velocity at the extraction point (in simple cases). The air velocity at the extraction point of the mechanical considerations, producing performance deficits within the extraction apparatus, make it essential that theoretical air velocities are multiplied by factors of 10 or more when extraction systems are installed or used. Personal protection Safety glasses with unperforated side shields may be used where continuous eye protection is desirable, as in laboratories; spectacles are not sufficient where extraction of specific spectacles are not sufficient where extractions on paratus, make it as advance of the material coming in contax with the eyes; goggles must be properly filted. Eye and face protection Safety glasses with unperforated side shields may be used where continuous eye protection of eyes; these efford face protection. Hatematevia ga again sam king replace splash goggles and face shields. Contention of specific		3: Intermittent, low production. 3: High production, heavy use			
of distance from the extraction point (in simple cases). Therefore the air speed at the extraction point should be adjusted, accordingly, after reference to distance from the contaminating source. The air velocity at the extraction point. Other mechanical considerations, produing performance defaits within the extraction apparetus, make it essential that theoretical air velocities are multiplied by factors of 10 or more when extraction systems are installed or used. Personal protection Version Ver		4: Large hood or large air mass in motion 4: Small hood-local control only			
Eve and face protection Safety glasses with unperforated side shields may be used where continuous eye protection is desirable, as in laboratories; spectacles are not sufficient where complete eye protection is needed such as when handling bulk-quantities, where there is a danger of splashing, or if the material may be under pressure. Eye and face protection • Chemical goggles, whenever there is a danger of the material coming in contact with the eyes; goggles must be properly fitted. • Lift face shield (20 orn, 8 in minimum) may be required for supplementary but never for primary protection of eyes; these afford face protection. • Alternatively agas mask may replace splash goggles and face shields. • 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 in a clean environment only after workers have washed hands thoroughly. (CDC NIOSH Current Intelligence Bulletin 59). [AS/NZS 1336 or national equivalent] Skin protection See Hand protection below Hands/feet protection • Diversitions Overails. • Ver Apron. •		of distance from the extraction point (in simple cases). Therefore the air speed at the extraction point should be adjusted, accordingly, after reference to distance from the contaminating source. The air velocity at the extraction fan, for example, should be a minimum of 1-2 m/s (200-400 f/min) for extraction of solvents generated in a tank 2 meters distant from the extraction point. Other mechanical considerations, producing performance deficits within the extraction			
Eye and face protection See Hand protection below Eye and face protection See Hand protection below Exercise Current Intelligence Bulletin 59], [AS/NZS 1336 or national equivalent] Skin protection See Other protection below Other protection See Other protection below Description See Other protection below Eye and face protection See Other protection below	Personal protection				
Hands/feet protection Elbow length PVC gloves When handling corrosive liquids, wear trousers or overalls outside of boots, to avoid spills entering boots. Body protection See Other protection below Other protection PVC Apron. PVC Apron. PVC protective suit may be required if exposure severe. Eyewash unit. Ensure there is ready access to a safety shower. 	Eye and face protection	 where complete eye protection is needed such as when handling bulk-quantities, where there is a danger of splashing, or if the material may be under pressure. Chemical goggles.whenever there is a danger of the material coming in contact with the eyes; goggles must be properly fitted. Full face shield (20 cm, 8 in minimum) may be required for supplementary but never for primary protection of eyes; these afford face protection. Alternatively a gas mask may replace splash goggles and face shields. 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 			
Handsreet protection When handling corrosive liquids, wear trousers or overalls outside of boots, to avoid spills entering boots. Body protection See Other protection below Overalls. PVC Apron. PVC Apron. PVC protective suit may be required if exposure severe. Eyewash unit. Ensure there is ready access to a safety shower. 	Skin protection	See Hand protection below			
Other protection Overalls. PVC Apron. PVC protective suit may be required if exposure severe. Eyewash unit. Ensure there is ready access to a safety shower. 	Hands/feet protection	► Elbow length PVC gloves			
Other protection PVC Apron. PVC protective suit may be required if exposure severe. Eyewash unit. Ensure there is ready access to a safety shower. 	Body protection	See Other protection below			
Thermal hazards Not Available	Other protection	 Overalls. PVC Apron. PVC protective suit may be required if exposure severe. Eyewash unit. 			
	Thermal hazards	Not Available			

Respiratory protection

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

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Information on basic physical and chemical properties

Appearance	light grey		
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)	<2	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
Conditions to avoid	See section 7
Incompatible materials	See section 7
Hazardous decomposition products	See section 5

SECTION 11 TOXICOLOGICAL INFORMATION

Information on toxicological 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. 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 considered as immediately dangerous to life and health. It is estimated that the lowest lethal concentration for a 5-minute human exposure to hydrogen fluoride is in the range of 50 to 250 parts per million. Exposure by either skin contact or inhalation may lead to low levels of calcium and magnesium in the blood, which may result in heart rhythm disturbances. Animal testing suggests that repeated exposure produces liver and kidney damage.
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. 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. 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, incoordination and irritation of the mucous membranes. 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. 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. 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. 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.

Eye	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. 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 comea occurred within 1 hour, followed by tissue death (necrosis) of structures of the front of the eye.					
Chronic	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. 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, and inflammation of lung tissue often occurs. 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 in the mouth and throat and blood calcium levels are dangerously reduced.					
QCS-21	TOXICITY	IRRITATION				
	Not Available	Not Available				
	TOXICITY			IRRITATION		
antimony	Dermal (rabbit) LD50: >8300 mg/kg ^[1]			Not Available		
	Oral (rat) LD50: 100 mg/kg ^[2]					
		1				
arsenic	TOXICITY		IRRIT	ATION		
	Oral (rat) LD50: 763 mg/kg ^[2]		Not Av	vailable		
	TOXICITY	IRRITATION				
calcium carbonate	dermal (rat) LD50: >2000 mg/kg ^[1]	Eye (rabbit): 0.75 mg/24h - S	EVER	Ξ		
	Oral (rat) LD50: >2000 mg/kg ^[1]	Skin (rabbit): 500 mg/24h-mo	derate			
beryllium acetate	TOXICITY	IRRITATION				
Del ynium acetate	Not Available	Not Available				
cadmium	тохісіту			IRRITATION		
	Oral (rat) LD50: >63<259 mg/kg> ^[1]		Not Available			
chromium	TOXICITY	IRRITATION				
	Not Available Not Available					
	TOXICITY		IF	RRITATION		
cobalt	dermal (rat) LD50: >2000 mg/kg ^[1]		N	lot Available		
	Oral (rat) LD50: 6170 mg/kgd ^[2]					
				IRRITATION		
	dermal (rat) LD50: >2000 mg/kg ^[1]			Not Available		
copper	Inhalation (rat) LC50: 0.733 mg/l/4hr ^[1]					
	Inhalation (rat) LC50: 1.03 mg/l/4hr ^[1]					
	Inhalation (rat) LC50: 1.67 mg/l/4hr ^[1]					
	Oral (rat) LD50: 300-500 mg/kg ^[1]					
iron	TOXICITY	-	RRITATION			
	Oral (rat) LD50: 98600 mg/kg] ^[2]		Not	Available		
	TOXICITY			IRRITATION		
lead	dermal (rat) LD50: >2000 mg/kg ^[1]			Not Available		
	Inhalation (rat) LC50: >5.05 mg/l/4hr ^[1]					

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	Oral (rat) LD50: >2000 mg/kg ^[1]				
	TOXICITY		IRRITATION		
lithium carbonate	dermal (rat) LD50: >2000 mg/kg ^[1]		Eye (rabbit) : Mo	oderate *	
	Oral (rat) LD50: 525 mg/kg ^[2]		Skin (rabbit) : N	ild *	
	TOXICITY			IRRITATION	
magnesium	Oral (rat) LD50: >2000 mg/kg ^[1]			Not Available	
	ΤΟΧΙΟΙΤΥ			IRRITATION	
molybdenum	dermal (rat) LD50: >2000 mg/kg ^[1]			Not Available	
	Oral (rat) LD50: >2000 mg/kg ^[1]				
nickel				IRRITATION	
	Oral (rat) LD50: 5000 mg/kg ^[2]			Not Available	
	TOXICITY			IRRITATION	
selenium	Oral (rat) LD50: 6700 mg/kgd ^[2]			Not Available	
	тохісіту			IRRITATION	
strontium nitrate	Oral (rat) LD50: 1892 mg/kg ^[2]			Not Available	
thallium.	ΤΟΧΙΟΙΤΥ	IRRITA	TION		
thallium	Not Available	Not Av	ailable		
	тохісіту			IRRITATION	
titanium	Oral (rat) LD50: >2000 mg/kg ^[1]			Not Available	
				IRRITATION	
ammonium metavanadate	dermal (rat) LD50: 2102 mg/kg ^[2]	Not Available			
	Oral (rat) LD50: 160 mg/kgd ^[2]				
	TOXICITY			IRRITATION	
zinc	Dermal (rabbit) LD50: 1130 mg/kg ^[2]			Not Available	
	Oral (rat) LD50: >2000 mg/kg ^[1]				
	ΤΟΧΙΟΙΤΥ			IRRITATION	
nitric acid	Inhalation (rat) LC50: 625 ppm/1h*t ^[2]			Not Available	
			DDITATION		
			RRITATION	SE//EDE	
hydrofluoric acid	Inhalation (rat) LC50: 1276 ppm/4hr ^[2]		Eye (human): 50 mg -		
	Inhalation (rat) LC50: 319 ppm/1hr ^[2]				
water	ΤΟΧΙΟΙΤΥ	IRRITA	TION		
water	Not Available	Not Av	ailable		
manganese(II) acetate	ΤΟΧΙΟΙΤΥ			IRRITATION	
manganese(II) acetate tetrahydrate	Oral (rat) LD50: 3730 mg/kga ^[2]			Not Available	
Legend:	1. Value obtained from Europe ECHA Registered Substances - A	cute toxicity 2.* Value o	btained from manufa	cturer's SDS. Unless otherwise specified	d data
- U	extracted from RTECS - Register of Toxic Effect of chemical Subs				

Chemwatch: 9-254352 Catalogue number: QCS-21 Version No: 3.3

ARSENIC

Arsenic compounds are classified by the European Union as toxic by inhalation and ingestion and toxic to aquatic life and long lasting in the environment. Tumorigenic - Carcinogenic by RTECS criteria.

	Tumorigenic - Carcinogenic by RTECS criteria.					
CALCIUM CARBONATE	No evidence of carcinogenic properties. teratogenic effects.					
CHROMIUM	On skin and inhalation exposure, chromium and its compounds (except hexavalent) can be a potent sensitiser, as particulates. Tenth Annual Report on Carcinogens: Substance known to be Carcinogenic [<i>National Toxicology Program: U.S. Dep.</i> 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.					
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", an acute industrial disease of short duration. tiredness, influenza like respiratory tract irritation with fever.					
LEAD	WARNING: Lead is a cumulative poison and has the potential to cause about	ortion and intellectual in	npairment to unborn children of pregnant workers.			
LITHIUM CARBONATE	Goitrogenic: Goitrogens are substances that suppress the function of the thyroid gland h thyroid (a goitre). Lacrimation, altered sleep times, hallucinations, distorted perception, toxic sytemic administration), foetoxicity and foetolethality and specific develop	psychosis, excitement	ataxia, respiratory depression, allergic dermatitis (after			
NICKEL	Tenth Annual Report on Carcinogens: Substance anticipated to be Carcino [<i>National Toxicology Program: U.S. Dep.</i> Oral (rat) TDLo: 500 mg/kg/5D-I Inhalation (rat) TCLo: 0.1 mg/m3/24H/17	0				
THALLIUM	Structural changes in nerves and sheath, changes in extraocular muscles	, hair loss recorded				
NITRIC ACID	For acid mists, aerosols, vapours Test results suggest that eukaryotic cells are susceptible to genetic damag The material may cause severe skin irritation after prolonged or repeated e 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)					
ARSENIC & BERYLLIUM ACETATE	WARNING: This substance has been classified by the IARC as Group 1: CARCINOGENIC TO HUMANS.					
CALCIUM CARBONATE & BERYLLIUM ACETATE & LITHIUM CARBONATE & STRONTIUM NITRATE & AMMONIUM METAVANADATE & NITRIC ACID & HYDROFLUORIC ACID & MANGANESE(II) ACETATE TETRAHYDRATE	Asthma-like symptoms may continue for months or even years after exposure to the material ends.					
CALCIUM CARBONATE & NITRIC ACID & HYDROFLUORIC ACID	The material may produce severe irritation to the eye causing pronounced inflammation.					
CALCIUM CARBONATE & ZINC	The material may cause 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.					
BERYLLIUM ACETATE & COBALT & NICKEL	The following information refers to contact allergens as a group and may r	not be specific to this pr	oduct.			
CHROMIUM & MOLYBDENUM & TITANIUM & HYDROFLUORIC ACID & WATER	No significant acute toxicological data identified in literature search.					
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 28	3: Possibly Carcinogen	ic to Humans.			
NITRIC ACID & HYDROFLUORIC ACID	The material may produce respiratory tract irritation, and result in damage to the lung including reduced lung function.					
Acute Toxicity	¥	Carcinogenicity	\otimes			
Skin Irritation/Corrosion	· · · · · · · · · · · · · · · · · · ·	Reproductivity	<u> </u>			
Serious Eye Damage/Irritation		- Single Exposure	0			
Respiratory or Skin sensitisation	STOT - R	Repeated Exposure	0			
Mutagenicity	\otimes	Aspiration Hazard	\otimes			
		Legend:	 Data available but does not fill the criteria for classification Data available to make classification Data Not Available to make classification 			

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Toxicity

QCS-21	ENDPOINT TEST DURATION (HR)			SPECIES VALUE			SOURCE	
QU3-21	Not Applicable	Not Appli	cable		Not Applicable	Not Appli	cable	Not Applicable
	ENDPOINT	TEST DURAT	ION (HR)	SPEC	IES		VALUE	SOURCE
	LC50	96		Fish			0.93mg/L	2
	EC50	48			Crustacea			2
antimony	EC50	72			or other aquatic plants		1mg/L >2.4mg/L	2
	EC50	96		Crusta			0.5mg/L	2
	NOEC	720		Fish			>0.0075mg/	
	ENDPOINT	TEST DURAT	TION (HR)	SPE	CIES		VALUE	SOURCE
	LC50	96		Fish			9.9mg/L	4
arsenic	EC50	336		Alga	e or other aquatic plant	ts	0.63mg/L	4
	NOEC	336		-	e or other aquatic plant		<0.75mg/	
				`				
	ENDPOINT	TEST DURAT	ION (HR)	SPEC	CIES		VALUE	SOURCE
coloium cort and	LC50	96		Fish			>56000mg/	L 4
calcium carbonate	EC50	72		Algae	or other aquatic plants	5	>14mg/L	2
	NOEC	72		Algae	or other aquatic plants	3	14mg/L	2
herullium acetato	ENDPOINT	TEST DU	JRATION (HR)		SPECIES	VALUE		SOURCE
beryllium acetate	Not Applicable	Not Appli	cable		Not Applicable	Not Appli	cable	Not Applicable
	ENDPOINT	TEST DURATION (HR)		SPECIE	SPECIES		VALUE	SOURCI
	LC50	96		Fish	Fish		0.001mg/L	4
	EC50	48		Crustac	Crustacea		0.0033mg/L	5
cadmium	EC50	72		Algae o	Algae or other aquatic plants		0.018mg/L	2
	BCF	960		Fish	Fish		500mg/L	4
	EC50	336		Crustac	ea	(0.00065mg/L	5
	NOEC	168		Fish		().00001821mg/l	_ 4
	ENDPOINT	TEST DURAT	ION (HR)	SPEC	CIES		VALUE	SOURCE
	LC50	96		Fish		13.9mg/L	4	
	EC50	48		Crust	acea		0.0225mg/L	5
chromium	EC50	72		Algae	or other aquatic plants		0.104mg/L	4
	BCF	1440		Algae	or other aquatic plants		0.0495mg/L	4
	EC50	48		Crust	acea		0.0245mg/L	5
	NOEC	672		Fish			0.00019mg/	L 4
	ENDPOINT	TEST DURAT	ION (HR)	SPE	CIES		VALUE	SOURCE
	LC50	96		Fish			1.406mg/L	
	EC50	48		Crus	tacea		>0.89mg/L	
cobalt	EC50	72			e or other aquatic plants	S	0.144mg/L	
	BCF	1344			Fish		0.99mg/L	4
	EC50	70		Alga	Algae or other aquatic plants		0.02mg/L	2
	NOEC	168		Alga	e or other aquatic plant	S	0.0018mg/	L 2
	ENDPOINT	TEST DURAT	ION (HR)		SPECIES		VALUE	SOURCE
	LC50	96		Fish			0.0028mg/L	2
	EC50	48		Crusta			0.001mg/L	5
copper	EC50	72		Algae	or other aquatic plants		0.013335mg/l	
				Fish 200mg/L				
	BCF EC50	960 96		Fish Crusta			200mg/L 0.001mg/L	4

Legend:

Extracted from 1. IUCLID Toxicity Data 2. Europe ECHA Registered Substances - Ecotoxicological Information - Aquatic Toxicity 3. EPIWIN Suite V3.12 (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. 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 ocean, 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 the atmosphere range from 500 to several thousand years. Fluorise in aerosols can be transported over large distances by wind or as a result of atmospheric trubulence. 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
lithium carbonate	LOW	LOW
ammonium metavanadate	HIGH	HIGH
water	LOW	LOW

Bioaccumulative potential

Ingredient

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	1			
lithium carbonate	LOW (LogKOW = -0.4605)			
ammonium metavanadate	LOW (LogKOW = 2.229)			
water	LOW (LogKOW = -1.38)			
Mahility in sail				

Mobility in soil

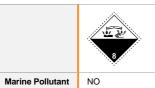
Ingredient	Mobility
lithium carbonate	HIGH (KOC = 1)
ammonium metavanadate	LOW (KOC = 35.04)
water	LOW (KOC = 14.3)

SECTION 13 DISPOSAL CONSIDERATIONS

aste treatment methods Product / Packaging disposal	 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. Recycle wherever possible. 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). Description on the follower of the product of the product.
	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. (contains nitric acid and hydrofluoric acid)
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	3264			
UN proper shipping name	Corrosive liquid, acidic	Corrosive liquid, acidic, inorganic, n.o.s. * (contains nitric acid and hydrofluoric acid)			fluoric acid)
Transport hazard class(es)	ICAO/IATA Class ICAO / IATA Subrisk ERG Code	8 Not Applicable 8L			
Packing group	Ш				
Environmental hazard	Not Applicable				
Special precautions for user	Special provisions Cargo Only Packing I Cargo Only Maximum			A3A803 855 30 L	

Passenger and Cargo Packing Instructions	851
Passenger and Cargo Maximum Qty / Pack	1 L
Passenger and Cargo Limited Quantity Packing Instructions	Y840
Passenger and Cargo Limited Maximum Qty / Pack	0.5 L

Sea transport (IMDG-Code / GGVSee)

UN number	3264
UN proper shipping name	CORROSIVE LIQUID, ACIDIC, INORGANIC, N.O.S. (contains nitric acid and hydrofluoric acid)
Transport hazard class(es)	IMDG Class8IMDG SubriskNot Applicable
Packing group	ll
Environmental hazard	Not Applicable
Special precautions for user	EMS NumberF-A, S-BSpecial provisions274Limited Quantities1 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

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)

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 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

CALCIUM CARBONATE(471-34-1) IS FOUND ON THE FOLLOWING REGULATORY LISTS

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

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 US ACGIH Threshold Limit Values (TLV) US ACGIH Threshold Limit Values (TLV) - Carcinogens 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) - Priority Pollutants US CWA (Clean Water Act) - Toxic Pollutants US EPCRA Section 313 Chemical List US National Toxicology Program (NTP) 14th Report Part A Known to be Human Carcinogens
- US NIOSH Recommended Exposure Limits (RELs) US OSHA Permissible Exposure Levels (PELs) - Table Z1
- US Toxic Substances Control Act (TSCA) Chemical Substance Inventory

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US - Alaska Limits for Air Contaminants	US - Rhode Island Hazardous Substance List
US - California Permissible Exposure Limits for Chemical Contaminants	US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants
US - Hawaii Air Contaminant Limits	US - Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminants
US - Idaho - Limits for Air Contaminants	US - Vermont Permissible Exposure Limits Table Z-1-A Transitional Limits for Air
US - Massachusetts - Right To Know Listed Chemicals	Contaminants
US - Michigan Exposure Limits for Air Contaminants	US - Washington Permissible exposure limits of air contaminants
US - Minnesota Permissible Exposure Limits (PELs)	US - Wyoming Toxic and Hazardous Substances Table Z1 Limits for Air Contaminants
US - Oregon Permissible Exposure Limits (Z-1)	US NIOSH Recommended Exposure Limits (RELs)
US - Pennsylvania - Hazardous Substance List	US OSHA Permissible Exposure Levels (PELs) - Table Z1
	US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory
BERYLLIUM ACETATE(543-81-7) IS FOUND ON THE FOLLOWING REGULATORY LIST	S
International Agency for Research on Cancer (IARC) - Agents Classified by the IARC	US - Washington Permissible exposure limits of air contaminants
Monographs	US - Washington Toxic air pollutants and their ASIL, SQER and de minimis emission values
US - Alaska Limits for Air Contaminants	US - Wyoming Toxic and Hazardous Substances Table Z1 Limits for Air Contaminants
US - California OEHHA/ARB - Chronic Reference Exposure Levels and Target Organs	US - Wyoming Toxic and Hazardous Substances Table Z-2 Acceptable ceiling concentration,
(CRELs)	Acceptable maximum peak above the acceptable ceiling concentration for an 8-hr shift
US - California Permissible Exposure Limits for Chemical Contaminants	US ACGIH Threshold Limit Values (TLV)
US - Hawaii Air Contaminant Limits	US ACGIH Threshold Limit Values (TLV) - Carcinogens
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 - Michigan Exposure Limits for Air Contaminants	US CWA (Clean Water Act) - Toxic Pollutants
US - Minnesota Permissible Exposure Limits (PELs)	US EPA Carcinogens Listing
US - Oregon Permissible Exposure Limits (Z-1)	US EPCRA Section 313 Chemical List
US - Oregon Permissible Exposure Limits (Z-2)	US National Toxicology Program (NTP) 14th Report Part A Known to be Human Carcinogens
US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants	US OSHA Permissible Exposure Levels (PELs) - Table Z1
US - Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminants	US OSHA Permissible Exposure Levels (PELs) - Table Z2
US - Vermont Permissible Exposure Limits Table Z-1-A Transitional Limits for Air	
International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Monographs US - Alaska 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 - California - Proposition 65 - Priority List for the Development of MADLs for Chemicals	US - Washington Permissible exposure limits of air contaminants
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 - Maximum Allowable Dose Levels (MADLs) for Chemicals	US ACGIH Threshold Limit Values (TLV) - Carcinogens
Causing Reproductive Toxicity	US ATSDR Minimal Risk Levels for Hazardous Substances (MRLs)
US - California Proposition 65 - No Significant Risk Levels (NSRLs) for Carcinogens	US Clean Air Act - Hazardous Air Pollutants
US - California Proposition 65 - Reproductive Toxicity	US CWA (Clean Water Act) - Priority Pollutants
US - Hawaii Air Contaminant Limits	US CWA (Clean Water Act) - Toxic Pollutants
US - Idaho - Acceptable Maximum Peak Concentrations	US EPA Carcinogens Listing
US - Idaho - Limits for Air Contaminants	US EPCRA Section 313 Chemical List
US - Massachusetts - Right To Know Listed Chemicals	US National Toxicology Program (NTP) 14th Report Part A Known to be Human Carcinogens
US - Michigan Exposure Limits for Air Contaminants	US NIOSH Recommended Exposure Limits (RELs)
US - Minnesota Permissible Exposure Limits (PELs)	US OSHA Carcinogens Listing
US - New Jersey Right to Know - Special Health Hazard Substance List (SHHSL):	US OSHA Permissible Exposure Levels (PELs) - Table Z1
Carcinogens	US OSHA Permissible Exposure Levels (PELs) - Table Z2
US - Oregon Permissible Exposure Limits (Z-1)	US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory
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	IS Verment Dermissible Experience Limits Table 7.4.4 Transitional Limits for Al-
International Agency for Research on Cancer (IARC) - Agents Classified by the IARC	US - Vermont Permissible Exposure Limits Table Z-1-A Transitional Limits for Air Contaminants
Monographs	
US - Alaska Limits for Air Contaminants	US - Washington Permissible exposure limits of air contaminants
US - California Permissible Exposure Limits for Chemical Contaminants	US - Wyoming Toxic and Hazardous Substances Table Z1 Limits for Air Contaminants
US - Hawaii Air Contaminant Limits	US ACGIH Threshold Limit Values (TLV)
US - Idaho - Limits for Air Contaminants	US ACGIH Threshold Limit Values (TLV) - Carcinogens
US - Massachusetts - Right To Know Listed Chemicals	US Clean Air Act - Hazardous Air Pollutants
US - Michigan Exposure Limits for Air Contaminants US - Oregon Permissible Exposure Limits (7-1)	US CWA (Clean Water Act) - Priority Pollutants
U.S CIEDOD PERMISSIDIE EXDOSURE LIMITS (Z-1)	

- US Michigan Exposure Limits for Air Contaminants
- 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

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

- 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

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International Agency for Research on Cancer (IARC) - Agents Classified by the IARC	LIC Versent Demissible Eveneure Limite Table 7.4.4 Transitional Limite for Air
Monographs	US - Vermont Permissible Exposure Limits Table Z-1-A Transitional Limits for Air Contaminants
US - Alaska Limits for Air 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 - California Proposition 65 - Carcinogens	US - Wyoming Toxic and Hazardous Substances Table Z1 Limits for Air Contaminants
US - Hawaii Air Contaminant Limits	US ACGIH Threshold Limit Values (TLV)
US - Idaho - Limits for Air Contaminants	US ACGIH Threshold Limit Values (TLV) - Carcinogens
US - Massachusetts - Right To Know Listed Chemicals	US ATSDR Minimal Risk Levels for Hazardous Substances (MRLs)
US - Michigan Exposure Limits for Air Contaminants	US Clean Air Act - Hazardous Air Pollutants
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 B.
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
US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants	Chemicals Causing Reproductive Toxicity
US - Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminants	US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory
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 - California Permissible Exposure Limits for Chemical Contaminants	US - Washington Permissible exposure limits of air contaminants
US - Hawaii Air Contaminant Limits	US - Washington Toxic air pollutants and their ASIL, SQER and de minimis emission values
US - Idaho - Limits for Air Contaminants	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 ATSDR Minimal Risk Levels for Hazardous Substances (MRLs)
US - Minnesota Permissible Exposure Limits (PELs)	US CWA (Clean Water Act) - Priority Pollutants
US - Oregon Permissible Exposure Limits (Z-1)	US CWA (Clean Water Act) - Toxic Pollutants
US - Pennsylvania - Hazardous Substance List	US EPA Carcinogens Listing
US - Rhode Island Hazardous Substance List	US EPCRA Section 313 Chemical List
US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants	US NIOSH Recommended Exposure Limits (RELs)
US - Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminants	US OSHA Permissible Exposure Levels (PELs) - Table Z1
	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 Parmissible Exposure Limite for Chamical Contaminante	, , , , , , , , , , , , , , , , , , , ,

- US California Permissible Exposure Limits for Chemical Contaminants
- 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 Monographs
- US Alaska Limits for Air Contaminants
- US California Proposition 65 Priority List for the Development of MADLs for Chemicals
- Causing Reproductive Toxicity
- US California Permissible Exposure Limits for Chemical Contaminants
- US California Proposition 65 Carcinogens
- US California Proposition 65 Maximum Allowable Dose Levels (MADLs) for Chemicals Causing Reproductive Toxicity
- US California Proposition 65 No Significant Risk Levels (NSRLs) for Carcinogens
- 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 Minnesota Permissible Exposure Limits (PELs)
- US New Jersey Right to Know Special Health Hazard Substance List (SHHSL): Carcinogens
- US Pennsylvania Hazardous Substance List
- US Rhode Island Hazardous Substance List

LITHIUM CARBONATE(554-13-2) IS FOUND ON THE FOLLOWING REGULATORY LISTS

- US California Proposition 65 Reproductive Toxicity
- US Massachusetts Right To Know Listed Chemicals
- US EPCRA Section 313 Chemical List

MAGNESIUM(7439-95-4) IS FOUND ON THE FOLLOWING REGULATORY LISTS

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

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

US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants

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

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

- US ACGIH Threshold Limit Values (TLV)
- US ACGIH Threshold Limit Values (TLV) Carcinogens
- 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 US EPCRA Section 313 Chemical List
- US National Toxicology Program (NTP) 14th Report Part B.
- US NIOSH Recommended Exposure Limits (RELs)
- US OSHA Permissible Exposure Levels (PELs) Table Z1
- US Toxic Substances Control Act (TSCA) Chemical Substance Inventory

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International Agency for Research on Cancer (IARC) - Agents Classified by the IARC	US - Pennsylvania - Hazardous Substance List
Monographs	US - Rhode Island Hazardous Substance List
US - California OEHHA/ARB - Chronic Reference Exposure Levels and Target Organs	US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants
(CRELs) US - California Permissible Exposure Limits for Chemical 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 Toxic Substances Control Act (TSCA) - Chemical Substance Inventory
US - Michigan Exposure Limits for Air Contaminants	
US - Oregon Permissible Exposure Limits (Z-1)	
MOLYBDENUM(7439-98-7) IS FOUND ON THE FOLLOWING REGULATORY LISTS	
US - Alaska Limits for Air Contaminants	US - Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminants
US - Hawaii Air Contaminant Limits	US - Vermont Permissible Exposure Limits Table Z-1-A Transitional Limits for Air
US - Idaho - Limits for Air Contaminants	Contaminants
US - Massachusetts - Right To Know Listed Chemicals	US - Washington Permissible exposure limits of air contaminants
US - Minnesota Permissible Exposure Limits (PELs)	US ACGIH Threshold Limit Values (TLV)
US - Pennsylvania - Hazardous Substance List	US ACGIH Threshold Limit Values (TLV) - Carcinogens
US - Rhode Island Hazardous Substance List	US NIOSH Recommended Exposure Limits (RELs)
US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants	US OSHA Permissible Exposure Levels (PELs) - Table Z1
	US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory
NICKEL(7440-02-0) 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 (REL	s) 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 ACGIH Threshold Limit Values (TLV)
US - California Proposition 65 - Carcinogens US - Hawaii Air Contaminant Limits	US ACGIH Threshold Limit Values (TLV) - Carcinogens US ATSDR Minimal Risk Levels for Hazardous Substances (MRLs)
US - Idaho - Limits for Air Contaminants	US Clean Air Act - Hazardous Air Pollutants
US - Massachusetts - Right To Know Listed Chemicals	US CWA (Clean Water Act) - Priority Pollutants
US - Michigan Exposure Limits for Air Contaminants	US CWA (Clean Water Act) - Toxic Pollutants
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 B.
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 Risl
US - Rhode Island Hazardous Substance List US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants	Levels (NSRLs) for Carcinogens and Maximum Allowable Dose Levels (MADLs) for Chemicals Causing Reproductive Toxicity
US - Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contamina	
SELENIUM(7782-49-2) IS FOUND ON THE FOLLOWING REGULATORY LISTS	
International Agency for Research on Cancer (IARC) - Agents Classified by the IARC	US - Washington Permissible exposure limits of air contaminants
Monographs US - Alaska Limits for Air Contaminants	US - Washington Toxic air pollutants and their ASIL, SQER and de minimis emission values
US - California OEHHA/ARB - Acute Reference Exposure Levels and Target Organs (REL	US ACGIH Threshold Limit Values (TLV) US ATSDR Minimal Risk Levels for Hazardous Substances (MRLs)
US - California OEHHA/ARB - Chronic Reference Exposure Levels and Target Organs	US Clean Air Act - Hazardous Air Pollutants
(CRELs)	US CWA (Clean Water Act) - Priority Pollutants
US - Hawaii Air Contaminant Limits	US CWA (Clean Water Act) - Toxic Pollutants
US - Idaho - Limits for Air Contaminants	US EPA Carcinogens Listing
US - Massachusetts - Right To Know Listed Chemicals	US EPCRA Section 313 Chemical List
US - Minnesota Permissible Exposure Limits (PELs)	US NIOSH Recommended Exposure Limits (RELs)
US - Pennsylvania - Hazardous Substance List	US OSHA Permissible Exposure Levels (PELs) - Table Z1
US - Rhode Island Hazardous Substance List 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	ints
US - Vermont Permissible Exposure Limits Table Z-1-A Transitional Limits for Air Contaminants	
	V LISTS
STRONTIUM NITRATE(10042-76-9) IS FOUND ON THE FOLLOWING REGULATOR	
US - Massachusetts - Right To Know Listed Chemicals	US EPCRA Section 313 Chemical List
US - Pennsylvania - Hazardous Substance List US - Rhode Island Hazardous Substance List	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 EPCRA Section 313 Chemical List
US - Pennsylvania - Hazardous Substance List US - Rhode Island Hazardous Substance List	US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory

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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 Fo	r Air Contaminants
International Air Transport Association (IATA) Dangerous Goods Regulations - Prohibited List	US - Washington Permissible exposure limits of air contar	
Passenger and Cargo Aircraft	US - Wyoming Toxic and Hazardous Substances Table Z1	
US - California OEHHA/ARB - Chronic Reference Exposure Levels and Target Organs (CRELs)	US Toxic Substances Control Act (TSCA) - Chemical Sub	stance Inventory
US - California Permissible Exposure Limits for Chemical Contaminants		
US - Hawaii Air Contaminant Limits		
US - Michigan Exposure Limits for Air Contaminants		
AMMONIUM METAVANADATE(7803-55-6) IS FOUND ON THE FOLLOWING REGULATORY	LISTS	
US - California OEHHA/ARB - Acute Reference Exposure Levels and Target Organs (RELs)	US EPCRA Section 313 Chemical List	
US - Massachusetts - Right To Know Listed Chemicals	US Toxic Substances Control Act (TSCA) - Chemical Sub	stance Inventory
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 Fo	r Air Contaminants
Monographs	US - Washington Permissible exposure limits of air contar	ninants
US - California OEHHA/ARB - Chronic Reference Exposure Levels and Target Organs	US - Wyoming Toxic and Hazardous Substances Table Z1	
(CRELs) US - California Permissible Exposure Limits for Chemical Contaminants	US ATSDR Minimal Risk Levels for Hazardous Substanc	es (MRLs)
US - Hawaii Air Contaminant Limits	US CWA (Clean Water Act) - Priority Pollutants	
US - Massachusetts - Right To Know Listed Chemicals	US CWA (Clean Water Act) - Toxic Pollutants US EPA Carcinogens Listing	
US - Michigan Exposure Limits for Air Contaminants	US EPCRA Section 313 Chemical List	
US - Oregon Permissible Exposure Limits (Z-1)	US Toxic Substances Control Act (TSCA) - Chemical Sub	stance Inventory
US - Pennsylvania - Hazardous Substance List		
US - Rhode Island Hazardous Substance List		
NITRIC ACID(7697-37-2) IS FOUND ON THE FOLLOWING REGULATORY LISTS		
International Air Transport Association (IATA) Dangerous Goods Regulations - Prohibited List	US - Vermont Permissible Exposure Limits Table Z-1-A Fi	nal Rule Limits for Air Contaminants
Passenger and Cargo Aircraft	US - Vermont Permissible Exposure Limits Table Z-1-A Tr	ransitional Limits for Air
US - Alaska Limits for Air Contaminants	Contaminants	ninente
US - California OEHHA/ARB - Acute Reference Exposure Levels and Target Organs (RELs) US - California Permissible Exposure Limits for Chemical Contaminants	US - Washington Permissible exposure limits of air contar US - Washington Toxic air pollutants and their ASIL, SQEF	
US - Hawaii Air Contaminant Limits	US - Wyoming Toxic and Hazardous Substances Table Z1	
US - Idaho - Limits for Air Contaminants	US ACGIH Threshold Limit Values (TLV)	
US - Massachusetts - Right To Know Listed Chemicals	US CWA (Clean Water Act) - List of Hazardous Substanc	es
US - Michigan Exposure Limits for Air Contaminants	US EPCRA Section 313 Chemical List	
US - Minnesota Permissible Exposure Limits (PELs)	US NIOSH Recommended Exposure Limits (RELs)	
US - Oregon Permissible Exposure Limits (Z-1) US - Pennsylvania - Hazardous Substance List	US OSHA Permissible Exposure Levels (PELs) - Table Z US SARA Section 302 Extremely Hazardous Substances	1
US - Rhode Island Hazardous Substance List	US Toxic Substances Control Act (TSCA) - Chemical Sub	stance Inventory
US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants		
HYDROFLUORIC ACID(7664-39-3) IS FOUND ON THE FOLLOWING REGULATORY LISTS International Agency for Research on Cancer (IARC) - Agents Classified by the IARC	US - Vermont Permissible Exposure Limits Table Z-1-A T	repetitional Limits for Air
Monographs	Contaminants	
US - Alaska Limits for Air Contaminants	US - Washington Permissible exposure limits of air contar	ninants
US - California OEHHA/ARB - Acute Reference Exposure Levels and Target Organs (RELs)	US - Washington Toxic air pollutants and their ASIL, SQEF	R and de minimis emission values
US - California OEHHA/ARB - Chronic Reference Exposure Levels and Target Organs	US - Wyoming Toxic and Hazardous Substances Table Z1	
(CRELs) US - California Permissible Exposure Limits for Chemical Contaminants	US - Wyoming Toxic and Hazardous Substances Table Z- Acceptable maximum peak above the acceptable ceiling co	1 0 ,
US - Hawaii Air Contaminant Limits	US ACGIH Threshold Limit Values (TLV)	
US - Idaho - Acceptable Maximum Peak Concentrations	US ACGIH Threshold Limit Values (TLV) - Carcinogens	
US - Idaho - Limits for Air Contaminants	US ATSDR Minimal Risk Levels for Hazardous Substanc	es (MRLs)
US - Massachusetts - Right To Know Listed Chemicals	US Clean Air Act - Hazardous Air Pollutants	
US - Michigan Exposure Limits for Air Contaminants	US CWA (Clean Water Act) - List of Hazardous Substanc	es
US - Minnesota Permissible Exposure Limits (PELs)	US EPCRA Section 313 Chemical List	
US - Oregon Permissible Exposure Limits (Z-1)	US NIOSH Recommended Exposure Limits (RELs)	1
US - Oregon Permissible Exposure Limits (Z-2) US - Pennsylvania - Hazardous Substance List	US OSHA Permissible Exposure Levels (PELs) - Table Z US OSHA Permissible Exposure Levels (PELs) - Table Z	
US - Rhode Island Hazardous Substance List	US SARA Section 302 Extremely Hazardous Substances	-
US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants	US Toxic Substances Control Act (TSCA) - Chemical Sub	stance Inventory
US - Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminants		
WATER(7732-18-5) IS FOUND ON THE FOLLOWING REGULATORY LISTS		

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

MANGANESE(II) ACETATE TETRAHYDRATE(6156-78-1) IS FOUND ON THE FOLLOWING REGULATORY LISTS

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US - Alaska Limits for Air Contaminants	US - Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminants
US - California OEHHA/ARB - Chronic Reference Exposure Levels and Target Organs (CRELs)	US - Vermont Permissible Exposure Limits Table Z-1-A Transitional Limits for Air Contaminants
US - California Permissible Exposure Limits for Chemical Contaminants	US - Washington Permissible exposure limits of air contaminants
US - Hawaii Air Contaminant Limits	US - Washington Toxic air pollutants and their ASIL, SQER and de minimis emission values
US - Idaho - Limits for Air Contaminants	US - Wyoming Toxic and Hazardous Substances Table Z1 Limits for Air Contaminants
US - Michigan Exposure Limits for Air Contaminants	US Clean Air Act - Hazardous Air Pollutants
US - Minnesota Permissible Exposure Limits (PELs)	US EPCRA Section 313 Chemical List
US - Oregon Permissible Exposure Limits (Z-1)	US OSHA Permissible Exposure Levels (PELs) - Table Z1
US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants	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 (lb)	Reportable Quantity in kg
Antimony	5000	2270
Arsenic	1	0.454
Cadmium	10	4.54
Chromium	5000	2270
Copper	5000	2270
Lead	10	4.54
Nickel	100	45.4
Selenium	100	45.4
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

Cadmium and cadmium compounds: Cadmium, Cobalt metal powder, Lead and lead compounds: Lead, Lithium carbonate, Nickel (Metallic) Listed

National Inventory	Status
Australia - AICS	N (beryllium acetate)
Canada - DSL	N (beryllium acetate)
Canada - NDSL	N (thallium; lead; zinc; lithium carbonate; strontium nitrate; ammonium metavanadate; titanium; magnesium; copper; water; antimony; selenium; molybdenum; arsenic; cobalt; nickel; manganese(II) acetate tetrahydrate; iron; chromium; hydrofluoric acid; beryllium acetate; cadmium; nitric acid)
China - IECSC	N (beryllium acetate)
Europe - EINEC / ELINCS / NLP	Y
Japan - ENCS	N (thallium; zinc; lithium carbonate; titanium; magnesium; copper; water; antimony; selenium; molybdenum; arsenic; cobalt; nickel; manganese(II) acetate tetrahydrate; iron; chromium; hydrofluoric acid; beryllium acetate; cadmium; nitric acid)
Korea - KECI	N (beryllium acetate)
New Zealand - NZIoC	N (beryllium acetate)
Philippines - PICCS	N (beryllium acetate)
USA - TSCA	N (beryllium acetate)
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

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Name	CAS No
calcium carbonate	471-34-1, 13397-26-7, 15634-14-7, 1317-65-3, 72608-12-9, 878759-26-3, 63660-97-9, 459411-10-0, 198352-33-9, 146358-95-4
copper	7440-50-8, 133353-46-5, 133353-47-6, 195161-80-9, 65555-90-0, 72514-83-1
strontium nitrate	10042-76-9, 13470-05-8
hydrofluoric acid	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 Chemwatch Classification committee using available literature references.

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

Definitions and abbreviations

PC-TWA: Permissible Concentration-Time Weighted Average

 ${\sf PC-STEL}: {\sf Permissible \ Concentration-Short \ Term \ Exposure \ Limit}$

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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