

ICP-MS-ICS-3B

High-Purity Standards

Version No: 1.1 Safety Data Sheet according to OSHA HazCom Standard (2012) requirements Chemwatch Hazard Alert Code: 3 Issue Date: 06/02/2017 Print Date: 06/02/2017

S.GHS.USA.EN

SECTION 1 IDENTIFICATION

Product Identifier

Product name	ICP-MS-ICS-3B
Synonyms	ICP-MS-ICS-3 Solution B
Proper shipping name	Corrosive liquid, acidic, inorganic, n.o.s. (contains nitric acid)
Other means of identification	ICP-MS-ICS-3 Solution B

Recommended use of the chemical and restrictions on use

Relevant identified uses Use according to manufacturer's directions.

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

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

Emergency phone number

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

SECTION 2 HAZARD(S) IDENTIFICATION

Classification	Metal Corrosion Category 1, Skin Corrosion/Irritation Category 1A		
abel elements			
Hazard pictogram(s)			
SIGNAL WORD	DANGER		
lazard statement(s)			
H290	May be corrosive to metals.		
H314	Causes severe skin burns and eye damage.		

Hazard(s) not otherwise specified

Not Applicable

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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.
Transutionary statement(a)	- Starage
Precautionary statement(s)	Storage
Precautionary statement(s) P405	Storage Store locked up.
	Store locked up.

Substances

See section below for composition of Mixtures

Mixtures

CAS No	%[weight]	Name
7697-37-2	2	nitric acid
7732-18-5	balance	water
7440-38-2	0.001	arsenic
7440-22-4	0.002	silver
7440-43-9	0.001	cadmium
7440-48-4	0.002	cobalt
7440-47-3	0.002	<u>chromium</u>
638-38-0	0.002 (as Mn)	manganese(II) acetate
7440-02-0	0.002	nickel
7803-55-6	0.002 (as V)	ammonium metavanadate
7440-66-6	0.001	zinc
7440-50-8	0.002	copper
7782-49-2	0.001	selenium

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	 If skin or hair contact occurs: Immediately flush body and clothes with large amounts of water, using safety shower if available. Quickly remove all contaminated clothing, including footwear. Wash skin and hair with running water. Continue flushing with water until advised to stop by the Poisons Information Centre. Transport to hospital, or doctor.
Inhalation	 If furnes or combustion products are inhaled remove from contaminated area. Lay patient down. Keep warm and rested. Prostheses such as false teeth, which may block airway, should be removed, where possible, prior to initiating first aid procedures. Apply artificial respiration if not breathing, preferably with a demand valve resuscitator, bag-valve mask device, or pocket mask as trained. Perform CPR if necessary. Transport to hospital, or doctor, without delay. Inhalation of vapours or aerosols (mists, fumes) may cause lung oedema. Corrosive substances may cause lung damage (e.g. lung oedema, fluid in the lungs). As this reaction may be delayed up to 24 hours after exposure, affected individuals need complete rest (preferably in semi-recumbent posture) and must be kept under medical observation even if no symptoms are (yet) manifested. Before any such manifestation, the administration of a spray containing a dexamethasone derivative or beclomethasone derivative may be considered. This must definitely be left to a doctor or person authorised by him/her. (ICSC13719)
Ingestion	 For advice, contact a Poisons Information Centre or a doctor at once. Urgent hospital treatment is likely to be needed. If swallowed do NOT induce vomiting. If vomiting occurs, lean patient forward or place on left side (head-down position, if possible) to maintain open airway and prevent aspiration. Observe the patient carefully. Never give liquid to a person showing signs of being sleepy or with reduced awareness; i.e. becoming unconscious. Give water to rinse out mouth, then provide liquid slowly and as much as casualty can comfortably drink. Transport to hospital or doctor without delay.

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See Section 11

Indication of any immediate medical attention and special treatment needed

For acute or short term repeated exposures to strong acids:

- Airway problems may arise from laryngeal edema and inhalation exposure. Treat with 100% oxygen initially.
- + Respiratory distress may require cricothyroidotomy if endotracheal intubation is contraindicated by excessive swelling
- Intravenous lines should be established immediately in all cases where there is evidence of circulatory compromise.

Strong acids produce a coagulation necrosis characterised by formation of a coagulum (eschar) as a result of the dessicating action of the acid on proteins in specific tissues. INGESTION:

- ▶ Immediate dilution (milk or water) within 30 minutes post ingestion is recommended.
- DO NOT attempt to neutralise the acid since exothermic reaction may extend the corrosive injury
- Be careful to avoid further vomit since re-exposure of the mucosa to the acid is harmful. Limit fluids to one or two glasses in an adult.
- Charcoal has no place in acid management.
- Some authors suggest the use of lavage within 1 hour of ingestion.

SKIN:

- Skin lesions require copious saline irrigation. Treat chemical burns as thermal burns with non-adherent gauze and wrapping.
- Deep second-degree burns may benefit from topical silver sulfadiazine.

EYE:

- Eye injuries require retraction of the eyelids to ensure thorough irrigation of the conjuctival cul-de-sacs. Irrigation should last at least 20-30 minutes. DO NOT use neutralising agents or any other additives. Several litres of saline are required.
- Cycloplegic drops, (1% cyclopentolate for short-term use or 5% homatropine for longer term use) antibiotic drops, vasoconstrictive agents or artificial tears may be indicated dependent on the severity of the injury.
- Steroid eye drops should only be administered with the approval of a consulting ophthalmologist).

[Ellenhorn and Barceloux: Medical Toxicology]

Both dermal and oral toxicity of manganese salts is low because of limited solubility of manganese. No known permanent pulmonary sequelae develop after acute manganese exposure. Treatment is supportive.

[Ellenhorn and Barceloux: Medical Toxicology]

In clinical trials with miners exposed to manganese-containing dusts, L-dopa relieved extrapyramidal symptoms of both hypo kinetic and dystonic patients. For short periods of time symptoms could also be controlled with scopolarnine and amphetamine. BAL and calcium EDTA prove ineffective.

[Gosselin et al: Clinical Toxicology of Commercial Products.]

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 furmes. May emit acrid smoke.

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

 b Drains for storage or use areas should have retention basins for pH adjustments and dilution of spills before discharge or disposal of material. b Check regularly for spills and leaks. b Clean up all spills immediately. b Avoid breathing vapours and contact with skin and eyes. b Control personal contact with the substance, by using protective equipment. b Contain and absorb spill with sand, earth, inert material or vermiculite. b Wipe up. b Place in a suitable, labelled container for waste disposal. 		
Major Spills #	Minor Spills	 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.
	Major Spills	#

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

SECTION 7 HANDLING AND STORAGE

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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 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. 		water and NEVER water to mate	erial.	

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

• Work clothes should be laundered separately. Launder contaminated clothing before re-use.

Observe manufacturer's storage and handling recommendations contained within this SDS.

Observe manufacturer's storage and handling recommendations contained within this SDS.

Always wash hands with soap and water after handling.

 Store away from incompatible materials and foodstuff containers. Protect containers against physical damage and check regularly for leaks.

• Use good occupational work practice.

 Store in original containers. Keep containers securely sealed. ▶ Store in a cool, dry, well-ventilated area.

Conditions for safe storage, including any incompatibilities

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Other information

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Suitable container	 DO NOT use aluminium or galvanised containers Check regularly for spills and leaks Lined metal can, lined metal pail/ can. Plastic pail. Polyliner drum. Packing as recommended by manufacturer. Check all containers are clearly labelled and free from leaks. For low viscosity materials Drums and jerricans must be of the non-removable head type. Where a can is to be used as an inner package, the can must have a screwed enclosure. For materials with a viscosity of at least 2680 cSt. (23 deg. C) and solids (between 15 C deg. and 40 deg C.): Removable head packaging; Cans with friction closures and low pressure tubes and cartridges may be used. Where combination packages are used, and the inner packages are of glass, porcelain or stoneware, there must be sufficient inert cushioning material in contact with inner and outer packages unless the outer packaging is a close fitting moulded plastic box and the substances are not incompatible with the plastic. 	
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 heat in small spaces. The dissolution of inorganic acids in water or the dilution of their concentrated solutions with additional water may generate significant heat. The addition of water to inorganic acids often generates sufficient heat in the small region of mixing to cause some of the water to boil explosively. The resulting "bumping" can spatter the acid. Inorganic acids react with active metals, including such structural metals as aluminum and iron, to release hydrogen, a flammable gas. Inorganic acids react with cyanide compounds to release gaseous hydrogen cyanide. Inorganic acids react with cyanide compounds to release gaseous hydrogen cyanide. Inorganic acids generate flammable and/or toxic gases in contact with dithiccarbamates, isocyanates, mercaptans, nitrides, nitriles, sulfides, and strong reducing agents. Additional gas-generating reactions occur with sulfites, nitrites, thiosulfates (to give H2S and SO3), dithionites (SO2), and even carbonates. Acids often catalyse (increase the rate of) chemical reactions. WARNING: Avoid or control reaction with peroxides. All <i>transition metal</i> peroxides should be considered as potentially explosive. For example transition meta complexes formed between chromium(0), vanadium(0) and other transition metals (haloarene-metal complexes) and mono-or poly-fluorobenzene show extreme sensitivity to heat and are explosive. Avoid reaction with borohydrides or cyanoborohydrides 	

SECTION 8 EXPOSURE CONTROLS / PERSONAL PROTECTION

Control parameters

OCCUPATIONAL EXPOSURE LIMITS (OEL)

INGREDIENT DATA

Source	Ingredient	Material name	TWA	STEL	Peak	Notes
US OSHA Permissible Exposure Levels (PELs) - Table Z1	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 furning nitric acid (RFNA), White furning nitric acid (WFNA)	5 mg/m3 / 2 ppm	4 ppm	Not Available	Not Available
US ACGIH Threshold Limit Values (TLV)	nitric acid	Nitric acid	2 ppm	Not Available	Not Available	Not Available
US 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)	silver	Silver metal: Argentum	0.01 mg/m3	Not Available	Not Available	Not Available

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sion No: 1.1								
US OSHA Permissible			0.005	Not	Net			
Exposure Levels (PELs) - lable Z1	cadmium	Cadmium	mg/m3	Available	Not Available	see 1910.1027;(as Cd)	
JS 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).]	
JS ACGIH Threshold Limit alues (TLV)	cadmium	Cadmium	Not Available	Not Available	Not Available	TLV® Basis: Kidney dam; BEI		
IS OSHA Permissible xposure Levels (PELs) - able Z1	cobalt	Cobalt metal, dust, and fume	0.1 mg/m3	Not Available	Not Available	(as Co)	(as Co)	
IS NIOSH Recommended Exposure Limits (RELs)	cobalt	Cobalt metal dust, Cobalt metal fume	0.05 mg/m3	Not Available	Not Available	TLV® Basis: Pneumonitis		
S ACGIH Threshold Limit alues (TLV)	cobalt	Hard metals containing Cobalt and Tungsten carbide, as Co	0.005 mg/m3	Not Available	Not Available	Not Available	Not Available	
IS NIOSH Recommended xposure Limits (RELs)	chromium	Chrome, Chromium	0.5 mg/m3	Not Available	Not Available	Not Available		
JS 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 do not apply to Nickel carbonyl.]		
JS ACGIH Threshold Limit /alues (TLV)	nickel	Nickel and inorganic compounds including Nickel subsulfide, as Ni - Elemental	1.5 mg/m3	Not Available	Not Available	TLV® Basis: Dermatitis; pneumoconios		
IS 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 Copp fume.]		
JS ACGIH Threshold Limit ′alues (TLV)	copper	Copper - Fume, as Cu	0.2 mg/m3	Not Available	Not Available	TLV® Basis: Irr; GI; metal fume fever; BI		
IS ACGIH Threshold Limit alues (TLV)	copper	Copper - Dusts and mists, as Cu	1 mg/m3	Not Available	Not Available	TLV® Basis: Irr; GI; metal fume fever; BE		
JS NIOSH Recommended Exposure Limits (RELs)	selenium	Elemental selenium, Selenium alloy	0.2 mg/m3	Not Available	Not Available	[*Note: The REL also a selenium compounds (Selenium hexafluoride	as Se) except	
MERGENCY LIMITS								
ngredient	Material nar	ne			TEEL-1	TEEL-2	TEEL-3	
tric acid	Nitric acid				Not Available	e Not Available	Not Available	
lver	Silver				0.3 mg/m3 170 mg/m3 990		990 mg/m3	
admium	Cadmium				Not Available	e Not Available	Not Available	
obalt	Cobalt				0.18 mg/m3 2 mg/m3		20 mg/m3	
hromium	Chromium				1.5 mg/m3 17 mg/m3		99 mg/m3	
anganese(II) acetate	Acetic acid, r	manganese(II) salt (2:1)			9.4 mg/m3 16 mg/m3		96 mg/m3	
ickel	Nickel				4.5 mg/m3	50 mg/m3	99 mg/m3	
mmonium metavanadate	Ammonium v	anadate; (Ammonium vanadium oxide; Ammonium n	netavanadate)		0.01 mg/m3	0.11 mg/m3	80 mg/m3	
inc	Zinc				6 mg/m3	21 mg/m3	120 mg/m3	
opper	Copper				3 mg/m3	33 mg/m3	200 mg/m3	
elenium	Selenium				0.6 mg/m3	6.6 mg/m3	40 mg/m3	
ngredient	Original IDL	H		Revis	ed IDLH			
itric acid	100 ppm			25 ppn	n			
vater	Not Available)		Not Av	vailable			
rsenic	100 mg/m3			5 mg/r	n3			
ilver	N.E. / N.E.			10 mg	/m3			
admium	50 mg/m3 / 9	mg/m3		9 mg/r	n3 / 9 [Unch] mg	/m3		
obalt	20 mg/m3			20 [Un	ich] mg/m3			
chromium	N.E. / N.E.			250 m	g/m3			
	IN.L. / IN.L.			230 11	9.110			

manganese(II) acetate

ammonium metavanadate

Exposure controls

nickel

zinc

copper

selenium

Engineering controls are used to remove a hazard or place a barrier between the worker and the hazard. Well-designed engineering controls can be highly effective in protecting workers and will typically be independent of worker interactions to provide this high level of protection.

500 mg/m3

10 mg/m3

Not Available

Not Available

100 mg/m3

1 mg/m3

Appropriate engineering controls

N.E. / N.E.

N.E. / N.E.

Not Available

Not Available

Unknown mg/m3 / Unknown ppm

N.E. / N.E.

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

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Event of approximation of the structure of		the particular process and chemical or contaminant in use. Employers may need to use multiple types of controls to prevent employee overexposure.			
Event Q25-0.5 m/s (60-100 brmin.) sectoret, vapours, degressing etc., eveporating from tank (in still air). Q25-0.5 m/s (60-100 brmin.) aerosols, turnes from pouring operations, intermittent container filling, low speed conveyer transfers, welding, spray drift, plaing Q15-1.7 m/s (100-200 trimin.) direct spray, spray, painting in stallow booths, drum filling, conveyer loading, cruster dusts, gas discharge (active generation into intrin.) Q15-10 m/s (500-200 trimin.) ginding, abravise blasting, turnbling, high speed wheel generated dusts (released at high initial velocity into zone of very high rapid generation doin). Q15-10 m/s (500-200 trimin.) Within each range the appropriate value depends or: Lower end of the range Upper end of the range 1. Boom air currents minimal of favourable to capture 1.0 float/bing room air currents 2. 2. Contaminants of how toxicity or of nuisance value only. 2. Contaminants of how toxicity 3. High production, heavy use 4. Large hood or large air mass in motion 4. Small hood-local control only 3. High production, heavy use of distance from the contraming gasars. The air wolcity fails rapidly with distance away from the astraction gine. Volcity generated distance from the extraction optic. The material concidues are motion at hard? needes data motion optic. The material concidues are multipled by factor of 10 or more when extraction popt. Volcity generated data we donot any and the extraction optic. The extrepolicity athe extraction optichastraction popt. Volcit		Supplied-air type respirator may be required in special circumstances. Correct fit is essential to ensu 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	re adequate protection. workplace possess varying "esca		
Event of approximation of the structure of		Type of Contaminant:		Air Speed:	
edd turnes, pickling (released at low velocity into zone of active generation) [trin,] direct spray, spray, parking in shallow boots, drum filling, conveyer loading, rusher dusts, gas discharge (active generation into min.) 1:2.5 m/s (200-500 (min.) girding, abrasive blasting, turnbling, high speed wheel generated dusts (released at high initial velocity into zone of very high rapid air motion). 25:10 m/s (500-2000 (min.) Within each range the appropriate value depends on: Upper end of the range 1:2.5 m/s (200-500 (min.) 1: Room air currents minimal or favourable to capture 1: Disturbing room air currents 2. Contaminants of high toxicity 2: Intermittent, low production. 3: High production, heavy use 4. Large hood or large air mass in motion 4: Small hood-local control only Simple theory shows that air velocity falls rapidly with distance away from the opening of a simple extraction pipe. Velocity generally decreases with the square of distance from the eortacino point. Other mechanical considerations, producing, speculared, accordingly, after reference to distance from the eortacino point. Other mechanical considerations, producing spectame, deficits within the extraction opiantaus, make the eostend in that theoretical air velocity at the estraction point. Other mechanical considerations, speculared, accordingly, after reference to distance from the contaminating source. The air velocity at the estraction point. Other mechanical considerations, producting systems are installed or used. Personal protection • Safety glasses with unpenforated side shields may be uade where continuous eye		solvent, vapours, degreasing etc., evaporating from tank (in still air).		0.25-0.5 m/s (50-100 f/min.)	
Even of rapid air motion) Image: An			, welding, spray drift, plating		
Personal protection 			charge (active generation into	,	
Even end of the range Upper end of the range 1: Room air currents minimal or favourable to capture 1: Disturbing room air currents 2: Contaminants of low toxicity or of nuisance value only. 2: Contaminants of high toxicity 3: Intermittent, low production. 3: High production, heavy use 4: Large hood or large air mass in motion 4: Small hood-local control only Simple theory shows that air velocity falls rapidly with distance away from the opening of a simple extraction pipe. Velocity generally decreases with the square of distance from the extraction point (ni simple cases). Therefore the air speed at the extraction point should be a dijusted, accordingly, after reference to distance from the extraction point. Other mechanical considerations, producing performance defaits within the extraction apparatus, make it essential that theoretical air velocity at the extraction point. Other mechanical considerations, producing performance defaits 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; spectades 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 Alternatively a gas mask may replace splash googies and face shields. Laring biold corm, B in minimum)			ocity into zone of very high rapid	2.5-10 m/s (500-2000 f/min.)	
I: Room air currents minimal or favourable to capture I: Disturbing room air currents I: Room air currents minimal or favourable to capture I: Disturbing room air currents I: Contaminants of low toxicity or of nuisance value only. I: Contaminants of high toxicity I: Intermittent, low production. I: High production, heavy use I: Large hood or large air mass in motion I: Small hood-local control only Simple theory shows that air velocity falls rapidly with distance away from the opening of a simple extraction ppic. Velocity generally decreases with the square of distance from the extraction point in the extraction point should be and usted, accordingly, after reference to distance from the extraction point. Other mechanical considerations, producing performance deficits within the extraction a sporatus, 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 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 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 prote		Within each range the appropriate value depends on:			
Event 2: Contaminants of low toxicity or of nuisance value only. 2: Contaminants of high toxicity 3: Intermittent, low production. 3: High production, heavy use 4: Large hood or large air mass in motion 4: Small hood-local control only Simple theory shows that air velocity falls rapidly with distance away from the opening of a simple extraction point, blevelog 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 1 of distance from the extraction point (in simple cases). Therefore the air speed at the extraction point should be adjusted, accordingly, after reference 1 of solvents generated in a tark 2 meters distant from the extraction point. Other mechanical considerations, producting 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 eve protection is desirable, as in laboratories; spectades are not sufficient necessarie. Eye and face protection Sinder y glasses with unperforated side shields may be used where continuous eve protection is desirable, as in laboratories; spectades are not sufficient pressure. A Chemical goggles. whenever there is a danger of the material coming in contact with the eyes; goggles must be properly fitted. Full face shield 2C or . Sin minimum miny may be required for supplementary but never for primary protection of eyes; these afford face protection.		Lower end of the range	Upper end of the range		
3: Intermittent, low production. 3: High production, heavy use 4: Large hood or large air mass in motion 4: Small hood-local control only Simple theory shows that air velocity falls rapidly with distance away from the opening of a simple extraction pipe. Velocity generally dacreases with the square of distance from the extraction point (in simple cases). Therefore the air speed at the extraction point should be adjusted, accordingly, after reference to distance from the contaminating source. The air velocity at the extraction form cmechanical 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 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 • Case in straigner of the material coming in contact with the eyes; goggles must be properly fitted. • Line as bind (20 cm. 8): minimum may be required for supplementary but never for primary protection of eyes; these afford face protection. • Atternatively ag as mask may replace splash goggles and face shields. • Contact lenses may pose a special hazard; soft contact lenses may abcord and concentrate initiants. A written policy document, describin for the class ochemicals in use and an account of injury sysperience. Medical affirst-aid personnel should be trained of the class so ochemi		1: Room air currents minimal or favourable to capture	1: Disturbing room air currents	1	
4: Large hood or large air mass in motion 4: Small hood-local control only Simple theory shows that air velocity falls rapidly with distance away from the opening of a simple extraction pipe. Velocity generally decreases with the square of distance from the contaminating source. The air velocity at the extraction point. Other mechanical considerations, producing performance deficits within the extraction apparatus, make it essential that theoretical air velocities are multiplied by factors of 10 or more when extraction systems are installed or used. Personal protection Safety glasses with upperforated 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 Cohemical 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 about do conclude are veive of lens absorption and adsorption for the class of definitions on use, should be created for each workplace or task. This should include a review of lens absorption and adsorption for the class of definitions on use, should be removed in a clean environment only after workers have washed hands thoroughy. [CDC NIC Current Intelligence Bulletin 59]. [ASNZS 1336 or national equivalent]		2: Contaminants of low toxicity or of nuisance value only.	2: Contaminants of high toxicity		
Even and face protection Safety glasses with unperforated side shields may be used where continuous ever protection is desirable, as in laboratories; spectacles are not sufficient. Even and face protection Safety glasses with unperforated side special special score of use, should be created for each workplace or task. This should include a review of lens absorption and adsorption for the class of a according the wearing on lenses or restrictions on use, should be created for each workplace or task. This should in their encover on the class of a according the wearing on the class of a according the wearing on the class of a distance from the extraction point. Other mechanical considerations, producing performance deficits within the extraction of solvents generated in a tank 2 meters distant from the extraction point. Other mechanical considerations, producing performance deficits within the extraction apparatus, make it essential that theoretical air velocities are multiplied by factors of 10 or more when extraction systems are installed or used. Personal protection Image: the science of the material coming in contact with the eyes; goggles must be properly fitted. Eye and face protection Safety glasses with unperforated side shields may be used where continuous even for primary protection of eyes; these afford face protection. Alternatively a gas mask may replace splash goggles and face shields. 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 bose a special hazard; soft contact lenses may absorb and concentrate iritiants. A written policy doc		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 fan, for example, should be a minimum of 1-2 m/s (200-400 f/min) for extraction of solvents generated in a tank 2 meters distant from the extraction point. Other mechanical considerations, producing performance deficits within the extraction apparatus, make it essential that theoretical air velocities are multiplied by factors of 10 or more when extraction systems are installed or used. Personal protection Image: Comparison of the extraction of solvents are installed or used. Selection solvents Image: Comparison of the extraction of the extraction of the extraction systems are installed or used. Personal protection Image: Comparison of the extraction systems are installed or used. Personal protection Image: Comparison of the extraction systems are installed or used. Personal protection Image: Selection 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 gogles.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 minim		4: Large hood or large air mass in motion	4: Small hood-local control only		
 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 or 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 or 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 NIC Current Intelligence Bulletin 59], [AS/NZS 1336 or national equivalent] 	Personal protection	of distance from the extraction point (in simple cases). Therefore the air speed at the extraction point s distance from the contaminating source. The air velocity at the extraction fan, for example, should be a solvents generated in a tank 2 meters distant from the extraction point. Other mechanical consideratio	should be adjusted, accordingly, a minimum of 1-2 m/s (200-400 f/m ns, producing performance deficit	fter reference to hin) for extraction of s within the extraction	
	Eye and face protection	 where complete eye protection is needed such as when handling bulk-quantities, where there is a pressure. Chemical goggles.whenever there is a danger of the material coming in contact with the eyes; go Full face shield (20 cm, 8 in minimum) may be required for supplementary but never for primary p 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 irrita lenses or restrictions on use, should be created for each workplace or task. This should include a chemicals in use and an account of injury experience. Medical and first-aid personnel should be treadily available. In the event of chemical exposure, begin eye irrigation immediately and remove at the first signs of eye redness or irritation - lens should be removed in a clean environment only a statement of the statement of the should be removed in a clean environment only a statement of the statement of the statement of the should be removed in a clean environment only a statement. 	a danger of splashing, or if the main oggles must be properly fitted. rotection of eyes; these afford fac ints. A written policy document, de review of lens absorption and ad rained in their removal and suitab contact lens as soon as practicabl	terial may be under e protection. escribing the wearing of isorption for the class of le equipment should be e. Lens should be remov	
	Skin protection	See Hand protection below			

Skin protection	See Hand protection below
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	 Overalls. PVC Apron. PVC protective suit may be required if exposure severe. Eyewash unit. Ensure there is ready access to a safety shower.
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)

SECTION 9 PHYSICAL AND CHEMICAL PROPERTIES

Information on basic physical and chemical properties

Appearance	Not Available		
Physical state	Liquid	Relative density (Water = 1)	Not Available
Odour	Not Available	Partition coefficient n-octanol / water	Not Available
Odour threshold	Not Available	Auto-ignition temperature (°C)	Not Available
pH (as supplied)	Not Available	Decomposition temperature	Not Available

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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	The material can cause respiratory irritation in some persons. The body's respondences concernsive acids can cause irritation of the respiratory tract, with coughing, choking nausea and weakness. The material has NOT been classified by EC Directives or other classification stanimal or human evidence.	ng and mucous membrane	damage. There r	may be dizziness, headache,	
Ingestion	Ingestion of acidic corrosives may produce burns around and in the mouth, the t speaking may also be evident. The material has NOT been classified by EC Directives or other classification s animal or human evidence. Poisonings rarely occur after oral administration of manganese salts because th	systems as "harmful by inge	stion". This is be	Ũ	
Skin Contact	Skin contact with acidic corrosives may result in pain and burns; these may be d Skin contact is not thought to have harmful health effects (as classified under Er through wounds, lesions or abrasions. 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, ma of the material and ensure that any external damage is suitably protected.	C Directives); the material m	nay still produce h	health damage following entry	
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.				
Chronic	Repeated or prolonged exposure to acids may result in the erosion of teeth, sw and inflammation of lung tissue often occurs. Long-term exposure to respiratory irritants may result in airways disease, involv Substance accumulation, in the human body, may occur and may cause some or Manganese is an essential trace element. Chronic exposure to low levels of ma slurred speech, disordered muscle tone, fatigue, anorexia, loss of strength and	ing difficulty breathing and ro oncern following repeated or inganese can include a mas	elated whole-bod long-term occup sk-like facial expre	dy problems. Dational exposure.	
	TOXICITY	IRRITATION			
ICP-MS-ICS-3B	Not Available	Not Available			
	TOXICITY		IRRITAT	ΓΙΟΝ	
nitric acid	Inhalation (rat) LC50: 625 ppm/1h*t ^[2]		Not Avai	ilable	
		1			
water	TOXICITY	IRRITATION			
	Not Available	Not Available			
arsenic	TOXICITY		IRRITATION		
	Oral (rat) LD50: 763 mg/kg ^[2]		Not Available		

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silver	TOXICITY		IRRITATION
	Oral (rat) LD50: >2000 mg/kg ^[1]		Not Available
cadmium	TOXICITY		IRRITATION
	Oral (rat) LD50: >63<259 mg/kg> ^[1]		Not Available
	ΤΟΧΙΟΙΤΥ		IRRITATION
cobalt	dermal (rat) LD50: >2000 mg/kg ^[1]		Not Available
	Oral (rat) LD50: 6170 mg/kgd ^[2]		
chromium	TOXICITY Nat Available	IRRITATION	
	Not Available	Not Available	
	ΤΟΧΙΟΙΤΥ		IRRITATION
manganese(II) acetate	Oral (rat) LD50: 2940 mg/kga ^[2]		Not Available
	Oral (ral) ED30. 2340 mg/kga		
	ΤΟΧΙΟΙΤΥ		IRRITATION
nickel	Oral (rat) LD50: 5000 mg/kg ^[2]		Not Available
	ΤΟΧΙΟΙΤΥ		IRRITATION
ammonium metavanadate	dermal (rat) LD50: 2102 mg/kg ^[2]		Not Available
	Oral (rat) LD50: 160 mg/kgd ^[2]		
	ΤΟΧΙΟΙΤΥ		IRRITATION
zinc	Dermal (rabbit) LD50: 1130 mg/kg ^[2]		Not Available
	Oral (rat) LD50: >2000 mg/kg ^[1]		
	ΤΟΧΙΟΙΤΥ		IRRITATION
	dermal (rat) LD50: >2000 mg/kg ^[1]		Not Available
	Inhalation (rat) LC50: 0.733 mg/l/4hr ^[1]		
copper	Inhalation (rat) LC50: 1.03 mg/l/4hr ^[1]		
	Inhalation (rat) LC50: 1.67 mg/l/4hr ^[1]		
	Oral (rat) LD50: 300-500 mg/kg ^[1]		
selenium	ΤΟΧΙΟΙΤΥ		IRRITATION
Scientum	Oral (rat) LD50: 6700 mg/kgd ^[2]		Not Available
Legend:	 Value obtained from Europe ECHA Registered Substances extracted from RTECS - Register of Toxic Effect of chemical S 		nufacturer's SDS. Unless otherwise specified data
	•		
	For acid mists, aerosols, vapours		
	Test results suggest that eukaryotic cells are susceptible to ge The material may produce severe irritation to the eye causing p		5.
NITRIC ACID	The material may produce respiratory tract irritation, and resul The material may cause severe skin irritation after prolonged o		
	vesicles, scaling and thickening of the skin.		
	Oral (?) LD50: 50-500 mg/kg * [Various Manufacturers] Arsenic compounds are classified by the European Union as t	oxic by inhalation and indestion and toxic to	aquatic life and long lasting in the environment
	WARNING: This substance has been classified by the IARC		
ARSENIC	were nunder uns substance has been classified by the IARC.	as GIUUD T. CARGINUGENIC TO HUMA	NO.
ARSENIC	Tumorigenic - Carcinogenic by RTECS criteria.		
ARSENIC		e to interactions between IgE antibodies an	nd allergens and occur rapidly.

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CHROMIUM	On skin and inhalation exposure, chromium and its compound Tenth Annual Report on Carcinogens: Substance known to b [<i>National Toxicology Program: U.S. Dep.</i> Gastrointestinal tumours, lymphoma, musculoskeletal tumour	e Carcinogenic	
MANGANESE(II) ACETATE	Laboratory tests have shown mutagenic effects: Positive B. re	ec.	
NICKEL	Tenth Annual Report on Carcinogens: Substance anticipated [<i>National Toxicology Program: U.S. Dep.</i> Oral (rat) TDLo: 500 mg/kg/5D-I Inhalation (rat) TCLo: 0.1 m	·	
ZINC	The material may cause skin irritation after prolonged or repe scaling and thickening of the skin.	ated exposure and may produce on	contact skin redness, swelling, the production of vesicles,
COPPER	for copper and its compounds (typically copper chloride): Acute toxicity: There are no reliable acute oral toxicity resul WARNING: Inhalation of high concentrations of copper fume like respiratory tract irritation with fever.		cute industrial disease of short duration. tiredness, influenza
NITRIC ACID & AMMONIUM METAVANADATE	Asthma-like symptoms may continue for months or even year	s after exposure to the material ends.	
WATER & CHROMIUM	No significant acute toxicological data identified in literature	search.	
COBALT & NICKEL	The following information refers to contact allergens as a gro	oup and may not be specific to this pr	oduct.
COBALT & NICKEL	WARNING: This substance has been classified by the IAR	C as Group 2B: Possibly Carcinogen	c to Humans.
CHROMIUM & SELENIUM	The substance is classified by IARC as Group 3: NOT classifiable as to its carcinogenicity to humans.		
Acute Toxicity	0	Carcinogenicity	0
Skin Irritation/Corrosion	 ✓ 	Reproductivity	\otimes
Serious Eye Damage/Irritation	0	STOT - Single Exposure	\odot
Respiratory or Skin sensitisation	\otimes	STOT - Repeated Exposure	\otimes
Mutagenicity	0	Aspiration Hazard	0
		v	– Data available but does not fill the criteria for classification – Data available to make classification – Data Not Available to make classification

SECTION 12 ECOLOGICAL INFORMATION

Toxicity ENDPOINT SPECIES VALUE TEST DURATION (HR) SOURCE ICP-MS-ICS-3B Not Applicable Not Applicable Not Applicable Not Applicable Not Applicable ENDPOINT TEST DURATION (HR) SPECIES SOURCE VALUE nitric acid NOEC 16 Crustacea 107mg/L 4 TEST DURATION (HR) SPECIES VALUE SOURCE ENDPOINT water Not Applicable Not Applicable Not Applicable Not Applicable Not Applicable ENDPOINT TEST DURATION (HR) SPECIES VALUE SOURCE LC50 96 Fish 9.9mg/L 4 arsenic EC50 336 4 Algae or other aquatic plants 0.63mg/L 4 NOEC 336 Algae or other aquatic plants <0.75mg/L ENDPOINT TEST DURATION (HR) SPECIES VALUE SOURCE LC50 0.00148mg/L 96 Fish 2 0.00024mg/L EC50 48 4 Crustacea EC50 96 0.001628837mg/L 4 silver Algae or other aquatic plants BCF 4 336 Crustacea 0.02mg/L EC50 48 0.00024mg/L 4 Crustacea NOEC 480 Crustacea 0.00031mg/L 2 TEST DURATION (HR) ENDPOINT SPECIES VALUE SOURCE cadmium LC50 96 Fish 0.001mg/L 4

Crustacea

48

EC50

Continued...

5

0.0033mg/L

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	1 1						1
	EC50	72		other aquatic plants		.018mg/L	2
	BCF	960	Fish			00mg/L	4
	EC50	336	Crustace	a	0	.00065mg/L	5
	NOEC	168	Fish		0	.00001821mg/L	4
	ENDPOINT	TEST DURATION (HR)	SPEC	IES		VALUE	SOURCE
	LC50	96	Fish			1.406mg/L	2
	EC50	48	Crust	2023		>0.89mg/L	2
cobalt	EC50	72		or other aquatic plant	re in the second se	0.144mg/L	2
CODAIL	BCF	1344	Fish		.5	0.144mg/L 0.99mg/L	4
		70		or other equatio plant		-	2
	EC50	168		or other aquatic plant		0.02mg/L	2
	NOEC	100	Aigae	or other aquatic plant	15	0.0018mg/L	2
	ENDPOINT	TEST DURATION (HR)	SPEC	IES		VALUE	SOURCE
	LC50	96	Fish			13.9mg/L	4
	EC50	48	Crusta	icea		0.0225mg/L	5
chromium	EC50	72	Algae	or other aquatic plants	3	0.104mg/L	4
	BCF	1440	-	or other aquatic plants		0.0495mg/L	4
	EC50	48	Crusta			0.0245mg/L	5
	NOEC	672	Fish			0.00019mg/L	4
						J	
	ENDPOINT	TEST DURATION (HR)		SPECIES	VALUE		SOURCE
manganese(II) acetate	Not Applicable			Not Applicable Not App		able	Not Applicable
	ENDPOINT	TEST DURATION (HR)	TEST DURATION (HR) SPECIES			VALUE	SOURCE
	LC50	96	Fish			0.0000475mg/L	4
	EC50	48	Crustac	Crustacea		0.013mg/L	5
nickel	EC50	72	Algae or other aquatic plants			0.0407mg/L	2
	BCF	1440	Algae o	Algae or other aquatic plants		0.47mg/L	4
	EC50	720	Crustac	Crustacea		0.0062mg/L	2
	NOEC	72	Algae o	r other aquatic plants		0.0035mg/L	2
	ENDPOINT	TEST DURATION (HR)	SPEC	IES		VALUE	SOURCE
	LC50	96	Fish			0.693mg/L	2
	EC50	48	Crust	acea		2.387mg/L	2
ammonium metavanadate	EC50	72		or other aquatic plant	ts	0.9894mg/L	2
	EC50	72		or other aquatic plant		1.162mg/L	2
	NOEC	72		or other aquatic plant		0.0168mg/L	2
							SOURCE
	ENDPOINT	TEST DURATION (HR)	SPEC	IES		VALUE	OCONOL
	ENDPOINT LC50	TEST DURATION (HR) 96	SPEC Fish	IES		VALUE 0.00272mg/L	4
zinc	LC50	96	Fish Crusta		S	0.00272mg/L	4
zinc	LC50 EC50	96 48	Fish Crusta Algae	acea		0.00272mg/L 0.04mg/L	4 5
zinc	LC50 EC50 EC50	96 48 72	Fish Crusta Algae	cea or other aquatic plants		0.00272mg/L 0.04mg/L 0.106mg/L	4 5 4
zinc	LC50 EC50 EC50 BCF	96 48 72 360	Fish Crusta Algae Algae Fish	cea or other aquatic plants	3	0.00272mg/L 0.04mg/L 0.106mg/L 9mg/L	4 5 4 4
zinc	LC50 EC50 EC50 BCF EC50	96 48 72 360 120	Fish Crusta Algae Algae Fish	icea or other aquatic plants or other aquatic plants	3	0.00272mg/L 0.04mg/L 0.106mg/L 9mg/L 0.00033mg/L	4 5 4 4 5 5
zinc	LC50 EC50 EC50 BCF EC50	96 48 72 360 120	Fish Crusta Algae Algae Fish	icea or other aquatic plants or other aquatic plants or other aquatic plants	3	0.00272mg/L 0.04mg/L 0.106mg/L 9mg/L 0.00033mg/L	4 5 4 4 5 5
zinc	LC50 EC50 EC50 BCF EC50 NOEC	96 48 72 360 120 336	Fish Crusta Algae Fish Algae	icea or other aquatic plants or other aquatic plants or other aquatic plants	3	0.00272mg/L 0.04mg/L 0.106mg/L 9mg/L 0.00033mg/L 0.00075mg/L	4 5 4 4 5 4
zinc	LC50 EC50 EC50 BCF EC50 NOEC ENDPOINT LC50 EC50	96 48 72 360 120 336 TEST DURATION (HR) 96 48	Fish Crusta Algae Fish Algae SPECI	icea or other aquatic plants or other aquatic plants or other aquatic plants ES	3	0.00272mg/L 0.04mg/L 0.106mg/L 9mg/L 0.00033mg/L 0.00075mg/L	4 5 4 4 5 4 4 5 4 SOURCE
zinc	LC50 EC50 EC50 BCF EC50 NOEC ENDPOINT LC50	96 48 72 360 120 336	Fish Crusta Algae Algae Fish Algae SPECI Fish Crusta	icea or other aquatic plants or other aquatic plants or other aquatic plants ES	5	0.00272mg/L 0.04mg/L 0.106mg/L 9mg/L 0.00033mg/L 0.00075mg/L	4 5 4 4 5 4 5 4 SOURCE 2
	LC50 EC50 EC50 BCF EC50 NOEC ENDPOINT LC50 EC50	96 48 72 360 120 336 TEST DURATION (HR) 96 48	Fish Crusta Algae Algae Fish Algae SPECI Fish Crusta	icea or other aquatic plants or other aquatic plants or other aquatic plants ES	5	0.00272mg/L 0.04mg/L 9mg/L 0.00033mg/L 0.00075mg/L VALUE 0.0028mg/L 0.001mg/L	4 5 4 4 5 4 5 4
	LC50 EC50 EC50 BCF EC50 NOEC ENDPOINT LC50 EC50 EC50	96 48 72 360 120 336 TEST DURATION (HR) 96 48 72	Fish Crusta Algae Fish Algae SPECI Fish Crusta Algae o	icea or other aquatic plants or other aquatic plants or other aquatic plants ES cea or other aquatic plants	5	0.00272mg/L 0.04mg/L 0.106mg/L 9mg/L 0.00033mg/L 0.00075mg/L VALUE 0.0028mg/L 0.001mg/L 0.013335mg/L	4 5 4 4 5 4 4

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	ENDPOINT	TEST DURATION (HR)	SPECIES	VALUE	SOURCE
	LC50	96	Fish	>0.0262mg/L	2
	EC50	48	Crustacea	>0.1603mg/L	2
selenium	EC50	72	Algae or other aquatic plants	>0.00173mg/L	2
	BCF	504	Crustacea	0.711mg/L	4
	EC50	96	Algae or other aquatic plants	0.355mg/L	2
	NOEC	72	Algae or other aquatic plants	0.000547mg/L	2
	NUEC	12	Aigae of other aquatic plants	0.000547 mg/L	2

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

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 Manganese and its Compounds:

Environmental Fate: Manganese is a naturally occurring element in the environment occurring as a result of weathering of geological material. It also occurs from its use in steel manufacture/ coal mining. The most commonly occurring of 11 possible oxidation states are +2, (e.g. manganese chloride or sulfate), +4, (e.g. manganese dioxide), and +7 (e.g. potassium permanganate), although the latter is unstable in the environment.

Atmospheric Fate: Elemental/inorganic manganese compounds may exist in air as suspended particulates from industrial emissions or soil erosion. Manganese-containing particles are mainly removed from the atmosphere by gravitational setting - large particles tend to fall out faster than small particles. The half-life of airborne particles is usually on the order of days, depending on the size of the particle and atmospheric conditions. Some removal by washout mechanisms such as rain may also occur, although it is of minor significance in comparison to dry deposition. Terrestrial Fate: Manganese in soil can migrate as particulate matter to air or water and soluble manganese compounds can be leached from the soil. High soil pH reduces manganese availability while low soil pH will increase availability, even to the point of toxicity. Soils high in organic matter Φ tie up Φ manganese such that high organic matter soils can be manganese to soil/sediments increases as positive ions increase, (cation), and organic matter increases. In some cases, adsorption of manganese to soils may not be a readily reversible process. At low concentrations, manganese may be fixed by clays and will not be released into solution readily. Bacteria and microflora can increase the mobility of manganese.

Aquatic Fate: Most manganese salts, with the exception of phosphates, carbonates, and oxides, are soluble in water. Solubility is controlled by the precipitation of insoluble forms, (species). In most oxygenated waters, the most common form is insoluble manganese oxide. Manganese chloride is the dominant form at pH 4-7, but may oxidize at pH>8 or 9.

Ecotoxicity: While lower organisms, (plankton, aquatic plants, and some fish), can significantly bioconcentrate manganese, higher organisms, (including humans), tend to maintain manganese balance. Manganese in water may be significantly concentrated at lower levels of the food chain.

Uptake of manganese by aquatic invertebrates and fish increases with temperature and decreases with pH. Fish and crustaceans appear to be the most sensitive to acute and chronic exposures. The substance has low toxicity to trout but, is moderately toxic to Coho salmon. The substance is toxic to Daphnia water fleas and moderately toxic to freshwater algae Pseudomonas putida and Photobacterium phosphoreum bacteria.

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.

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
water	LOW	LOW
ammonium metavanadate	HIGH	HIGH

Bioaccumulative potential

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

Mobility in soil

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

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SECTION 13 DISPOSAL CONSIDERATIONS

Waste treatment methods Product / Packaging disposal 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). 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



Marine Pollutant

Land transport (DOT)

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

Air transport (ICAO-IATA / DGR)

UN number	3264				
UN proper shipping name	Corrosive liquid, acidio	Corrosive liquid, acidic, inorganic, n.o.s. * (contains nitric 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	Passenger and Cargo Passenger and Cargo		A3A803 855 30 L 851 1 L Y840 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)		
Transport hazard class(es)	IMDG Class8IMDG SubriskNot Applicable		
Packing group	Ш		
Environmental hazard	Not Applicable		

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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

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 Final Rule Limits for Air Contaminants
Passenger and Cargo Aircraft	US - Vermont Permissible Exposure Limits Table Z-1-A Transitional Limits for Air
US - Alaska Limits for Air Contaminants	Contaminants
US - California OEHHA/ARB - Acute Reference Exposure Levels and Target Organs (RELs)	US - Washington Permissible exposure limits of air contaminants
US - California Permissible Exposure Limits for Chemical Contaminants	US - Washington Toxic air pollutants and their ASIL, SQER and de minimis emission values
US - Hawaii Air Contaminant Limits	US - Wyoming Toxic and Hazardous Substances Table Z1 Limits for Air Contaminants
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 Substances
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 OSHA Permissible Exposure Levels (PELs) - Table Z1
US - Pennsylvania - Hazardous Substance List	US SARA Section 302 Extremely Hazardous Substances
US - Rhode Island Hazardous Substance List	US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory
US - Tennessee Occupational Exposure Limits - 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
ARSENIC(7440-38-2) 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 - Washington Toxic air pollutants and their ASIL, SQER and de minimis emission values
US - Alaska Limits for Air Contaminants	US ACGIH Threshold Limit Values (TLV)
US - California OEHHA/ARB - Acute Reference Exposure Levels and Target Organs (RELs)	US ACGIH Threshold Limit Values (TLV) - Carcinogens
US - California OEHHA/ARB - Chronic Reference Exposure Levels and Target Organs	US ATSDR Minimal Risk Levels for Hazardous Substances (MRLs)
(CRELs)	US Clean Air Act - Hazardous Air Pollutants
US - California Permissible Exposure Limits for Chemical Contaminants US - Hawaii Air Contaminant Limits	US CWA (Clean Water Act) - Priority Pollutants
US - Idaho - Limits for Air Contaminants	US CWA (Clean Water Act) - Toxic Pollutants
US - Massachusetts - Right To Know Listed Chemicals	US EPCRA Section 313 Chemical List
US - Minnesota Permissible Exposure Limits (PELs)	US National Toxicology Program (NTP) 14th Report Part A Known to be Human Carcinogens
US - New Jersey Right to Know - Special Health Hazard Substance List (SHHSL):	US NIOSH Recommended Exposure Limits (RELs)
Carcinogens	US OSHA Permissible Exposure Levels (PELs) - Table Z1
US - Pennsylvania - Hazardous Substance List	US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory
US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants	
US - Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminants	
US - Vermont Permissible Exposure Limits Table Z-1-A Transitional Limits for Air	
Contaminants	
SILVER(7440-22-4) IS FOUND ON THE FOLLOWING REGULATORY LISTS	
US - Alaska Limits for Air Contaminants	US - Vermont Permissible Exposure Limits Table Z-1-A Transitional Limits for Air
US - California Permissible Exposure Limits for Chemical Contaminants	Contaminants
US - Hawaii Air Contaminant Limits	US - Washington Permissible exposure limits of air contaminants
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 CWA (Clean Water Act) - Priority Pollutants
US - Minnesota Permissible Exposure Limits (PELs)	US CWA (Clean Water Act) - Toxic Pollutants
US - Oregon Permissible Exposure Limits (Z-1)	US EPA Carcinogens Listing

- 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

CADMIUM(7440-43-9) IS FOUND ON THE FOLLOWING REGULATORY LISTS

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

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

US EPCRA Section 313 Chemical List

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International Agency for Research on Cancer (IARC) - Agents Classified by the IARC	US - Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminar
Monographs	US - Vermont Permissible Exposure Limits Table Z-1-A Transitional Limits for Air
US - Alaska Limits for Air Contaminants	Contaminants
US - California - Proposition 65 - Priority List for the Development of MADLs for Chemicals Causing Reproductive Toxicity	US - Washington Permissible exposure limits of air contaminants 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 (CRELs)	US - Wyoming Toxic and Hazardous Substances Table Z1 Limits for Air Contaminants 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 Causing Reproductive Toxicity	US ACGIH Threshold Limit Values (TLV) - Carcinogens 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 Carcinoger
US - Michigan Exposure Limits for Air Contaminants	US NIOSH Recommended Exposure Limits (RELs)
US - Minnesota Permissible Exposure Limits (PELs) US - New Jersey Right to Know - Special Health Hazard Substance List (SHHSL):	US OSHA Carcinogens Listing
Carcinogens	US OSHA Permissible Exposure Levels (PELs) - Table Z1
US - Oregon Permissible Exposure Limits (Z-1)	US OSHA Permissible Exposure Levels (PELs) - Table Z2
US - Oregon Permissible Exposure Limits (Z-2)	US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory
US - Pennsylvania - Hazardous Substance List	
US - Rhode Island Hazardous Substance List	
US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants	
COBALT(7440-48-4) IS FOUND ON THE FOLLOWING REGULATORY LISTS	
International Agency for Research on Cancer (IARC) - Agents Classified by the IARC 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 - Oregon Permissible Exposure Limits (Z-1)	US NIOSH Recommended Exposure Limits (RELs) 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 R
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
CHROMIUM(7440-47-3) IS FOUND ON THE FOLLOWING REGULATORY LISTS	
International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Monographs	US - 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 - 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 CWA (Clean Water Act) - Priority Pollutants US CWA (Clean Water Act) - Toxic Pollutants

US NIOSH Recommended Exposure Limits (RELs)

US Clean Air Act - Hazardous Air Pollutants

US EPCRA Section 313 Chemical List

US OSHA Permissible Exposure Levels (PELs) - Table Z1

US - Washington Permissible exposure limits of air contaminants

US OSHA Permissible Exposure Levels (PELs) - Table Z1

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

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

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

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

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

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

US EPCRA Section 313 Chemical List

Contaminants

- 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

MANGANESE(II) ACETATE(638-38-0) IS FOUND ON THE FOLLOWING REGULATORY LISTS

- US Alaska Limits for Air Contaminants
- US California OEHHA/ARB Chronic Reference Exposure Levels and Target Organs (CRELs)
- US California Permissible Exposure Limits for Chemical Contaminants
- US Hawaii Air Contaminant Limits
- US Idaho Limits for Air Contaminants
- US Michigan Exposure Limits for Air Contaminants
- US Minnesota Permissible Exposure Limits (PELs)
- US Oregon Permissible Exposure Limits (Z-1)
- US Tennessee Occupational Exposure Limits Limits For Air Contaminants

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

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rsion No: 1.1		
US - Alaska Limits for Air Contaminants	US - Vermont Permissible Exposure Limits Table Z-1	I-A Transitional Limits for Air
US - California OEHHA/ARB - Acute Reference Exposure Levels and Target Organs (RELs)	Contaminants	
US - California OEHHA/ARB - Chronic Reference Exposure Levels and Target Organs	US - Washington Permissible exposure limits of air c	
(CRELs)	US - Wyoming Toxic and Hazardous Substances Tab	ble 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 ACGIH Threshold Limit Values (TLV) - Carcinog	jens
US - Hawaii Air Contaminant Limits	US ATSDR Minimal Risk Levels for Hazardous Sub-	stances (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) - Tal	ble Z1
US - Pennsylvania - Hazardous Substance List	US Priority List for the Development of Proposition 6	5 Safe Harbor Levels - No Significant Risl
US - Rhode Island Hazardous Substance List	Levels (NSRLs) for Carcinogens and Maximum Allow	wable 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) - Chemica	I Substance Inventory
AMMONIUM METAVANADATE(7803-55-6) IS FOUND ON THE FOLLOWING REGULATOR	Y LISTS	
US - California OEHHA/ARB - Acute Reference Exposure Levels and Target Organs (RELs)	US EPCRA Section 313 Chemical List	
US - Massachusetts - Right To Know Listed Chemicals	US Toxic Substances Control Act (TSCA) - Chemica	Substance 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 - Lim	its For Air Contaminants
Monographs	US - Washington Permissible exposure limits of air c	
US - California OEHHA/ARB - Chronic Reference Exposure Levels and Target Organs	US - Wyoming Toxic and Hazardous Substances Tab	
(CRELs)	US ATSDR Minimal Risk Levels for Hazardous Substances has	
US - California Permissible Exposure Limits for Chemical Contaminants		stances (WIRES)
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 - Michigan Exposure Limits for Air Contaminants	US EPA Carcinogens Listing	
US - Oregon Permissible Exposure Limits (Z-1)	US EPCRA Section 313 Chemical List	
US - Pennsylvania - Hazardous Substance List	US Toxic Substances Control Act (TSCA) - Chemica	I Substance Inventory
US - Rhode Island Hazardous Substance List		
COPPER(7440-50-8) IS FOUND ON THE FOLLOWING REGULATORY LISTS		
	US - Vermont Permissible Exposure Limits Table Z-1	A Transitional Limits for Air
US - Alaska Limits for Air Contaminants	Contaminants	I-A Transitional Limits for All
US - California OEHHA/ARB - Acute Reference Exposure Levels and Target Organs (RELs)	US - Washington Permissible exposure limits of air c	ontominanto
US - California Permissible Exposure Limits for Chemical Contaminants		
US - Hawaii Air Contaminant Limits	US - Washington Toxic air pollutants and their ASIL,	
US - Idaho - Limits for Air Contaminants	US - Wyoming Toxic and Hazardous Substances Tab	ble 21 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 Sub	stances (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) - Tal US Toxic Substances Control Act (TSCA) - Chemica	
	03 Toxic Substances Control Act (TOCA) - Chemica	Substance inventory
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 c	
Monographs	US - Washington Toxic air pollutants and their ASIL,	SQER and de minimis emission values
US - Alaska Limits for Air Contaminants	US ACGIH Threshold Limit Values (TLV)	
US - California OEHHA/ARB - Acute Reference Exposure Levels and Target Organs (RELs)	US ATSDR Minimal Risk Levels for Hazardous Sub	stances (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) - Tal	ble Z1
US - Rhode Island Hazardous Substance List	US Toxic Substances Control Act (TSCA) - Chemica	
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		

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

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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
Nitric acid	1000	454
Arsenic	1	0.454
Silver	1000	454
Cadmium	10	4.54
Chromium	5000	2270
Nickel	100	45.4
Ammonium vanadate	1000	454
Zinc	1000	454
Copper	5000	2270
Selenium	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, Nickel (Metallic) Listed

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

SECTION 16 OTHER INFORMATION

Other information

Ingredients with multiple cas numbers

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

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

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

Definitions and abbreviations

PC – TWA: Permissible Concentration-Time Weighted Average PC – STEL: Permissible Concentration-Short Term Exposure Limit IARC: International Agency for Research on Cancer ACGIH: American Conference of Governmental Industrial Hygienists

STEL: Short Term Exposure Limit

TEEL: Temporary Emergency Exposure Limit。

IDLH: Immediately Dangerous to Life or Health Concentrations

OSF: Odour Safety Factor

NOAEL :No Observed Adverse Effect Level

LOAEL: Lowest Observed Adverse Effect Level

TLV: Threshold Limit Value

LOD: Limit Of Detection OTV: Odour Threshold Value

BCF: BioConcentration Factors

BEI: Biological Exposure Index

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Chemwatch: 9-405776 Catalogue number: ICP-MS-ICS-3 Solution B Version No: 1.1

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Issue Date: 06/02/2017 Print Date: 06/02/2017