

ICP-WS-2

High-Purity Standards

Catalogue number: ICP-WS-2

Version No: 3.3 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

i louuct identifier	
Product name	ICP-WS-2
Synonyms	50µg/mL Calcium, Magnesium, Potassium, Sodium 10µg/mL Boron, Barium, Bismuth, Cadmium, Cobalt, Chromium, Copper, Magnesium, Manganese, Nickel, Strontium, Vanadium in 2% HNO3
Proper shipping name	Corrosive liquid, acidic, inorganic, n.o.s. (contains nitric acid)
Other means of identification	ICP-WS-2

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 / C	Organisation	INFOTRAC
Emergenc	cy telephone numbers	1-800-535-5053
Other emergenc	y telephone numbers	1-352-323-3500

SECTION 2 HAZARD(S) IDENTIFICATION

Classification of the substance or mixture

Classification	Metal Corrosion Category 1, Skin Corrosion/Irritation Category 1A, Serious Eye Damage Category 1
Label elements	
Hazard pictogram(s)	
SIGNAL WORD	DANGER
Hazard 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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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)	Storage
P405	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
10022-31-8	0.001 (as Ba)	barium nitrate
7440-69-9	0.001	bismuth
10043-35-3	0.001 (as B)	boric acid
7440-43-9	0.001	cadmium
471-34-1	0.005 (as Ca)	calcium carbonate
7440-47-3	0.001	<u>chromium</u>
7440-48-4	0.001	cobalt
7440-50-8	0.001	copper
7439-95-4	0.005	magnesium
6156-78-1	0.001 (as Mn)	manganese(II) acetate tetrahydrate
7440-02-0	0.001	nickel
7757-79-1	0.005 (as K)	potassium nitrate
497-19-8	0.005 (as Na)	sodium carbonate
10042-76-9	0.001 (as Sr)	strontium nitrate
7803-55-6	0.001 (as V)	ammonium metavanadate
7697-37-2	2	nitric acid
7732-18-5	balance	water

SECTION 4 FIRST-AID MEASURES

Description of first aid measures

Eye Contact	If this product comes in contact with the eyes: Immediately hold eyelids apart and flush the eye continuously with running water. Ensure complete irrigation of the eye by keeping eyelids apart and away from eye and moving the eyelids by occasionally lifting the upper and lower lids. Continue flushing until advised to stop by the Poisons Information Centre or a doctor, or for at least 15 minutes. Transport to hospital or doctor without delay. Removal of contact lenses after an eye injury should only be undertaken by skilled personnel.
Skin Contact	 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 fumes or combustion products are inhaled remove from contaminated area. Lay patient down. Keep warm and rested. Prostheses such as false teeth, which may block airway, should be removed, where possible, prior to initiating first aid procedures. Apply artificial respiration if not breathing, preferably with a demand valve resuscitator, bag-valve mask device, or pocket mask as trained. Perform CPR if necessary. Transport to hospital, or doctor, without delay. Inhalation of vapours or aerosols (mists, fumes) may cause lung oedema. Corrosive substances may cause lung damage (e.g. lung oedema, fluid in the lungs). As this reaction may be delayed up to 24 hours after exposure, affected individuals need complete rest (preferably in semi-recumbent posture) and must be kept under medical observation even if no symptoms are (yet) manifested. Before any such manifestation, the administration of a spray containing a dexamethasone derivative or beclomethasone derivative may be considered. This must definitely be left to a doctor or person authorised by him/her. (ICSC13719)
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.
	Continued

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ent forward or place on left side (head-down position, if possible) to maintain y.	open airway and prevent aspiration.
n showing signs of being sleepy or with reduced awareness; i.e. becoming	
th, then provide liquid slowly and as much as casualty can comfortably drink. tor without delay.	
t	ICP-WS-2 ent forward or place on left side (head-down position, if possible) to maintain y. n showing signs of being sleepy or with reduced awareness; i.e. becoming th, then provide liquid slowly and as much as casualty can comfortably drink.

Most important symptoms and effects, both acute and delayed

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]

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.

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	#

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

SECTION 7 HANDLING AND STORAGE

Precautions for safe handling

Safe handling

Avoid all personal contact, including inhalation.

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	 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 scoap 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	ge, including any incompatibilities
Suitable container	 DO NOT use aluminium or galvanised containers Check regularly for spills and leaks Lined metal can, lined metal pail/ can. Plastic pail. Polyliner drum. Packing as recommended by manufacturer. Check all containers are clearly labelled and free from leaks. For low viscosity materials Drums and jerricans must be of the non-removable head type. Where a can is to be used as an inner package, the can must have a screwed enclosure. For materials with a viscosity of at least 2680 cSt. (23 deg. C) and solids (between 15 C deg. and 40 deg C.): Removable head packaging; Cans with friction closures and low pressure tubes and cartridges may be used. Where combination packages are used, and the inner packages are of glass, porcelain or stoneware, there must be sufficient inert cushioning material in contact with inner and outer packages unless the outer packaging is a close fitting moulded plastic box and the substances are not incompatible with the plastic.
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 flammable and/or toxic gases in contact with dithiocarbamates, isocyanates, mercaptans, nitrides, nitriles, sulfides, and strong reducing agents. Additional gas-generating reactions occur with sulfites, nitrites, thiosulfates (to give H2S and SO3), dithionites (SO2), and even carbonates. Acids often catalyse (increase the rate of) chemical reactions. WARNING: Avoid or control reaction with peroxides. All <i>transition meta</i>l peroxides should be considered as potentially explosive. For example transition metal complexes of alkyl hydroperoxides may decompose explosively. The pi-complexes formed between chromium(0), vanadium(0) and other transition metals (haloarene-metal complexes) and mono-or poly-fluorobenzene show extreme sensitivity to heat and are explosive.

SECTION 8 EXPOSURE CONTROLS / PERSONAL PROTECTION

Control parameters

OCCUPATIONAL EXPOSURE LIMITS (OEL)

INGREDIENT DATA

Source	Ingredient	Material name	TWA	STEL	Peak	Notes
US NIOSH Recommended Exposure Limits (RELs)	barium nitrate	Barium dinitrate, Barium(II) nitrate (1:2), Barium salt of nitric acid	0.5 mg/m3	Not Available	Not Available	[*Note: The REL also applies to other soluble barium compounds (as Ba) except Barium sulfate.]
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)	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

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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 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)	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 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

EMERGENCY LIMITS

Ingredient	Material name		TEEL-1	TEEL-2	TEEL-3
barium nitrate	Barium nitrate		2.9 mg/m3	350 mg/m3	2,100 mg/m3
bismuth	Bismuth		15 mg/m3	170 mg/m3	990 mg/m3
boric acid	Boric acid		6 mg/m3	23 mg/m3	830 mg/m3
cadmium	Cadmium		Not Available	Not Available	Not Available
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
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
magnesium	Magnesium		18 mg/m3	200 mg/m3	1,200 mg/m3
manganese(II) acetate tetrahydrate	Acetic acid, manganese(2+) salt, tetrahydrate		13 mg/m3	22 mg/m3	740 mg/m3
manganese(II) acetate tetrahydrate	Acetic acid, manganese(II) salt (2:1)		9.4 mg/m3	16 mg/m3	96 mg/m3
nickel	Nickel		4.5 mg/m3	50 mg/m3	99 mg/m3
potassium nitrate	Potassium nitrate		9 mg/m3	100 mg/m3	600 mg/m3
sodium carbonate	Sodium carbonate		7.6 mg/m3	83 mg/m3	500 mg/m3
strontium nitrate	Strontium nitrate		5.7 mg/m3	62 mg/m3	370 mg/m3
ammonium metavanadate	Ammonium vanadate; (Ammonium vanadium oxide; Ammonium metava	nadate)	0.01 mg/m3	0.11 mg/m3	80 mg/m3
nitric acid	Nitric acid		Not Available	Not Available	Not Available
Ingredient	Original IDLH	Revised IDLH			
barium nitrate	1,100 mg/m3	50 mg/m3			
bismuth	Not Available	Not Available			
boric acid	Not Available	Not Available			

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cadmium	50 mg/m3 / 9 mg/m3	9 mg/m3 / 9 [Unch] mg/m3
calcium carbonate	Not Available	Not Available
chromium	N.E. / N.E.	250 mg/m3
cobalt	20 mg/m3	20 [Unch] mg/m3
copper	N.E. / N.E.	100 mg/m3
magnesium	Not Available	Not Available
manganese(II) acetate tetrahydrate	N.E. / N.E.	500 mg/m3
nickel	N.E. / N.E.	10 mg/m3
potassium nitrate	Not Available	Not Available
sodium carbonate	Not Available	Not Available
strontium nitrate	Not Available	Not Available
ammonium metavanadate	Not Available	Not Available
nitric acid	100 ppm	25 ppm
water	Not Available	Not Available

Exposure controls

Appropriate engineering The of Contaminant		Engineering controls are used to remove a hazard or place a barrier between the worker and the haza effective in protecting workers and will typically be independent of worker interactions to provide this hig 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 th "removes" air in the work environment. Ventilation can remove or dilute an air contaminant if designed p 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. Cor Supplied-air type respirator may be required in special circumstances. Correct fit is essential to ensure 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 w turn, determine the "capture velocities" of fresh circulating air required to effectively remove the contami	gh level of protection. The worker and ventilation that stra properly. The design of a ventilation prect fit is essential to obtain adeq re adequate protection. workplace possess varying "esca	tegically "adds" and on system must match uate protection.					
Appropriate engineering 0.25-0.5 m/s (20-100 tmm.) account, vapours, degressing etc., evaporating from tark (in still air). 0.25-0.5 m/s (20-100 tmm.) account, vapours, degressing etc., evaporating from tark (in still air). 0.25-0.5 m/s (20-200 tmm.) account, vapours, degressing etc., evaporating from tark (in still air). 0.25-0.5 m/s (20-200 tmm.) account, vapours, degressing etc., evaporating from tark (in still air). 0.25-0.5 m/s (20-200 tmm.) grading abraive blasting, umbing in haldw booths, drum filing, conveyer backing, crusher dusts, gas dscharge (active generation into 1.25 m/s (20-200 tmm.) 1.25 m/s (20-200 tmm.) grading abraive blasting, umbing in haldw booths, drum filing, conveyer backing, crusher dusts (released at high initial velocity into zone of very high rapid at motion). 2.5-10 m/s (50-200 tmm.) With each range Upper end of the range 2.5-10 m/s (50-200 tmm.) 1: Room air ourrents minimal of 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, two production. 3: High production heavy shows that air velocity falls rapid/will will distance away from the opering of a simple correction opti. 4: Small hood-local control only Strugt how oy shows that air velocity falls rapid/will will distance away from the opering of a simple correction post. 4: Large hood or large air mass in motion				Air Speed:					
Appropriate engineering controls and funces, picking (released at low velocity into zone of active generation) (fmin.) drect spray, spray painting in shallow bodhs, drum filling, conveyer loading, crusher dusts, gas discharge (active generation into into active discrete spray) 1-25 m/s (200-500 (min.) grinding, abrasive blasting, tumbling, high speed wheel generated dusts (released at high initial velocity into zone of very high rapid ir motion). 25-10 m/s (500-2000 (min.) With each range the appropriate value depends on: Upper end of the range 1 Lower end of the range 1.0 Elstarting room air currents 2 2. Contaminants of two toxicity or of nuisance value only. 2. Contaminants of high toxicity 3 3. Intermittent, low production. 4. Signal hood-local control only. 3 Stringe heavy, show that at velocity falls rapidly with distance away from the opering of a simple extraction pixt boxid heavy shows that at velocity at the extraction opint. Other mechanical consideration pixt for mix (200-00 (min.)) for extraction of obserts 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 at velocities are nulpiled by factors of 10 or mee when extraction systems are installed or used. Personal protection • Singly glasses with unperforated side shick may be used where continuous eye protection is desirable, as in laboratorine; spectacles are not sufficient where complete eye protection is needed such				0.25-0.5 m/s (50-100					
Identicity spray, spray painting in shalow boots, drum htting, conveyer foading, drucher dusts, gas discharge (active generation into 12-25 ms (200-900 tmm), grinding, abrasive blasting, turbling, high speed wheel generated dusts (released at high initial velocity into zone of very high rapid 25-10 m/s (500-2000 tmm), grinding, abrasive blasting, turbling, high speed wheel generated dusts (released at high initial velocity into zone of very high rapid 25-10 m/s (500-2000 tmm), grinding, abrasive blasting, turbling, high speed wheel generated dusts (released at high initial velocity into zone of very high rapid 25-10 m/s (500-2000 tmm), grinding, abrasive blasting, turbling at the appropriate value depends on: Uwer end of the range Upper end of the range 2-10 m/s (500-2000 tmm), grinding, abrasive blasting, turbling in structures in the appropriate value depends on: I. Room air currents minimal of favorusble to capture 1: Disturbing room air currents 2. Contaminants of high toxidity 3: Intermittent, low production. 3: High production, heavy use 4: Large hood or large air mass in motion 4: Small hood-local control only Struct P heory shows that air velocity falls rapidly with distance away from the extraction points shuld be adjusted, acconding, after reference to distance from the contaminants groups. Therefore the air speed at the extraction points shuld be adjusted, acconding, with the extraction apparatus, make It essential that theoretical air velocity at the extraction for the extraction systems are installed or used. Personal protection • Safety glasses with unperforated side shelichs may be used where continuous eye protection is desirable, as in laboratories; spectades are not sufficient w			welding, spray drift, plating						
Bit motion). It min.) Image: the appropriate value depends on: Lower end of the range Upper end of the range 1: Boom 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 heavy shows that air velocity falls rapidly with distance away from the opening of a simple 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 contaminant gource. The air velocity at the extraction function considerations, producting performance deficits within 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 (in the ext	controls		charge (active generation into						
Lower end of the range Upper end of the range 1: Boarn 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 point should be adjusted, accordingly, after reference to distance from the extraction point is found to adjusted, accordingly, after reference to distance from the extraction point should be adjusted, accordingly, after reference to distance from 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 uperforated side shields may be used where continuous eye protection is desirable, as in laboratories; spectades are not sufficient pressure. • Chemical goggles.whenever there is a danger of the material corning in contact with the eyes; goggles must be properly fitted. • Aulf face protection • Safety glasses with uperforated side shields may be used where continuous eye protection of eyes; these afford face protection. • Abernical goggles.whenever there is a danger of the material corning in contact with the eyes; goggles must be properly fitted. <t< td=""><th></th><td colspan="5"></td></t<>									
I: Room air currents minimal of favourable to capture I: Disturbing room air currents I: Room air currents minimal of favourable to capture I: Disturbing room air currents I: Contaminants of low toxicity or of nuisance value only. I: Contaminants of high toxicity I: Itage hood or large air mass in motion I: Bigh 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 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 a divisued, accordingly, after reference to distance from the extraction point (in simple cases). Therefore the air speed at the extraction point should be a minimum of 1-2 ms (200-d00 trimi) for extraction of aparatus, 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,		Within each range the appropriate value depends on:							
Eventaminants of low toxicity or divisance 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. 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 extrantianting source. The air velocity at the extraction point. Other mechanical considerations, producing performance deficits within the extraction aparatus, 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 of eyes; these aftor face protection. • Large biologies whenever there is a danger of the material coming in contact with the eyes; goggles must be properly fitted. • Full face protection • Contact lenses may may reas a special hazard; soft contact lenses may absoch and concentrate infrants. A written policy document, describing the wearing of chemical spouse, begin yee		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 decreases with the square of distance from the extraction point (in simple cases). Therefore the air speed at the extraction point should be adjusted, accordingly, after reference to distance from the contaminating source. The air velocity at the extraction point on thou the adjusted, accordingly, after reference to 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 Importance deficits within the extraction factors of 10 or more when extraction systems are installed or used. Life protection Importance 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. Life protection Importance deficits 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. Lift face shield (20, m, 8 in minimum) may be required for supplementary but never for primary protection of eyes; these afford face protection. High hood-local control or injury approace splashing on the cases of chemical syn top space as a protical hard may be under pressure. Lorenical		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 falls rapidly with distance away from the opening of a simple extraction pipe. Velocity generally decreases with the square of distance from the extraction point in simple cases). Therefore the air speed at the extraction point should be adjusted, accordingly, after reference to distance from the contaminating source. The air velocity at the extraction point should be a minimum of 1-2 m/s (200-400 t/min) for extraction of solvenits 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 complete expression of 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 complete expression. • Safety glasses with upperforated side shields may be used where continuous expertection is desirable, as in laboratories; spectacles are not sufficient where complete expression. • Lemical googles.whenever there is a danger of the material complit in contact with the eyes; goggles must be properly fitted. • Chemical googles.whenever there is a danger of primary protection of eyes; these afford face protection. • Atternatively ag amask may replace splash goggles and face shields. • Contact lienese may pose a special hazard; soft contact wi		2: Contaminants of low toxicity or of nuisance value only.	2: Contaminants of high toxicity						
Eye and face 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 Safety glasses with unperforated side shields may be required for supplementary but never for spinary protection of eyes; these alford face protection. Full face protection Vietnatively a gas mask may replace splash goggles and face shields. Contact lenses or prestrictions on use, should be remeded for each workplace or task. This should include a review of lens absorption and adsorption for the deas of chemicage or task. This should include a review of lens absorption and adsorption for the deas of chemicage explash end of the remedel and first-aid personnel should be trained in their removal and suitable equipment should be trained in their removal and suitable equipment should be trained in their removal and suitable equipment should be trained in their removal and suitable equipment should be remove at the first signs of eye rentees or irritation - lens should be removed in a clean environment only after workers have washed hands thoroughly. [CDC NIOSH Current Intelligence Bulletin 59], [AS/NZS 1336 or national equivalent]		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 fmin) 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 the extraction of the extraction systems are installed or used. Personal protection Image: Comparison of the extraction of the extraction of the extraction 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 Full face shield (20 orm, 8 in minimum) may be required for supplementary but never for primary protection of eyes; these alford face protection. Alternatively a gas mask may replace splash goggles and face shields. Contact lenses may pascrift, 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 removeed in a clean environment only after workers have washed hands		4: Large hood or large air mass in motion	4: Small hood-local control only						
Eye and face protection Alternatively a gas mask may replace splash goggles and face shields. 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 removed in a clean environment only after workers have washed hands thoroughly. [CDC NIOSH Current Intelligence Bulletin 59], [AS/NZS 1336 or national equivalent]		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 consideration	hould 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 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. 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 Current Intelligence Bulletin 59], [AS/NZS 1336 or national equivalent] 	Personal protection								
Skin protection See Hand protection below	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 prevailed (20 cm, 8 in minimum) may be required for supplementary but never for primary prevails and face shields. Contact lenses may pose a special hazard; soft contact lenses may absorb and concentrate irritar 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 travel available. In the event of chemical exposure, begin eye irrigation immediately and remove c at the first signs of eye redness or irritation - lens should be removed in a clean environment only a 	danger of splashing, or if the mat ggles must be properly fitted. otection of eyes; these afford face nts. A written policy document, de review of lens absorption and ad ained in their removal and suitabl ontact lens as soon as practicable	erial may be under e protection. scribing the wearing of sorption for the class of le equipment should be e. Lens should be removed					
	Skin protection	See Hand protection below							

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Hands/feet protection	Elbow length PVC glovesWhen handling corrosive liquids, wear	r trousers or overalls outside of boots, to avoid spills	s entering boots.
Body protection	See Other protection below		
Other protection	 Overalls. PVC Apron. PVC protective suit may be required if Eyewash unit. Ensure there is ready access to a safe 		
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	colorless		
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	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. The material has NOT been classified by EC Directives or other classification systems as "harmful by inhalation". This is because of the lack of corroborating animal or human evidence.
Ingestion	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. The material has NOT been classified by EC Directives or other classification systems as "harmful by ingestion". This is because of the lack of corroborating animal or human evidence. Ingestion or skin absorption of boric acid causes nausea, abdominal pain, diarrhoea and profuse vomiting which may be blood stained, headache, weakness, reddened lesions on the skin. In severe cases, it may cause shock, with fall in blood pressure, increase in heart rate, blue skin colour, brain and nervous irritation, reduced urine volume or even absence of urine. Borate poisoning causes nausea, vomiting, diarrhoea and pain in the upper abdomen. Often persistent vomiting occurs, and there may be blood in the facees.
Skin Contact	Skin contact with acidic corrosives may result in pain and burns; these may be deep with distinct edges and may heal slowly with the formation of scar tissue. Skin contact is not thought to have harmful health effects (as classified under EC Directives); the material may still produce health damage following entry through wounds, lesions or abrasions. Boric acid is not absorbed via intact skin but absorbed on broken or inflamed skin.

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	Open cuts, abraded or irritated skin should not be exposed to this r Entry into the blood-stream, through, for example, cuts, abrasions of of the material and ensure that any external damage is suitably pro	or lesions, ma	y produce systemic injury with	n harmf	ul effects. Examine the skin prior to the use
Eye	If applied to the eyes, this material causes severe eye damage. Direct eye contact with acid corrosives may produce pain, tears, s completely.	sensitivity to lig	ght and burns. Mild burns of th	ne epith	elia generally recover rapidly and
Chronic	Repeated or prolonged exposure to acids may result in the erosio and inflammation of lung tissue often occurs. Long-term exposure to respiratory irritants may result in airways di Substance accumulation, in the human body, may occur and may o Chronic boric acid poisoning is characterized by mild gastrointest irregular and discoloured rash. Dryness of skin, reddening of tongu Borate can accumulate in the testes and deplete germ cells and o stomach ulcer and anaemia can all occur.	lisease, involvi cause some co tinal irritation, ue, loss of hair	ing difficulty breathing and rel oncern following repeated or la loss of appetite, disturbed dig r, inflammation of conjunctiva,	ated wh ong-terr gestion, , and kie	nole-body problems. m occupational exposure. nausea, possibly vomiting and a hard dney injury have also been reported.
			IDDITATION		
ICP-WS-2	TOXICITY Not Available		IRRITATION Not Available		
			Not Available		
	TOYICITY	100			
h				oto	
barium nitrate	Oral (rat) LD50: 355 mg/kgd ^[2]		e (rabbit):100 mg/24h - moder	ate	
		SKI	n (rabbit): 500 mg/24h - mild		
bismuth					ATION
	Oral (rat) LD50: 2000 mg/kg ^[1]			NOT AV	vailable
			IDDITATION		
boric acid	TOXICITY IRRITATION				
	Oral (rat) LD50: 2500 mg/kg ^[2] Skin (human): 15 mg/3d -I- mild			nild	
cadmium					RRITATION
	Oral (rat) LD50: >63<259 mg/kg> ^[1]			r	Not Available
calcium carbonate	dermal (rat) LD50: >2000 mg/kg ^[1]		Eye (rabbit): 0.75 mg/24h - SB		
	Oral (rat) LD50: >2000 mg/kg ^[1]		Skin (rabbit): 500 mg/24h-moo	derate	
			IRRITATION		
chromium	TOXICITY Not Available		Not Available		
			Not Available		
	ΤΟΧΙΟΙΤΥ			IR	RITATION
cobalt	dermal (rat) LD50: >2000 mg/kg ^[1]				ot Available
	Oral (rat) LD50: 6170 mg/kgd ^[2]				
	ΤΟΧΙΟΙΤΥ				IRRITATION
	dermal (rat) LD50: >2000 mg/kg ^[1]				Not Available
	Inhalation (rat) LC50: 0.733 mg/l/4hr ^[1]				
copper	Inhalation (rat) LC50: 1.03 mg//4fr ^[1]				
	Inhalation (rat) LC50: 1.03 mg//4nr - Inhalation (rat) LC50: 1.67 mg//4nr ^[1]				
	Oral (rat) LD50: 300-500 mg/kg ^[1]				
					TATION
magnesium					
	Oral (rat) LD50: >2000 mg/kg ^[1]			Not /	Available
manganese(II) acetate tetrahydrate					TATION
tetranyurate	Oral (rat) LD50: 3730 mg/kga ^[2]			Not A	Available

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	L. C.						
	ΤΟΧΙΟΙΤΥ			IRF	RITATION		
nickel	Oral (rat) LD50: 5000 mg/kg ^[2]				Not Available		
	ΤΟΧΙΟΙΤΥ				IRRITATION		
potassium nitrate	dermal (rat) LD50: >5000 mg/kg ^[1]				Not Available		
	Oral (rat) LD50: >2000 mg/kg ^[1]						
	ΤΟΧΙΟΙΤΥ		IRRITATION				
	dermal (rat) LD50: >2000 mg/kg*E ^[2]		Eye (rabbit): 100 mg/24h	mod	lerate		
sodium carbonate	Oral (rat) LD50: 2800 mg/kg*d ^[2]		Eye (rabbit): 100 mg/30s				
			Eye (rabbit): 50 mg SEVI				
			Skin (rabbit): 500 mg/24h				
	TOXICITY			IRF	RITATION		
strontium nitrate	Oral (rat) LD50: 1892 mg/kg ^[2]			Not	Not Available		
	тохісіту				RRITATION		
ammonium metavanadate	dermal (rat) LD50: 2102 mg/kg ^[2]			r	Not Available		
	Oral (rat) LD50: 160 mg/kgd ^[2]						
	ΤΟΧΙΟΙΤΥ				IRRITATION		
nitric acid	Inhalation (rat) LC50: 625 ppm/1h*t ^[2]				Not Available		
			IDDITATION				
water	TOXICITY Not Available		IRRITATION Not Available				
Legend:	1. Value obtained from Europe ECHA Registered Substances - Acute toxic extracted from RTECS - Register of Toxic Effect of chemical Substances	ty 2.*	Value obtained from manufa	actui	rer's SDS. Unless otherwise specified date		
BARIUM NITRATE	The material may produce moderate eye irritation leading to inflammation.						
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. The substance is classified by IARC as Group 3: NOT classifiable as to its carcinogenicity to humans. 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. Excogenous allergic alveolitis is induced essentially by allergen specific immune-complexes of the IgG type; cell-mediated reactions (T lymphocytes) may be imported.

involved. for copper and its compounds (typically copper chloride): Acute toxicity: There are no reliable acute oral toxicity results available. COPPER 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. Tenth Annual Report on Carcinogens: Substance anticipated to be Carcinogen NICKEL [National Toxicology Program: U.S. Dep. Oral (rat) TDLo: 500 mg/kg/5D-I Inhalation (rat) TCLo: 0.1 mg/m3/24H/17W-C For sodium carbonate: SODIUM CARBONATE Sodium carbonate has little potential for skin irritation, but is irritating to the eyes. For acid mists, aerosols, vapours Test results suggest that eukaryotic cells are susceptible to genetic damage when the pH falls to about 6.5. The material may produce respiratory tract irritation, and result in damage to the lung including reduced lung function. NITRIC ACID The material may cause severe skin irritation after prolonged or repeated exposure and may produce on contact skin redness, swelling, the production of vesicles, scaling and thickening of the skin. Oral (?) LD50: 50-500 mg/kg * [Various Manufacturers] **BARIUM NITRATE & BORIC** ACID & CALCIUM The material may cause skin irritation after prolonged or repeated exposure and may produce on contact skin redness, swelling, the production of vesicles, **CARBONATE & SODIUM** scaling and thickening of the skin. CARBONATE

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CALCIUM CARBONATE & MANGANESE(II) ACETATE TETRAHYDRATE & SODIUM CARBONATE & STRONTIUM NITRATE & AMMONIUM METAVANADATE & NITRIC ACID	Asthma-like symptoms may continue for months or even years	after exposure to the material ends.					
CALCIUM CARBONATE & NITRIC ACID	The material may produce severe irritation to the eye causing pronounced inflammation.						
CHROMIUM & WATER	No significant acute toxicological data identified in literature search.						
COBALT & NICKEL	The following information refers to contact allergens as a group and may not be specific to this product.						
COBALT & NICKEL	WARNING: This substance has been classified by the IARC as Group 2B: Possibly Carcinogenic to Humans.						
Acute Toxicity	\otimes	Carcinogenicity	\otimes				
Skin Irritation/Corrosion	×	Reproductivity	0				
Serious Eye Damage/Irritation	✓	STOT - Single Exposure	0				
Respiratory or Skin sensitisation	0	STOT - Repeated Exposure	0				
Mutagenicity	0	Aspiration Hazard	0				
		0	 Data available but does not fill the criteria for classification Data available to make classification 				

Data available to make classification

 \bigcirc – Data Not Available to make classification

SECTION 12 ECOLOGICAL INFORMATION

	ENDPOINT TEST DURATION (H		TEST DURATION (HR)) SPECIES		VALUE	VALUE		RCE
ICP-WS-2	Not Applicable Not Applicable			Not Applicable	Not Applica	ble	Not A	Applicable	
	ENDPOINT	TE	TEST DURATION (HR)		SPECIES		VALUE		SOURCE
barium nitrate	LC50	96		Fis	Fish		>3.5mg/L		2
	EC50	72		Alg	Algae or other aquatic plants		>1.92mg/L		2
	EC50	72		Alg	Algae or other aquatic plants		>34.31mg/L	-	2
	NOEC	72		Alg	ae or other aquatic plants	3	>=1.92mg/l	L	2
			1		1				
bismuth	ENDPOINT		TEST DURATION (HR)		SPECIES	VALUE		SOURCE	
	Not Applicable		Not Applicable		Not Applicable	Not Applica	ble	Not A	Applicable
	ENDPOINT	TE	TEST DURATION (HR)		SPECIES		VALUE		SOURCE
	LC50	96		F	Fish		74mg/L		2
	EC50	48		С	Crustacea		133mg/L		4
boric acid	EC50	72		A	gae or other aquatic plan	ts	54mg/L		2
	EC50	72		A	Algae or other aquatic plants		66mg/L		2
	NOEC	768			sh		0.009mg/	Ľ	2
	ENDPOINT	TEST DURATION (HR)		SPEC	IES	VA	LUE		SOURCE
	LC50	96	96		Fish 0				4
	EC50	48	48		Crustacea).0033mg/L		5
cadmium	EC50	72		Algae	Algae or other aquatic plants 0		0.018mg/L		2
	BCF	960		Fish	Fish		500mg/L		4
	EC50	336		Crust	Crustacea)0065mg/L		5
	NOEC	168		Fish	Fish 0.0)0001821mg/L	-	4
	ENDPOINT		ST DURATION (HR)		SPECIES		VALUE		SOURCE
calcium carbonate	LC50	96		Fis			>56000mg/l	<u> </u>	4
	EC50	72		Alg	ae or other aquatic plants	5	>14mg/L		2

	ENDPOINT	TEST DURATION (HR)	SPECIES		VA	ALUE	SOURCE
	LC50	96	Fish	Fish		3.9mg/L	4
	EC50	48	Crustacea	Crustacea		0225mg/L	5
chromium	EC50	72	Algae or other aqua	Algae or other aquatic plants		104mg/L	4
	BCF	1440		Algae or other aquatic plants		0495mg/L	4
	EC50	48	Crustacea			0245mg/L	5
	NOEC	672	Fish			00019mg/L	4
		012	1 1311		0.0	000 TBITIg/L	4
	ENDPOINT	TEST DURATION (HR)	SPECIES		1	/ALUE	SOURCE
				Fish			
	LC50	96				1.406mg/L	2
	EC50	48	Crustacea			>0.89mg/L	2
cobalt	EC50	72	Algae or other aqu	atic plants).144mg/L	2
	BCF	1344	Fish).99mg/L	4
	EC50	70	Algae or other aqu	atic plants	0).02mg/L	2
	NOEC	168	Algae or other aqua	atic plants	0).0018mg/L	2
			00000				0011707
	ENDPOINT	TEST DURATION (HR)	SPECIES			LUE	SOURCE
	LC50	96	Fish			028mg/L	2
	EC50	48	Crustacea			01mg/L	5
copper	EC50	72	Algae or other aquat	ic plants	0.0	13335mg/L	4
	BCF	960	Fish		200)mg/L	4
	EC50	96	Crustacea		0.00	01mg/L	5
	NOEC	96	Crustacea	Crustacea		008mg/L	4
	ENDPOINT TEST DURATION (HR)		SPECIES			VALUE	SOURCE
	LC50	96	Fish			541mg/L	2
magnesium	EC50	72	Algae or other aqu	atic plants		>20mg/L	2
	EC50	72	Algae or other aqu	latic plants		>20mg/L	2
	NOEC	72	Algae or other aqu	atic plants		>25.5mg/L	2
manganese(II) acetate	ENDPOINT TEST DURATION (HR)		SPECIES	VAL	UE	S	OURCE
tetrahydrate	Not Applicable	Not Applicable	Not Applica	ole Not A	Applicable	N	lot Applicable
	ENDPOINT	TEST DURATION (HR)	SPECIES		VAL		SOURCE
	LC50	96	Fish			00475mg/L	4
	EC50	48	Crustacea		0.013	3mg/L	5
nickel	EC50	72	Algae or other aquati	c plants	0.040	07mg/L	2
	BCF			Algae or other aquatic plants			
	BCF	1440	Algae or other aquati	c plants	0.471	mg/L	4
	EC50	1440 720	Algae or other aquati Crustacea	c plants		mg/L 62mg/L	4 2
				•	0.006	-	
	EC50 NOEC	720 72	Crustacea Algae or other aquati	•	0.000	62mg/L 35mg/L	2 2
	EC50 NOEC ENDPOINT	720 72 TEST DURATION (HR)	Crustacea Algae or other aquati	•	0.000 0.000	52mg/L 35mg/L LUE	2 2 SOURCE
potassium nitrate	EC50 NOEC	720 72	Crustacea Algae or other aquati	•	0.000 0.000	62mg/L 35mg/L	2 2
potassium nitrate	EC50 NOEC ENDPOINT	720 72 TEST DURATION (HR)	Crustacea Algae or other aquati	c plants	0.000 0.000 VAI 22.0	52mg/L 35mg/L LUE	2 2 SOURCE
potassium nitrate	EC50 NOEC ENDPOINT LC50	720 72 TEST DURATION (HR) 96	Crustacea Algae or other aquati SPECIES Fish	c plants	0.000 0.000 VAI 22.0 118	52mg/L 35mg/L LUE 5mg/L	2 2 SOURCE 4
potassium nitrate	EC50 NOEC ENDPOINT LC50 EC50 EC50	720 72 TEST DURATION (HR) 96 96 384	Crustacea Algae or other aquati SPECIES Fish Algae or other aquat Crustacea	c plants	0.000 0.000 VAI 22. 118 49.	62mg/L 35mg/L LUE 5mg/L 11.887mg/L 116mg/L	2 2 SOURCE 4 3 3
potassium nitrate	EC50 NOEC ENDPOINT LC50 EC50 EC50 EC50	720 72 TEST DURATION (HR) 96 96 384 TEST DURATION (HR)	Crustacea Algae or other aquati SPECIES Fish Algae or other aquat Crustacea SPECIES SPECIES	c plants	0.000 0.000 22. 118 49.	62mg/L 35mg/L LUE 5mg/L 11.887mg/L 116mg/L	2 2 SOURCE 4 3 3 3 SOURCE
potassium nitrate	EC50 NOEC ENDPOINT LC50 EC50 EC50 EC50 EC50	720 72 TEST DURATION (HR) 96 96 384 TEST DURATION (HR) 96	Crustacea Algae or other aquati SPECIES Fish Algae or other aquat Crustacea	c plants	0.000 0.000 22. 118 49. VA	62mg/L 35mg/L LUE 5mg/L 11.887mg/L 11.6mg/L ALUE 00mg/L	2 2 SOURCE 4 3 3 3 SOURCE 4
	EC50 NOEC ENDPOINT LC50 EC50 EC50 EC50	720 72 TEST DURATION (HR) 96 96 384 TEST DURATION (HR)	Crustacea Algae or other aquati SPECIES Fish Algae or other aquat Crustacea SPECIES SPECIES	c plants	0.000 0.000 22. 118 49. VA	62mg/L 35mg/L LUE 5mg/L 11.887mg/L 116mg/L	2 2 SOURCE 4 3 3 3 SOURCE
potassium nitrate	EC50 NOEC ENDPOINT LC50 EC50 EC50 EC50 EC50	720 72 TEST DURATION (HR) 96 96 384 TEST DURATION (HR) 96	Crustacea Algae or other aquati SPECIES Fish Algae or other aquat Crustacea SPECIES Fish Fish	c plants	0.000 0.000 22. 118 49. VAI 22. 118 49.	62mg/L 35mg/L LUE 5mg/L 11.887mg/L 11.6mg/L ALUE 00mg/L	2 2 SOURCE 4 3 3 3 SOURCE 4
	EC50 NOEC ENDPOINT LC50 EC50 EC50 EC50 EC50 EC50 EC50	720 72 TEST DURATION (HR) 96 96 384 TEST DURATION (HR) 96 48	Crustacea Algae or other aquati SPECIES Fish Algae or other aquat Crustacea SPECIES Fish Crustacea	c plants	0.000 0.000 22. 118 49. VAI 24.	62mg/L 62mg/L 135mg/L 10E 5mg/L 116mg/L 116mg/L 176mg/L	2 2 SOURCE 4 3 3 3 SOURCE 4 1
	EC50 NOEC ENDPOINT LC50 EC50 EC50 EC50 EC50 EC50 EC50	720 72 TEST DURATION (HR) 96 96 384 TEST DURATION (HR) 96 48 96 48 96	Crustacea Algae or other aquati SPECIES Fish Algae or other aquati Crustacea SPECIES Fish Crustacea Algae or other aquat	c plants	0.000 0.003 22. 118 49. 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7	62mg/L 62mg/L 35mg/L 5mg/L 11.887mg/L 116mg/L ALUE 20mg/L 176mg/L 12mg/L	2 2 3 4 3 3 3 5 5 0 URCE 4 1 4 4
	EC50 NOEC ENDPOINT LC50 EC50 EC50 EC50 EC50 EC50 EC50 EC50	720 72 TEST DURATION (HR) 96 96 384 TEST DURATION (HR) 96 48 96 384	Crustacea Algae or other aquati SPECIES Fish Algae or other aquati Crustacea SPECIES Fish Crustacea Algae or other aquat Crustacea Algae or other aquat Crustacea	c plants	0.000 0.003 22. 118 49. 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7	62mg/L 62mg/L 35mg/L LUE 5mg/L 11.887mg/L 116mg/L 116mg/L 00mg/L 176mg/L 12mg/L 19.200mg/L	2 2 3 3 3 3 5 5 5 0 7 7 4 1 4 3 3
	EC50 NOEC ENDPOINT LC50 EC50 EC50 EC50 EC50 EC50 EC50 EC50	720 72 TEST DURATION (HR) 96 96 384 TEST DURATION (HR) 96 48 96 384	Crustacea Algae or other aquati SPECIES Fish Algae or other aquati Crustacea SPECIES Fish Crustacea Algae or other aquat Crustacea Algae or other aquat Crustacea	c plants	0.000 0.000 22. 118 49. 24 30 = 1 24 14 24	62mg/L 62mg/L 35mg/L LUE 5mg/L 11.887mg/L 116mg/L 116mg/L 00mg/L 176mg/L 12mg/L 19.200mg/L	2 2 3 3 3 3 5 5 5 0 7 7 4 1 4 3 3

72 2 96 POINT TE 96 96 96 96 96 97 96 97 97 72 72 72 72 72 72 72 72 72 7	EST DURATION (HR)		aquatic plants		>43.3mg/L >=40.3mg/L VALUE 0.693mg/L 2.387mg/L 0.9894mg/L 1.162mg/L	2 2 SO 2 2 2 2 2 2	OURCE
POINT TE 96 48 72 72	EST DURATION (HR)	SPECIES Fish Crustacea Algae or other	• •		VALUE 0.693mg/L 2.387mg/L 0.9894mg/L	2 2 2 2	URCE
96 48 72 72	2 2	Fish Crustacea Algae or other	• •		0.693mg/L 2.387mg/L 0.9894mg/L	2 2 2 2	URCE
96 48 72 72	2 2	Fish Crustacea Algae or other	• •		0.693mg/L 2.387mg/L 0.9894mg/L	2 2 2 2	URCE
48 72 72	3 2 2	Crustacea Algae or other	• •		2.387mg/L 0.9894mg/L	2	
72	2	Algae or other	• •		0.9894mg/L	2	
72	2		• •				
		Algae or other	aquatic plants		1.162mg/L	2	
C 72	,		Algae or other aquatic plants				
		Algae or other aquatic plants		0.0168mg/L	2		
POINT	TEST DURATION (HR)		SPECIES	VA	LUE	SOURCI	E
C	16		Crustacea	107	rmg/L	4	
POINT	TEST DURATION (HR)	SPECI	S	VALUE		SOURCE	
pplicable	Not Applicable		-	-		Not Applical	able
	POINT	16 POINT TEST DURATION (HR) pplicable Not Applicable	Test Duration (HR) SPECIA pplicable Not Applicable Not Applicable	C 16 Crustacea POINT TEST DURATION (HR) SPECIES pplicable Not Applicable Not Applicable	C 16 Crustacea 107 POINT TEST DURATION (HR) SPECIES VALUE pplicable Not Applicable Not Applicable Not Applicable	Count TEST DURATION (HR) SPECIES VALUE pplicable Not Applicable Not Applicable Not Applicable	C 16 Crustacea 107mg/L 4 POINT TEST DURATION (HR) SPECIES VALUE SOURCE

(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 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.

for Boron and Borates:

Environmental Fate - Boron is generally found in nature bound to oxygen and is never found as the free element. As an element, boron itself cannot be degraded in the environment, however, it may undergo various reactions that change the form of boron (e.g., precipitation, polymerization, and acid-base reactions) depending on conditions such as its concentration in water and pH. As boron is a natural component of the environment, individuals will have some exposure from foods and drinking water.

Atmospheric Fate: Atmospheric boron may be in the form of particulate matter or aerosols as borides, boron oxides, borates, organoboron compounds, trihalide boron compounds, or borazines. Boron and borates will probably be removed from the atmosphere by precipitation and dry deposition. The half-life of airborne particles is usually on the order of days, depending on the size of the particle and atmospheric conditions.

Aquatic Fate: Borates are relatively soluble in water. Boron readily hydrolyses in water and, in concentrated solutions, may polymerize. The mineral content of water is not likely to control the fate of boron in water. Boron was found to not be significantly removed during the conventional treatment of waste water. Boron may, however; be co-precipitated with aluminium, silicon, or iron to form hydroxyborate compounds on the surfaces of minerals. Waterborne boron may be adsorbed by soils and sediments. Adsorption-desorption reactions are expected to be the only significant mechanism that will influence the fate of boron in water.

Terrestrial Fate: Soil - Boron is added to farmland as a soil improving agent, but there is not sufficient data to evaluate its effect on soil organisms. The extent of boron adsorption depends on the pH of the water and the chemical composition of the soil. The greatest adsorption is generally observed at pH 7.5-9.0. The single most important property of soil that will influence the mobility of boron is the abundance of amorphous aluminium oxide. The extent of boron adsorption has also been attributed to the levels of iron oxide, and to a lesser extent, the organic matter present in the soil, although other studies found that the amount of organic matter present was not important. The adsorption of boron may not be reversible in some soils. Most boron compounds are transformed to borates in soil due to the presence of moisture. Borates themselves are not further degraded in soil, however; borates can exist in a variety of forms in soil. Borates are removed from soils by water leaching and by assimilation by plants. Surface soil, unpolluted waterways and seawater all typically contain significant amounts of boron as borate. Plants - Boron is an essential micronutrient for healthy growth of plants, however, it can be harmful to boron sensitive plants in higher quantities. In some areas such as the American Southwest, boron occurs naturally in surface waters in concentrations that have been shown to be toxic to commercially important plants.

Ecotoxicity: It is unlikely that boron is bioconcentrated significantly by organisms from water. Boron is not expected to bioaccumulate and bioconcentration factors for fish, plants and invertebrates are low. Boron is not regarded to be dangerous to aquatic organisms. In aquatic environments low concentrations of borates generally promote the growth of algae, whereas higher concentrations inhibited algal growth. Boron has little effect on freshwater algae and water fleas. The toxicity of boron in fish is often higher in soft water than in hard water. Zebra fish and rainbow trout are the most sensitive species to the effects of boron.

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
boric acid	LOW	LOW

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Catalogue number: ICP-WS-2 Version No: 3.3

potassium nitrate	LOW	LOW
sodium carbonate	LOW	LOW
ammonium metavanadate	HIGH	HIGH
water	LOW	LOW

Bioaccumulative potential

Ingredient	Bioaccumulation			
boric acid	LOW (BCF = 0)			
potassium nitrate	LOW (LogKOW = 0.209)			
sodium carbonate	LOW (LogKOW = -0.4605)			
ammonium metavanadate	LOW (LogKOW = 2.229)			
water	LOW (LogKOW = -1.38)			

Mobility in soil

Ingredient	Mobility
boric acid	LOW (KOC = 35.04)
potassium nitrate	LOW (KOC = 14.3)
sodium carbonate	HIGH (KOC = 1)
ammonium metavanadate	LOW (KOC = 35.04)
water	LOW (KOC = 14.3)

SECTION 13 DISPOSAL CONSIDERATIONS

Waste treatment methods

	 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.
Product / Packaging	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
disposal	specifically licensed to accept chemical and / or pharmaceutical wastes or Incineration in a licensed apparatus (after admixture with suitable combustible material).
	 Decontaminate empty containers with 5% aqueous sodium hydroxide or soda ash, followed by water. Observe all label safeguards until containers are cleaned and destroyed.

SECTION 14 TRANSPORT INFORMATION

Labels Required



Land transport (DOT)

UN number	64							
UN proper shipping name	orrosive liquid, acidic, inorganic, n.o.s. (contains nitric acid)							
Transport hazard class(es)	Class8SubriskNot Applicable							
Packing group								
Environmental hazard	Not Applicable							
Special precautions for user	Hazard Label8Special provisions386, B2, IB2, T11, TP2, TP27							

Air transport (ICAO-IATA / DGR)

	-						
UN number	3264	164					
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					

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	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

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 ACGIH Threshold Limit Values (TLV) - Carcinogens
US ATSDR Minimal Risk Levels for Hazardous Substances (MRLs)
US EPA Carcinogens Listing
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 EPA Carcinogens Listing
US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory
US - Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminants
US - Vermont Permissible Exposure Limits Table Z-1-A Transitional Limits for Air Contaminants
US - Washington Permissible exposure limits of air contaminants
US - 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 - Wyoming Toxic and Hazardous Substances Table Z-2 Acceptable ceiling concentration,
Acceptable maximum peak above the acceptable ceiling concentration for an 8-hr shift
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) - Toxic Pollutants
US EPA Carcinogens Listing
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 Carcinogens Listing
US OSHA Permissible Exposure Levels (PELs) - Table Z1
US OSHA Permissible Exposure Levels (PELs) - Table Z2
US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory

US - Rhode Island Hazardous Substance List

US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants

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atalogue number: ICP-WS-2	ICP-WS-2	Print Date: 06/02/201
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	IETE	
CALCIUM CARBONATE(471-34-1) IS FOUND ON THE FOLLOWING REGULATORY L		
US - Alaska Limits for Air Contaminants	US - Rhode Island Hazardous Substance List	Limite For Air Conteminants
US - California Permissible Exposure Limits for Chemical Contaminants	US - Tennessee Occupational Exposure Limits -	
US - Hawaii Air Contaminant Limits	US - Vermont Permissible Exposure Limits Table	
US - Idaho - Limits for Air Contaminants	US - Vermont Permissible Exposure Limits Table Contaminants	2-1-A Transitional Limits for Air
US - Massachusetts - Right To Know Listed Chemicals US - Michigan Exposure Limits for Air Contaminants	US - Washington Permissible exposure limits of a	air contaminants
US - Minnesota Permissible Exposure Limits (PELs)	US - Wyoming Toxic and Hazardous Substances	
US - Oregon Permissible Exposure Limits (Z-1)	US NIOSH Recommended Exposure Limits (REL	
US - Pennsylvania - Hazardous Substance List	US OSHA Permissible Exposure Levels (PELs) -	,
US - Perinsylvania - Hazarduus Substance List	US Toxic Substances Control Act (TSCA) - Cherr	
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 Contaminants	Z-1-A Transitional Limits for Air
US - Alaska Limits for Air Contaminants	US - Washington Permissible exposure limits of a	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) - Carci	nogens
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) - Priority Pollutants	
US - Oregon Permissible Exposure Limits (Z-1)	US CWA (Clean Water Act) - Toxic Pollutants	
US - Pennsylvania - Hazardous Substance List	US EPCRA Section 313 Chemical List	
US - Rhode Island Hazardous Substance List	US NIOSH Recommended Exposure Limits (REL	_s)
US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants	US OSHA Permissible Exposure Levels (PELs) -	- Table Z1
COBALT(7440-48-4) 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 Transitional Limits for Air
Monographs	Contaminants	
US - Alaska Limits for Air Contaminants	US - Washington Permissible exposure limits of a	
US - California Permissible Exposure Limits for Chemical Contaminants	US - Washington Toxic air pollutants and their AS	
US - California Proposition 65 - Carcinogens	US - Wyoming Toxic and Hazardous Substances	Table 21 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) - Carci	-
US - Massachusetts - Right To Know Listed Chemicals	US ATSDR Minimal Risk Levels for Hazardous S	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	and Dad D
US - New Jersey Right to Know - Special Health Hazard Substance List (SHHSL): Carcinogens	US National Toxicology Program (NTP) 14th Rep	
US - Oregon Permissible Exposure Limits (Z-1)	US NIOSH Recommended Exposure Limits (REL	
	US OSHA Permissible Exposure Levels (PELs) -	
US - Pennsylvania - Hazardous Substance List US - Rhode Island Hazardous Substance List	US Priority List for the Development of Propositio Levels (NSRLs) for Carcinogens and Maximum A	
US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants	Chemicals Causing Reproductive Toxicity	Rilowable Dose Levels (MADLS) 101
US - Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminants		nical 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)		
US - California Permissible Exposure Limits for Chemical Contaminants	US - Washington Permissible exposure limits of a	air contaminants
US - Hawaii Air Contaminant Limits	US - Washington Toxic air pollutants and their AS	IL, 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 S	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 Tennessee Occupational Exposure Limits Limits For Air Contaminants
- US Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminants

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

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

- 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 Massachusetts Right To Know Listed Chemicals
- US Michigan Exposure Limits for Air Contaminants
- US Oregon Permissible Exposure Limits (Z-1)

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

- 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 Pennsylvania Hazardous Substance List
- US Rhode Island Hazardous Substance List
- US Tennessee Occupational Exposure Limits 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 Toxic Substances Control Act (TSCA) Chemical Substance Inventory

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atalogue number: ICP-WS-2	ICP-WS-2	Print Date: 06/02/2
rsion No: 3.3		
US - Alaska Limits for Air Contaminants	US - Vermont Permissit	ole Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminants
US - California OEHHA/ARB - Chronic Reference Exposure Levels and Target Organs		ble Exposure Limits Table Z-1-A Transitional Limits for Air
(CRELs)	Contaminants	
US - California Permissible Exposure Limits for Chemical Contaminants		issible exposure limits of air contaminants
US - Hawaii Air Contaminant Limits	-	air pollutants and their ASIL, SQER and de minimis emission values
US - Idaho - Limits for Air Contaminants	-	d Hazardous Substances Table Z1 Limits for Air Contaminants
US - Michigan Exposure Limits for Air Contaminants	US Clean Air Act - Haz	
US - Minnesota Permissible Exposure Limits (PELs)	US EPCRA Section 313	3 Chemical List
US - Oregon Permissible Exposure Limits (Z-1)		Exposure Levels (PELs) - Table Z1
US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants		Control Act (TSCA) - Chemical Substance Inventory
NICKEL(7440-02-0) IS FOUND ON THE FOLLOWING REGULATORY LISTS US - Alaska Limits for Air Contaminants		ole Exposure Limits Table Z-1-A Transitional Limits for Air
	Contaminants	Die Exposure Limits Table 2-1-A Transitional Limits for All
US - California OEHHA/ARB - Acute Reference Exposure Levels and Target Organs (RELs)		issible exposure limits of air contaminants
US - California OEHHA/ARB - Chronic Reference Exposure Levels and Target Organs (CRELs)	-	d Hazardous Substances Table Z1 Limits for Air Contaminants
US - California Permissible Exposure Limits for Chemical Contaminants	US ACGIH Threshold L	
US - California Proposition 65 - Carcinogens		.imit Values (TLV) - Carcinogens
US - Hawaii Air Contaminant Limits		
US - Idaho - Limits for Air Contaminants		sk Levels for Hazardous Substances (MRLs)
	US Clean Air Act - Haz	Act) - Priority Pollutants
US - Massachusetts - Right To Know Listed Chemicals	· ·	, .
US - Michigan Exposure Limits for Air Contaminants	US CWA (Clean Water	
US - Minnesota Permissible Exposure Limits (PELs)	US EPCRA Section 31:	
US - New Jersey Right to Know - Special Health Hazard Substance List (SHHSL):		Program (NTP) 14th Report Part B.
Carcinogens		ded Exposure Limits (RELs)
US - Oregon Permissible Exposure Limits (Z-1)		Exposure Levels (PELs) - Table Z1
US - Pennsylvania - Hazardous Substance List US - Rhode Island Hazardous Substance List		Development of Proposition 65 Safe Harbor Levels - No Significant Ris
	Chemicals Causing Re	cinogens and Maximum Allowable Dose Levels (MADLs) for
US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants US - Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminants		Control Act (TSCA) - Chemical Substance Inventory
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SECTION 311/312 HAZARD CATEGORIES

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

Reactivity hazard

No

US. EPA CERCLA HAZARDOUS SUBSTANCES AND REPORTABLE QUANTITIES (40 CFR 302.4)		
Name	Reportable Quantity in Pounds (Ib)	Reportable Quantity in kg
Cadmium	10	4.54
Chromium	5000	2270
Copper	5000	2270
Nickel	100	45.4
Ammonium vanadate	1000	454
Nitric acid	1000	454

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	Y
Canada - DSL	Υ
Canada - NDSL	N (bismuth; strontium nitrate; ammonium metavanadate; barium nitrate; magnesium; copper; water; cobalt; nickel; manganese(II) acetate tetrahydrate; boric acid; chromium; potassium nitrate; sodium carbonate; cadmium; nitric acid)
China - IECSC	Υ
Europe - EINEC / ELINCS / NLP	Y
Japan - ENCS	N (bismuth; magnesium; copper; water; cobalt; nickel; manganese(II) acetate tetrahydrate; boric acid; chromium; sodium carbonate; cadmium; nitric acid)
Korea - KECI	Υ
New Zealand - NZIoC	Υ
Philippines - PICCS	Υ
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
barium nitrate	10022-31-8, 34053-87-7
boric acid	10043-35-3, 11113-50-1, 41685-84-1
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
sodium carbonate	497-19-8, 7542-12-3, 1314087-39-2, 1332-57-6
strontium nitrate	10042-76-9, 13470-05-8

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

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

Definitions and abbreviations

PC-TWA: Permissible Concentration-Time Weighted Average

PC-STEL: Permissible Concentration-Short Term Exposure Limit

IARC: International Agency for Research on Cancer

ACGIH: American Conference of Governmental Industrial Hygienists

STEL: Short Term Exposure Limit

TEEL: Temporary Emergency Exposure Limit。 IDLH: Immediately Dangerous to Life or Health Concentrations

OSF: Odour Safety Factor

NOAEL : No Observed Adverse Effect Level

LOAEL: Lowest Observed Adverse Effect Level

TLV: Threshold Limit Value

LOD: Limit Of Detection

OTV: Odour Threshold Value

BCF: BioConcentration Factors

BEI: Biological Exposure Index

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