

High-Purity Standards

Catalogue number: ICP-MS-TS-4

Version No: 1.1

Safety Data Sheet according to OSHA HazCom Standard (2012) requirements

Chemwatch Hazard Alert Code: 3

Issue Date: **05/19/2017**Print Date: **05/19/2017**S GHS USA EN

SECTION 1 IDENTIFICATION

Product Identifier

Product name	ICP-MS Tuning Solution 4
Synonyms	ICP-MS-TS-4
Proper shipping name	Corrosive liquid, acidic, inorganic, n.o.s.
Other means of identification	ICP-MS-TS-4

Recommended use of the chemical and restrictions on use

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 / Organisation	INFOTRAC
Emergency telephone numbers	1-800-535-5053
Other emergency 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

Label elements

Hazard pictogram(s)



SIGNAL WORD D

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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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
7429-90-5	0.001	aluminium
7440-39-3	0.001	<u>barium</u>
543-81-7	0.001	<u>beryllium acetate</u>
7440-69-9	0.001	<u>bismuth</u>
7440-45-1	0.001	cerium
7440-48-4	0.001	cobalt
7440-74-6	0.001	<u>indium</u>
7439-92-1	0.001	<u>lead</u>
7439-95-4	0.001	<u>magnesium</u>
7440-02-0	0.001	nickel
7803-55-6	0.001	ammonium metavanadate
7440-65-5	0.001	<u>yttrium</u>
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:

- $\blacksquare \ \ \, \text{Immediately hold eyelids apart and flush the eye continuously with running water.}$
- ▶ 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.

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.

For thermal burns:

- ► Decontaminate area around burn.
- ▶ Consider the use of cold packs and topical antibiotics.

For first-degree burns (affecting top layer of skin)

- Hold burned skin under cool (not cold) running water or immerse in cool water until pain subsides.
- ▶ Use compresses if running water is not available.
- Cover with sterile non-adhesive bandage or clean cloth.
- ▶ Do NOT apply butter or ointments; this may cause infection.
- ▶ Give over-the counter pain relievers if pain increases or swelling, redness, fever occur.

Skin Contact

- For second-degree burns (affecting top two layers of skin)
- ► Cool the burn by immerse in cold running water for 10-15 minutes
- Use compresses if running water is not available.
 Do NOT apply ice as this may lower body temperature and cause further damage.
- ▶ Do NOT break blisters or apply butter or ointments; this may cause infection.
- ▶ Protect burn by cover loosely with sterile, nonstick bandage and secure in place with gauze or tape.
- To prevent shock: (unless the person has a head, neck, or leg injury, or it would cause discomfort):
- ▶ Lay the person flat.
- ► Elevate feet about 12 inches.
- ▶ Elevate burn area above heart level, if possible.
- ► Cover the person with coat or blanket.
- Seek medical assistance.

For third-degree burns

Seek immediate medical or emergency assistance.

In the mean time:

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Protect burn area cover loosely with sterile, nonstick bandage or, for large areas, a sheet or other material that will not leave lint in wound. Separate burned toes and fingers with dry, sterile dressings. Do not soak burn in water or apply ointments or butter; this may cause infection. To prevent shock see above. For an airway burn, do not place pillow under the person's head when the person is lying down. This can close the airway. Have a person with a facial burn sit up ▶ Check pulse and breathing to monitor for shock until emergency help arrives. • If 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 Inhalation of vapours or aerosols (mists, fumes) may cause lung oedema. ▶ Corrosive substances may cause lung damage (e.g. lung oedema, fluid in the lungs). As this reaction may be delayed up to 24 hours after exposure, affected individuals need complete rest (preferably in semi-recumbent posture) and must be kept under medical observation even if no symptoms are (yet) manifested. • Before any such manifestation, the administration of a spray containing a dexamethasone derivative or beclomethasone derivative may be considered. This must definitely be left to a doctor or person authorised by him/her. (ICSC13719) ▶ For advice, contact a Poisons Information Centre or a doctor at once. Urgent hospital treatment is likely to be needed. wed 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. Ingestion Observe the patient carefully Never give liquid to a person showing signs of being sleepy or with reduced awareness; i.e. becoming unconscious. Give water to rinse out mouth, then provide liquid slowly and as much as casualty can comfortably drink. Transport to hospital or doctor without delay.

Most important symptoms and effects, both acute and delayed

See Section 11

Indication of any immediate medical attention and special treatment needed

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 furnes. May emit acrid smoke. When aluminium oxide dust is dispersed in air, firefighters should wear protection against inhalation of dust particles, which can also contain hazardous substances from the fire absorbed on the alumina particles.

SECTION 6 ACCIDENTAL RELEASE MEASURES

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See section 8

Environmental precautions

See section 12

Methods and material for containment and cleaning up

▶ 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

Minor Spills

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

SECTION 7 HANDLING AND STORAGE

Precautions for safe handling

- Avoid all personal contact, including inhalation.
- Wear protective clothing when risk of exposure occurs.
- Use in a well-ventilated area.
- WARNING: To avoid violent reaction, ALWAYS add material to water and NEVER water to material
- Avoid smoking, naked lights or ignition sources.
- Avoid contact with incompatible materials.
- When handling, DO NOT eat, drink or smoke
- Keep containers securely sealed when not in use.
- Avoid physical damage to containers.
- Always wash hands with soap and water after handling.
- Work clothes should be laundered separately. Launder contaminated clothing before re-use.
- Use good occupational work practice.
- Observe manufacturer's storage and handling recommendations contained within this SDS.
- Atmosphere should be regularly checked against established exposure standards to ensure safe working conditions are maintained.

Other information

Safe handling

- Store in original containers Keep containers securely sealed.
- Store in a cool, dry, well-ventilated area.
- Store away from incompatible materials and foodstuff containers.
- Protect containers against physical damage and check regularly for leaks. Observe manufacturer's storage and handling recommendations contained within this SDS

Conditions for safe storage, including any incompatibilities

► DO NOT use aluminium or galvanised containers

- ▶ Check regularly for spills and leaks
- ▶ Lined metal can, lined metal pail/ can.
- Plastic pail.
- Polvliner drum.
- ► Packing as recommended by manufacturer.
- Check all containers are clearly labelled and free from leaks.

For low viscosity materials

- Suitable container
- Drums and ierricans 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.

For aluminas (aluminium oxide):

Incompatible with hot chlorinated rubber.

In the presence of chlorine trifluoride may react violently and ignite.

-May initiate explosive polymerisation of olefin oxides including ethylene oxide.

-Produces exothermic reaction above 200 C with halocarbons and an exothermic reaction at ambient temperatures with halocarbons in the presence of other metals.

-Produces exothermic reaction with oxygen difluoride.

-May form explosive mixture with oxygen difluoride

-Forms explosive mixtures with sodium nitrate.

-Reacts vigorously with vinyl acetate. Storage incompatibility

Aluminium oxide is an amphoteric substance, meaning it can react with both acids and bases, such as hydrofluoric acid and sodium hydroxide, acting as an acid with a base and a base with an acid, neutralising the other and producing a salt

- ▶ 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.

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- ▶ Inorganic acids can initiate the polymerisation of certain classes of organic compounds.
- ► Inorganic acids react with cyanide compounds to release gaseous hydrogen cyanide.
- Inorganic acids generate flammable and/or toxic gases in contact with dithiocarbamates, isocyanates, mercaptans, nitrides, nitrides, sulfides, and strong reducing agents. Additional gas-generating reactions occur with sulfites, nitrites, thiosulfates (to give H2S and SO3), dithionites (SO2), and even carbonates.
- ► Acids often catalyse (increase the rate of) chemical reactions.
- WARNING: Avoid or control reaction with peroxides. All transition metal peroxides should be considered as potentially explosive. For example transition metal complexes of alkyl hydroperoxides may decompose explosively.
- ► The pi-complexes formed between chromium(0), 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	aluminium	Aluminum, metal	15 mg/m3	Not Available	Not Available	Total dust; (as Al)
US OSHA Permissible Exposure Levels (PELs) - Table Z1	aluminium	Aluminum, metal- Respirable fraction	5 mg/m3	Not Available	Not Available	(as Al)
US NIOSH Recommended Exposure Limits (RELs)	aluminium	Aluminium, Aluminum metal, Aluminum powder, Elemental aluminum	10 (total), 5 (resp) mg/m3	Not Available	Not Available	Not Available
US 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)	indium	Indium metal	0.1 mg/m3	Not Available	Not Available	[*Note: The REL also applies to other indium compounds (as In).]
US NIOSH Recommended Exposure Limits (RELs)	lead	Lead metal, Plumbum	0.050 mg/m3	Not Available	Not Available	See Appendix C [*Note: The REL also applies to other lead compounds (as Pb) see Appendix C.]
US NIOSH Recommended Exposure Limits (RELs)	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	yttrium	Yttrium	1 mg/m3	Not Available	Not Available	[*Note: The REL also applies to other yttrium compounds (as Y).]
US NIOSH Recommended Exposure Limits (RELs)	yttrium	Yttrium metal	1 mg/m3	Not Available	Not Available	Not Available
US OSHA Permissible Exposure Levels (PELs) - Table Z1	nitric acid	Nitric acid	5 mg/m3 / 2 ppm	10 mg/m3 / 4 ppm	Not Available	TLV® Basis: URT & eye irr; dental erosion
US NIOSH Recommended Exposure Limits (RELs)	nitric acid	Aqua fortis, Engravers acid, Hydrogen nitrate, Red fuming nitric acid (RFNA), White fuming nitric acid (WFNA)	5 mg/m3 / 2 ppm	4 ppm	Not Available	Not Available
US ACGIH Threshold Limit Values (TLV)	nitric acid	Nitric acid	2 ppm	Not Available	Not Available	Not Available

| EMERGENCY LIMITS

Ingredient	Material name	TEEL-1	TEEL-2	TEEL-3
barium	Barium	1.5 mg/m3	180 mg/m3	1,100 mg/m3
bismuth	Bismuth	15 mg/m3	170 mg/m3	990 mg/m3
cerium	Cerium	30 mg/m3	330 mg/m3	2,000 mg/m3
cobalt	Cobalt	0.18 mg/m3	2 mg/m3	20 mg/m3
indium	Indium	0.3 mg/m3	3.3 mg/m3	20 mg/m3
lead	Lead	0.15 mg/m3	120 mg/m3	700 mg/m3
magnesium	Magnesium	18 mg/m3	200 mg/m3	1,200 mg/m3
nickel	Nickel	4.5 mg/m3	50 mg/m3	99 mg/m3
ammonium metavanadate	Ammonium vanadate; (Ammonium vanadium oxide; Ammonium metavanadate)	0.01 mg/m3	0.11 mg/m3	80 mg/m3
yttrium	Yttrium	3 mg/m3	33 mg/m3	200 mg/m3
nitric acid	Nitric acid	Not Available	Not Available	Not Available

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Ingredient	Original IDLH	Revised IDLH
aluminium	Not Available	Not Available
barium	1,100 mg/m3	50 mg/m3
beryllium acetate	10 mg/m3	4 mg/m3
bismuth	Not Available	Not Available
cerium	Not Available	Not Available
cobalt	20 mg/m3	20 [Unch] mg/m3
indium	Not Available	Not Available
lead	700 mg/m3	100 mg/m3
magnesium	Not Available	Not Available
nickel	N.E. / N.E.	10 mg/m3
ammonium metavanadate	Not Available	Not Available
yttrium	Not Available	Not Available
nitric acid	100 ppm	25 ppm
water	Not Available	Not Available

Exposure controls

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

The basic types of engineering controls are:

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

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

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

Local exhaust ventilation usually required. If risk of overexposure exists, wear approved respirator. Correct fit is essential to obtain adequate protection. Supplied-air type respirator may be required in special circumstances. Correct fit is essential to ensure adequate protection.

An approved self contained breathing apparatus (SCBA) may be required in some situations.

Provide adequate ventilation in warehouse or closed storage area. Air contaminants generated in the workplace possess varying "escape" velocities which, in turn, determine the "capture velocities" of fresh circulating air required to effectively remove the contaminant.

Appropriate engineering controls

Type of Contaminant:	Air Speed:
solvent, vapours, degreasing etc., evaporating from tank (in still air).	0.25-0.5 m/s (50-100 f/min.)
aerosols, fumes from pouring operations, intermittent container filling, low speed conveyer transfers, welding, spray drift, plating acid fumes, pickling (released at low velocity into zone of active generation)	0.5-1 m/s (100-200 f/min.)
direct spray, spray painting in shallow booths, drum filling, conveyer loading, crusher dusts, gas discharge (active generation into zone of rapid air motion)	1-2.5 m/s (200-500 f/min.)
grinding, abrasive blasting, tumbling, high speed wheel generated dusts (released at high initial velocity into zone of very high rapid air motion).	2.5-10 m/s (500-2000 f/min.)

Within each range the appropriate value depends on:

Lower end of the range	Upper end of the range
1: Room air currents minimal or favourable to capture 1: Disturbing room air currents	
2: Contaminants of low toxicity or of nuisance value only.	2: Contaminants of high toxicity
3: Intermittent, low production.	3: High production, heavy use
4: Large hood or large air mass in motion	4: Small hood-local control only

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

Personal protection











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- ► 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.
- 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.

Eye and 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]

Skin protection

See Hand protection below

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

SECTION 10 STABILITY AND REACTIVITY

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

Information on toxicologic	al 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. Exposure to vapours of some rare earth salts can cause sensitivity to heat, itching, and increased sensitivity of smell and taste. Other effects include inflamed airways and lung, emphysema, regional narrowing of terminal airways and cell changes. 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.
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. Though considered non-harmful, slight irritation may result from contact because of the abrasive nature of the aluminium oxide particles. Thus it may cause itching and skin reaction and inflammation. Open cuts, abraded or irritated skin should not be exposed to this material

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	Entry into the blood-stream, through, for example, cuts, abrasions or of the material and ensure that any external damage is suitably prote If applied to the eyes, this material causes severe eye damage.		roduce systemic injury with	harmful effects.	Examine the skin prior to the use
Eye	Direct eye contact with acid corrosives may produce pain, tears, se completely.	ensitivity to light a	and burns. Mild burns of the	e epithelia gene	rally recover rapidly and
Chronic	Repeated or prolonged exposure to acids may result in the erosion and inflammation of lung tissue often occurs. Long-term exposure to respiratory irritants may result in airways dis Substance accumulation, in the human body, may occur and may ca Animal testing shows long term exposure to aluminium oxides may the greater the tendencies of causing harm. Yttrium is a rare earth metal - heavy type (yttrium family). There hav abnormalities due to its high density.	sease, involving ause some conc cause lung dise	difficulty breathing and rela ern following repeated or lo ease and cancer, depending	ted whole-body ng-term occupa g on the size of	problems. tional exposure. the particle. The smaller the size,
ICP-MS Tuning Solution 4	TOXICITY Not Available		ot Available		
	TVOCAVAIRABLE	14	Ot Available		
	TOXICITY			IRRITATION	
aluminium	Oral (rat) LD50: >2000 mg/kg ^[1]			Not Available	
				I	
	TOXICITY	IR	RRITATION		
barium	Not Available	N	ot Available		
beryllium acetate	TOXICITY	IR	RRITATION		
•	Not Available	N	ot Available		
	TOWN TO THE TOWN T				
bismuth	TOXICITY			Not Available	
	Oral (rat) LD50: 2000 mg/kg ^[1]			Not Available	
	TOXICITY			IRRITATION	
cerium	Oral (rat) LD50: >5000 mg/kg ^[1]			Not Available	
	Cia (ta) 2500. 20000 mg/lg				
	TOXICITY			IRRITATIO	N
cobalt	dermal (rat) LD50: >2000 mg/kg ^[1]			Not Availab	le
	Oral (rat) LD50: 6170 mg/kgd ^[2]				
indium	TOXICITY	IR	RRITATION		
indium	Not Available	N	ot Available		
	-				
	TOXICITY			IRRITAT	
lead	dermal (rat) LD50: >2000 mg/kg ^[1]			Not Ava	ilable
	Inhalation (rat) LC50: >5.05 mg/l/4hr ^[1]				
	Oral (rat) LD50: >2000 mg/kg ^[1]				
	TOXICITY			IRRITATION	
magnesium	Oral (rat) LD50: >2000 mg/kg ^[1]			Not Available	
	Grai (rai) ED50. >2000 Highly				
	TOXICITY			IRRITATION	
nickel	Oral (rat) LD50: 5000 mg/kg ^[2]			Not Available	
	, , , ,		I		
	тохісіту			IRRITATION	
ammonium metavanadate	dermal (rat) LD50: 2102 mg/kg ^[2]			Not Available	
	Oral (rat) LD50: 160 mg/kgd ^[2]				
yttrium	TOXICITY	IR	RRITATION		
yandiii	Not Available	N	ot Available		
	<u> </u>				

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Legend:

1. Value obtained from Europe ECHA Registered Substances - Acute toxicity 2.* Value obtained from manufacturer's SDS. Unless otherwise specified data extracted from RTECS - Register of Toxic Effect of chemical Substances

BERYLLIUM ACETATE	WARNING: This substance has been classified by the IARC as Group 1: CARCINOGENIC TO	HUMANS.		
COBALT	Allergic reactions involving the respiratory tract are usually due to interactions between IgE antibodies and allergens and occur rapidly. Attention should be paid to atopic diathesis, characterised by increased susceptibility to nasal inflammation, asthma and eczema. Exogenous allergic alveolitis is induced essentially by allergen specific immune-complexes of the IgG type; cell-mediated reactions (T lymphocytes) may be involved.			
LEAD	WARNING: Lead is a cumulative poison and has the potential to cause abortion and intellectual in	npairment to unborn children of pregnant workers.		
NICKEL	Tenth Annual Report on Carcinogens: Substance anticipated to be Carcinogen [National Toxicology Program: U.S. Dep. Oral (rat) TDLo: 500 mg/kg/5D-I Inhalation (rat) TCLo: 0.1 mg/m3/24H/17W-C			
YTTRIUM	For typical lanthanides: Symptoms of toxicity from rare earth elements include writhing, inco-ordination, laboured breathing	g, and sedation.		
NITRIC ACID	For acid mists, aerosols, vapours Test results suggest that eukaryotic cells are susceptible to genetic damage when the pH falls to about 6.5. The material may produce severe irritation to the eye causing pronounced inflammation. The material may produce respiratory tract irritation, and result in damage to the lung including reduced lung function. 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)			
ALUMINIUM & BARIUM & CERIUM & INDIUM & YTTRIUM & WATER	No significant acute toxicological data identified in literature search.			
BARIUM & BERYLLIUM ACETATE & AMMONIUM METAVANADATE & NITRIC ACID	Asthma-like symptoms may continue for months or even years after exposure to the material ends.			
BERYLLIUM ACETATE & COBALT & NICKEL	The following information refers to contact allergens as a group and may not be specific to this product.			
CERIUM & YTTRIUM	Lanthanide poisoning causes immediate defaecation, writhing, inco-ordination, laboured breathing	g, and inactivity.		
COBALT & NICKEL	WARNING: This substance has been classified by the IARC as Group 2B: Possibly Carcinogeni	ic to Humans.		
Acute Toxicity	Carcinogenicity	0		
Skin Irritation/Corrosion	Reproductivity	0		
Serious Eye Damage/Irritation	STOT - Single Exposure	0		
Respiratory or Skin sensitisation	STOT - Repeated Exposure	0		
Mutagenicity		0		

Legend:

X − Data available but does not fill the criteria for classification
 ✓ − Data available to make classification

O - Data Not Available to make classification

SECTION 12 ECOLOGICAL INFORMATION

Toxicity

ICP-MS Tuning Solution 4	ENDPOINT	TEST DURATION (HR)		SPECIES	VALUE	Ē	SOURCE
	Not Applicable	Not Applicable	Not Applicable Not		Not Ap	pplicable Not Applicab	
				-		1	
	ENDPOINT	TEST DURATION (HR)	SPECIES			VALUE	SOURCE
	LC50	96	Fish			0.078-0.108mg/L	2
	EC50	48	Crustac	ea		0.7364mg/L	2
aluminium	EC50	96	Algae or other aquatic plants		0.0054mg/L	2	
	BCF	360	Algae or other aquatic plants		9mg/L	4	
	EC50	120	Fish	Fish		0.000051mg/L	5
	NOEC	72	Algae or other aquatic plants		>=0.004mg/L	2	
	ENDPOINT	TEST DURATION (HR)	SPEC	IES		VALUE	SOURCE
	LC50	96	Fish	Fish >500mg/L		>500mg/L	4
barium	EC50	96	Algae	Algae or other aquatic plants 26mg/L		26mg/L	4
barium	BCF	24	Crusta	icea		0.000002mg/L	4
	EC50	240	Algae	or other aquatic plants		8.10306mg/L	4
	NOEC	48	Crusta	icea		68mg/L	4

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96 48 72 1440 720 72 TEST DU 96 48 72 72 72 72 TESS Not	URATION (HR) URATION (HR) ST DURATION (HR) Applicable EST DURATION (HR)	SPECIE Fish Crustac Algae o Crustac Algae o SPEC Fish Crust Algae Algae Algae	cea or other aquatic plants or other aquatic plants cea or other aquatic plants	ts ts ts Not Appli	>25.5mg VALUE 0.0000475mg 0.013mg/L 0.0407mg/L 0.0062mg/L 0.0035mg/L VALUE 0.693mg/L 0.9894mg 1.162mg/L 0.0168mg cable	yL L L g/L Not A	2 SOURCE 4 5 2 4 2 2 2 2 2 2 2 2
TEST DU 96 48 72 1440 720 72 TEST DU 96 48 72 72 72 72 TEST DU 72	URATION (HR)	SPECIE Fish Crustac Algae o Crustac Algae o SPEC Fish Crust Algae Algae Algae	cea or other aquatic plants or other aquatic plants or other aquatic plants or other aquatic plants cea or other aquatic plants decea e or other aquatic plan	ts ts ts ts VALUE	VALUE 0.0000475mg 0.013mg/L 0.0407mg/L 0.0062mg/L 0.0035mg/L VALUE 0.693mg/L 2.387mg/ 0.9894mg/L 1.162mg/L 0.0168mg/L	yL L L g/L L g/L	SOURCE 4 5 2 4 2 2 2 2 2 2 2 2
TEST DU 96 48 72 1440 720 72 TEST DU 96 48 72 72 72 72 TEST DU 72	URATION (HR)	SPECIE Fish Crustac Algae o Crustac Algae o SPEC Fish Crust Algae Algae Algae	cea or other aquatic plants or other aquatic plants or other aquatic plants or other aquatic plants cea or other aquatic plants decea e or other aquatic plan	ts ts ts ts VALUE	VALUE 0.0000475mg 0.013mg/L 0.0407mg/L 0.0062mg/L 0.0035mg/L VALUE 0.693mg/L 2.387mg/ 0.9894mg/L 1.162mg/L 0.0168mg/L	yL L L g/L L g/L	SOURCE 4 5 2 4 2 2 2 2 2 2 2 2
TEST DU 96 48 72 1440 720 72 TEST DU 96 48 72 72 72 72	URATION (HR)	SPECIE Fish Crustac Algae o Crustac Algae o SPEC Fish Crust Algae Algae Algae	cea or other aquatic plants or other aquatic plants or other aquatic plants or other aquatic plants cea or other aquatic plants tacea e or other aquatic plan	ts ts ts	VALUE 0.0000475mg 0.013mg/L 0.0407mg/L 0.47mg/L 0.0062mg/L 0.0035mg/L VALUE 0.693mg/ 2.387mg/ 0.9894mg/ 1.162mg/L	yL L L g/L L	SOURCE 4 5 2 4 2 2 SOURCE 2 2 2 2 2
TEST DU 96 48 72 1440 720 72 TEST DU 96 48 72 72		SPECIE Fish Crustac Algae o Crustac Algae o SPEC Fish Crust Algae Algae Algae	cea or other aquatic plants or other aquatic plants or other aquatic plants or other aquatic plants cea or other aquatic plants dacea e or other aquatic plan e or other aquatic plan e or other aquatic plan	ts ts	VALUE 0.0000475mg 0.013mg/L 0.0407mg/L 0.47mg/L 0.0062mg/L 0.0035mg/L VALUE 0.693mg/ 2.387mg/ 0.9894mg/ 1.162mg/L	\(\) \(\)	SOURCE 4 5 2 4 2 2 2 2 2 2 2 2
TEST DU 96 48 72 1440 720 72 TEST DU 96 48 72 72		SPECIE Fish Crustac Algae o Crustac Algae o SPEC Fish Crust Algae Algae Algae	cea or other aquatic plants or other aquatic plants or other aquatic plants or other aquatic plants cea or other aquatic plants dacea e or other aquatic plan e or other aquatic plan e or other aquatic plan	ts ts	VALUE 0.0000475mg 0.013mg/L 0.0407mg/L 0.47mg/L 0.0062mg/L 0.0035mg/L VALUE 0.693mg/ 2.387mg/ 0.9894mg/ 1.162mg/L	\(\) \(\)	SOURCE 4 5 2 4 2 2 2 2 2 2 2 2
72 1440 720 72 TEST DU 96 48 72 72 72		SPECIE Fish Crustac Algae o Algae o Crustac Algae o SPEC Fish Crust	cea or other aquatic plants or other aquatic plants cea or other aquatic plants cea or other aquatic plants cles tacea e or other aquatic plan	nts	VALUE 0.0000475mg/L 0.0407mg/L 0.47mg/L 0.0062mg/L 0.0035mg/L VALUE 0.693mg/L 0.9894mg/L	yL L L	SOURCE 4 5 2 4 2 2 SOURCE 2 2 2 2
72 1440 720 72 TEST DU 96 48		SPECIE Fish Crustac Algae o Algae o Crustac Algae o SPEC Fish Crust	cea or other aquatic plants or other aquatic plants cea or other aquatic plants cea or other aquatic plants	nts	VALUE 0.0000475mg 0.013mg/L 0.0407mg/L 0.47mg/L 0.0062mg/L 0.0035mg/L VALUE 0.693mg/ 2.387mg/	yL L	SOURCE 4 5 2 4 2 2 2 SOURCE 2 2
72 1440 720 72 TEST DU 96		SPECIE Fish Crustace Algae o Crustace Algae o SPECIE Fish	cea or other aquatic plants or other aquatic plants cea or other aquatic plants	nts	VALUE 0.0000475mg 0.013mg/L 0.0407mg/L 0.47mg/L 0.0062mg/L 0.0035mg/L VALUE 0.693mg/	yL L	SOURCE 4 5 2 4 2 2 2 SOURCE 2
72 1440 720 72 TEST DU		SPECIE Fish Crustace Algae o Crustace Algae o SPECIE	cea or other aquatic plants or other aquatic plants cea or other aquatic plants	nts	VALUE 0.0000475mg 0.013mg/L 0.0407mg/L 0.47mg/L 0.0062mg/L 0.0035mg/L VALUE	y'L	SOURCE 4 5 2 4 2 2 2 SOURCE
72 1440 720	JRATION (HR)	SPECIE Fish Crustace Algae o Algae o Crustace	cea or other aquatic plants or other aquatic plants or other aquatic plants	nts	VALUE 0.0000475mg 0.013mg/L 0.0407mg/L 0.47mg/L 0.0062mg/L		\$0URCE 4 5 2 4 4 2 2 2
72 1440 720	JRATION (HR)	SPECIE Fish Crustace Algae o Algae o Crustace	cea or other aquatic plants or other aquatic plants or other aquatic plants	nts	VALUE 0.0000475mg 0.013mg/L 0.0407mg/L 0.47mg/L 0.0062mg/L		SOURCE 4 5 2 4 2
72 1440 720	JRATION (HR)	SPECIE Fish Crustace Algae o Algae o Crustace	cea or other aquatic plants or other aquatic plants or other aquatic plants	nts	VALUE 0.0000475mg 0.013mg/L 0.0407mg/L 0.47mg/L 0.0062mg/L		SOURCE 4 5 2 4 2
7EST DU 96 48 72 1440	JRATION (HR)	SPECIE Fish Crustac Algae o	cea or other aquatic plants or other aquatic plants	nts	VALUE 0.0000475mg 0.013mg/L 0.0407mg/L 0.47mg/L		SOURCE 4 5 2 4
72 TEST DU	JRATION (HR)	SPECIE Fish Crustac Algae o	ea or other aquatic plants	nts	VALUE 0.0000475mg 0.013mg/L 0.0407mg/L		SOURCE 4 5 2
TEST DU 96 48	JRATION (HR)	SPECIE Fish Crustac	ES	nts	VALUE 0.0000475mg 0.013mg/L		SOURCE 4 5
TEST DU	IRATION (HR)	SPECIE Fish	ES		VALUE 0.0000475mg		SOURCE 4
TEST DU	JRATION (HR)	SPECIE			VALUE		SOURCE
						g/L	
72		Alga	e or other aquatic plar		>25.5m	g/L	2
70		A1.	o or other		. 05.5	œ/I	
EC50 72		Alga	e or other aquatic plar	າເຮ	>20mg/	L	2
72		Algae or other aquatic plants		>20mg/	-		
96		Fish			541mg/l		2
	UKATION (HR)						SOURCE
672		Fish			0.00003mg	g/L	4
48			S	0.0217mg/L		2	
8		Fish			-		4
72		Algae or other aquatic plants		-		2	
			Crustacea		_		2
96		Fish			-		2
	JRATION (HR)		IES				SOURCE
TECT :	IDATION (UD)	6777	150		V411:-		00115.07
INOT	луріісаріє		ινοι ΑμμισαύΙΕ	Not Appli	CADIC	INOL A	ppiicable
					cable		pplicable
TES	ST DURATION (HR)		SPECIES	VALUE		SOU	RCE
168		Algae	e or other aquatic plan	ts	0.0018m	g/L	2
70		Algae	e or other aquatic plan	ts	0.02mg/L		2
1344		Fish			0.99mg/L		4
72		Algae	e or other aquatic plan	ts	0.144mg/	'L	2
48		Crust	tacea		>0.89mg	/L	2
96		Fish			1.406mg/	L L	2
TEST DI	URATION (HR)	SPEC	CIES		VALUE		SOURCE
			1	'			
			Not Applicable		cable		pplicable
TES	ST DURATION (HR)		SPECIES	VALUE		SOUI	RCE
				1			
					cable		pplicable
TES	ST DUPATION (HP)		SDECIES	VALUE		SOLII	PCE
NOC	Арріісавіе		Not Applicable	1401 Арріі	cable	NOLA	pplicable
					coblo		
	TEST DI 96 48 72 1344 70 168 TEST DI 96 48 72 8 48 672	48 72 1344 70 168 TEST DURATION (HR) Not Applicable TEST DURATION (HR) 96 48 72 8 48	TEST DURATION (HR) Not Applicable	Not Applicable Not Applicable	Not Applicable Not Applicable	Not Applicable Not Applicable Not Applicable	Not Applicable Not

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wate

ENDPOINT	TEST DURATION (HR)	SPECIES	VALUE	SOURCE
Not Applicable	Not Applicable	Not Applicable	Not Applicable	Not Applicable

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:

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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 Lanthanoids (Formerly Lanthanides: Synonym Rare Earth Metals and their Salts):

Environmental Fate: Rare earths, such as the lanthanoids, are relatively abundant in the crust of the Earth. These elements are not • rare • -scientists once thought these substances were only found in very small amounts on the Earth Most of the lanthanides occur together in nature, and they are very difficult to separate from each other. The lanthanides form alloys, (mixtures), with many other metals, and these alloys exhibit a wide range of physical properties. Lanthanoid emissions to the environment have increased as a result of the growing industrial applications of these elements; however, robust data to evaluate the environmental fate of lanthanoids are scarce.

Atmospheric Fate: These substances react with oxygen in the atmosphere to form an oxide residue which tamishes surfaces exposed to these elements. They burn readily in air to form oxides. Terrestrial Fate: Soil - Lanthanoids can be found in most soils. These substances are expected to strongly sorb to soil and are not expected to evaporate from soil surfaces. Plants • These substances are expected to accumulate in plants, especially duckweed.

Aquatic Fate: Rare earth chlorides are very poorly soluble in water. These substances will bind to carbonated and dissolved organic matter in water. The lanthanides react slowly with cold water and more rapidly with hot water to form hydrogen gas. The lanthanum ion is expected to have high attraction to the negatively charged humic material present in most natural waters. This mechanism will also remove lanthanum from the water column.

Ecotoxicity: These elements have a high tendency to accumulate in plants and organisms. A typical oxide of this group, cerium oxide, has low toxicity to the fathead minnow, green algae, and Daphnia water fleas. Rare earth chlorides exhibit acute aquatic toxicity at concentrations exceeding 100 ppm and chronic toxicity, persisting for more than 21 days, at concentrations greater than 30 ppm. Industrial processes have little impact on altering background levels. Lanthanum 3+ is toxic to some aquatic organisms. Dissolved lanthanum is very toxic to species of Daphnia in both chronic and acute tests and may also be toxic to other species. There seems little doubt that dissolved lanthanum has at least high acute and chronic toxicity to fresh water fish and to various species of Daphnia in soft water, although water quality appears to have a very large effect on the toxicity.

Prevent, by any means available, spillage from entering drains or water courses.

DO NOT discharge into sewer or waterways

Persistence and degradability

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

Bioaccumulative potential

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

Mobility in soil

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

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

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- 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

NO

Land transport (DOT)

UN number	3264
UN proper shipping name	Corrosive liquid, acidic, inorganic, n.o.s.
Transport hazard class(es)	Class 8 Subrisk Not Applicable
Packing group	
Environmental hazard	Not Applicable
Special precautions for user	Hazard Label 8 Special provisions 386, B2, IB2, T11, TP2, TP27

Air transport (ICAO-IATA / DGR)

• • • • • • • • • • • • • • • • • • • •	•	
UN number	3264	
UN proper shipping name	CORROSIVE LIQUID, ACIDIC, INORGANIC, N.O.S.	
Transport hazard class(es)	ICAO/IATA Class 8 ICAO / IATA Subrisk Not Applicable ERG Code 8L	
Packing group	П	
Environmental hazard	Not Applicable	
Special precautions for user	Special provisions Cargo Only Packing Instructions Cargo Only Maximum Qty / Pack Passenger and Cargo Packing Instructions Passenger and Cargo Maximum Qty / Pack Passenger and Cargo Limited Quantity Packing Instructions	A3A803 855 30 L 851 1 L Y840
	Passenger and Cargo Limited Maximum Qty / Pack	0.5 L

Sea transport (IMDG-Code / GGVSee)

UN number	3264
UN proper shipping name	Corrosive liquid, acidic, inorganic, n.o.s. *
Transport hazard class(es)	IMDG Class 8 IMDG Subrisk Not Applicable
Packing group	Ш
Environmental hazard	Not Applicable
Special precautions for user	EMS Number F-A, S-B Special provisions 274 Limited Quantities 1 L

Transport in bulk according to Annex II of MARPOL and the IBC code

Source	Product name	Pollution Category	Ship Type

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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 - Alaska Limits for Air Contaminants

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 - Minnesota Permissible Exposure Limits (PELs)

US - Oregon Permissible Exposure Limits (Z-1)

US - Pennsylvania - Hazardous Substance List

US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants

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

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

US - Alaska Limits for Air Contaminants

US - Hawaii Air Contaminant Limits

US - Idaho - Limits for Air Contaminants

US - Massachusetts - Right To Know Listed Chemicals

US - Minnesota Permissible Exposure Limits (PELs)

US - Pennsylvania - Hazardous Substance List

US - Rhode Island Hazardous Substance List

US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants

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

BERYLLIUM ACETATE(543-81-7) IS FOUND ON THE FOLLOWING REGULATORY LISTS

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

Monographs

US - Alaska Limits for Air Contaminants

US - California OEHHA/ARB - Chronic Reference Exposure Levels and Target Organs (CRELs)

US - California Permissible Exposure Limits for Chemical Contaminants

US - Hawaii Air Contaminant Limits

US - Idaho - Acceptable Maximum Peak Concentrations

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 - Oregon Permissible Exposure Limits (Z-2)

US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants

 ${\it US-Vermont\ Permissible\ Exposure\ Limits\ Table\ Z-1-A\ Final\ Rule\ Limits\ for\ Air\ Contaminants}$

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

BISMUTH(7440-69-9) IS FOUND ON THE FOLLOWING REGULATORY LISTS

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

CERIUM(7440-45-1) IS FOUND ON THE FOLLOWING REGULATORY LISTS

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

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

US - Washington Permissible exposure limits of air contaminants

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

US ACGIH Threshold Limit Values (TLV)

US ACGIH Threshold Limit Values (TLV) - Carcinogens

US ATSDR Minimal Risk Levels for Hazardous Substances (MRLs)

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

US - Washington Permissible exposure limits of air contaminants

US ACGIH Threshold Limit Values (TLV)

US ACGIH Threshold Limit Values (TLV) - Carcinogens

Contaminants

US EPA Carcinogens Listing
US EPCRA Section 313 Chemical List

US OSHA Permissible Exposure Levels (PELs) - Table Z1

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

US - 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 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 OSHA Permissible Exposure Levels (PELs) - Table Z1 $\,$

US OSHA Permissible Exposure Levels (PELs) - Table Z2 $\,$

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 - Alaska Limits for Air Contaminants

US - California Permissible Exposure Limits for Chemical Contaminants

US - California Proposition 65 - Carcinogens

US - Hawaii Air Contaminant Limits

US - Idaho - Limits for Air Contaminants

US - Massachusetts - Right To Know Listed Chemicals

US - Michigan Exposure Limits for Air Contaminants

US - Minnesota Permissible Exposure Limits (PELs)

US - New Jersey Right to Know - Special Health Hazard Substance List (SHHSL): Carcinogens

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

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 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 EPCRA Section 313 Chemical List

US National Toxicology Program (NTP) 14th Report Part B.

US NIOSH Recommended Exposure Limits (RELs)

US OSHA Permissible Exposure Levels (PELs) - Table Z1

US Priority List for the Development of Proposition 65 Safe Harbor Levels - No Significant Risk Levels (NSRLs) for Carcinogens and Maximum Allowable Dose Levels (MADLs) for Chemicals Causing Reproductive Toxicity

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

INDIUM(7440-74-6) IS FOUND ON THE FOLLOWING REGULATORY LISTS

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

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International Agency for Research on Cancer (IARC) - Agents Classified by the IARC US - Rhode Island Hazardous Substance List Monographs US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants US - Alaska Limits for Air Contaminants $\label{lem:us-vermont} \textbf{US-Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminants}$ US - California OEHHA/ARB - Chronic Reference Exposure Levels and Target Organs US - Vermont Permissible Exposure Limits Table Z-1-A Transitional Limits for Air (CRELs) US - California Permissible Exposure Limits for Chemical Contaminants US - Washington Permissible exposure limits of air contaminants US - Hawaii Air Contaminant Limits US - Wyoming Toxic and Hazardous Substances Table Z1 Limits for Air Contaminants US - Massachusetts - Right To Know Listed Chemicals US ACGIH Threshold Limit Values (TLV) US - Michigan Exposure Limits for Air Contaminants US NIOSH Recommended Exposure Limits (RELs)

US - Minnesota Permissible Exposure Limits (PELs)

US - Oregon Permissible Exposure Limits (Z-1) US - Pennsylvania - Hazardous Substance List

LEAD(7439-92-1) IS FOUND ON THE FOLLOWING REGULATORY LISTS		
International Agency for Research on Cancer (IARC) - Agents Classified by the IARC	US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants	
Monographs	US - Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminants	
US - Alaska Limits for Air Contaminants	US - Vermont Permissible Exposure Limits Table Z-1-A Transitional Limits for Air	
US - California - Proposition 65 - Priority List for the Development of MADLs for Chemicals	Contaminants	
Causing Reproductive Toxicity	US - Washington Permissible exposure limits of air contaminants	
US - California Permissible Exposure Limits for Chemical Contaminants	US - Washington Toxic air pollutants and their ASIL, SQER and de minimis emission values	
US - California Proposition 65 - Carcinogens	US ACGIH Threshold Limit Values (TLV)	
US - California Proposition 65 - Maximum Allowable Dose Levels (MADLs) for Chemicals	US ACGIH Threshold Limit Values (TLV) - Carcinogens	
Causing Reproductive Toxicity	US Clean Air Act - Hazardous Air Pollutants	
US - California Proposition 65 - No Significant Risk Levels (NSRLs) for Carcinogens	US CWA (Clean Water Act) - Priority Pollutants	
US - California Proposition 65 - Reproductive Toxicity	US CWA (Clean Water Act) - Toxic Pollutants	
US - Hawaii Air Contaminant Limits	US EPA Carcinogens Listing	
US - Idaho - Acceptable Maximum Peak Concentrations	US EPCRA Section 313 Chemical List	
US - Idaho - Limits for Air Contaminants	US National Toxicology Program (NTP) 14th Report Part B.	

US - Massachusetts - Right To Know Listed Chemicals US NIOSH Recommended Exposure Limits (RELs) US - Minnesota Permissible Exposure Limits (PELs) US OSHA Permissible Exposure Levels (PELs) - Table Z1 US - New Jersey Right to Know - Special Health Hazard Substance List (SHHSL): US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory Carcinogens US - Pennsylvania - Hazardous Substance List US - Rhode Island Hazardous Substance List

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

International Agency for Research on Cancer (IARC) - Agents Classified by the IARC	US - Pennsylvania - Hazardous Substance List	
Monographs	US - Rhode Island Hazardous Substance List	
US - California OEHHA/ARB - Chronic Reference Exposure Levels and Target Organs	US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants	
(CRELs)	US - 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 Toxic Substances Control Act (TSCA) - Chemical Substance Inventory	
US - Massachusetts - Right To Know Listed Chemicals	,	
US - Michigan Exposure Limits for Air Contaminants		

US - Oregon Permissible Exposure Limits (Z-1)

NICKEL(7440-02-0) IS FOUND ON THE FOLLOWING REGULATORY LISTS	
US - Alaska Limits for Air Contaminants	US - Vermont Permissible Exposure Limits Table Z-1-A Transitional Limits for Air
US - California OEHHA/ARB - Acute Reference Exposure Levels and Target Organs (RELs)	Contaminants
US - California OEHHA/ARB - Chronic Reference Exposure Levels and Target Organs	US - Washington Permissible exposure limits of air contaminants
(CRELs)	US - Wyoming Toxic and Hazardous Substances Table Z1 Limits for Air Contaminants
US - California Permissible Exposure Limits for Chemical Contaminants	US ACGIH Threshold Limit Values (TLV)
US - California Proposition 65 - Carcinogens	US ACGIH Threshold Limit Values (TLV) - Carcinogens
US - Hawaii Air Contaminant Limits	US ATSDR Minimal Risk Levels for Hazardous Substances (MRLs)
US - Idaho - Limits for Air Contaminants	US Clean Air Act - Hazardous Air Pollutants
US - Massachusetts - Right To Know Listed Chemicals	US CWA (Clean Water Act) - Priority Pollutants
US - Michigan Exposure Limits for Air Contaminants	US CWA (Clean Water Act) - Toxic Pollutants
US - Minnesota Permissible Exposure Limits (PELs)	US EPCRA Section 313 Chemical List
US - New Jersey Right to Know - Special Health Hazard Substance List (SHHSL):	US National Toxicology Program (NTP) 14th Report Part B.
Carcinogens	US NIOSH Recommended Exposure Limits (RELs)
US - Oregon Permissible Exposure Limits (Z-1)	US OSHA Permissible Exposure Levels (PELs) - Table Z1
US - Pennsylvania - Hazardous Substance List	US Priority List for the Development of Proposition 65 Safe Harbor Levels - No Significant Risk
US - Rhode Island Hazardous Substance List	Levels (NSRLs) for Carcinogens and Maximum Allowable Dose Levels (MADLs) for
US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants	Chemicals Causing Reproductive Toxicity
US - Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminants	US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory

AMMONIUM METAVANADATE(7803-55-6) IS FOUND ON THE FOLLOWING REGULATORY LISTS

US - California OEHHA/ARB - Acute Reference Exposure Levels and Target Organs (RELs)	US EPCRA Section 313 Chemical List
US - Massachusetts - Right To Know Listed Chemicals	US Toxic Substances Control Act (TSCA) - Chemical Substance Inventory
US - Pennsylvania - Hazardous Substance List	

YTTRIUM(7440-65-5) IS FOUND ON THE FOLLOWING REGULATORY LISTS

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International Air Transport Association (IATA) Dangerous Goods Regulations - Prohibited List	US - Tennessee Occupational Exposure Limits - Limits For Air Contaminants	
Passenger and Cargo Aircraft	US - Vermont Permissible Exposure Limits Table Z-1-A Final Rule Limits for Air Contaminants	
US - Alaska Limits for Air Contaminants	US - Vermont Permissible Exposure Limits Table Z-1-A Transitional Limits for Air	
US - Hawaii Air Contaminant Limits	Contaminants	
US - Idaho - Limits for Air Contaminants	US - Washington Permissible exposure limits of air contaminants	
US - Massachusetts - Right To Know Listed Chemicals	US - Wyoming Toxic and Hazardous Substances Table Z1 Limits for Air Contaminants	
US - Michigan Exposure Limits for Air Contaminants	US ACGIH Threshold Limit Values (TLV)	
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 Toxic Substances Control Act (TSCA) - Chemical Substance Inventory	
US - Rhode Island Hazardous Substance List		

NITRIC ACID(7697-37-2) IS FOUND ON THE FOLLOWING REGULATORY LISTS

International Air Transport Association (IATA) Dangerous Goods Regulations - Prohibited List	US - Vermont Permissible Exposure Limits Table Z-1-A 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

Federal Regulations

Superfund Amendments and Reauthorization Act of 1986 (SARA)

SECTION 311/312 HAZARD CATEGORIES

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

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

Name	Reportable Quantity in Pounds (lb)	Reportable Quantity in kg
Lead	10	4.54
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

Cobalt metal powder, Lead and lead compounds: Lead, Nickel (Metallic) Listed

National Inventory	Status
Australia - AICS	N (yttrium; beryllium acetate)
Canada - DSL	N (beryllium acetate)
Canada - NDSL	N (bismuth; lead; cerium; indium; ammonium metavanadate; magnesium; water; barium; yttrium; aluminium; cobalt; nickel; beryllium acetate; nitric acid)
China - IECSC	N (cerium; beryllium acetate)
Europe - EINEC / ELINCS / NLP	Υ
Japan - ENCS	N (bismuth; cerium; indium; magnesium; water; barium; yttrium; aluminium; cobalt; nickel; beryllium acetate; nitric acid)
Korea - KECI	N (beryllium acetate)
New Zealand - NZIoC	N (beryllium acetate)
Philippines - PICCS	N (yttrium; beryllium acetate)
USA - TSCA	N (beryllium acetate)
Legend:	Y = All ingredients are on the inventory N = Not determined or one or more ingredients are not on the inventory and are not exempt from listing(see specific ingredients in brackets)

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SECTION 16 OTHER INFORMATION

Other information

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Ingredients with multiple cas numbers

Name	CAS No
aluminium	7429-90-5, 91728-14-2

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

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

Definitions and abbreviations

PC-TWA: Permissible Concentration-Time Weighted Average

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

IARC: International Agency for Research on Cancer

ACGIH: American Conference of Governmental Industrial Hygienists

STEL: Short Term Exposure Limit

TEEL: Temporary Emergency Exposure Limit $_{\circ}$

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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