# Vortex 71T

#### Weldwell New Zealand

Chemwatch: 4622-89 Version No: 3.1.1.1 Safety Data Sheet according to WHS and ADG requirements emwatch Hazard Alert Code: 3

Issue Date: 27/06/2017 Print Date: 20/06/2019 L.GHS.AUS.EN

# SECTION 1 IDENTIFICATION OF THE SUBSTANCE / MIXTURE AND OF THE COMPANY / UNDERTAKING

#### **Product Identifier**

Product name	Vortex 71T
Synonyms	flux cored electrode; 71 T; rutile wire
Other means of identification	Not Available

#### Relevant identified uses of the substance or mixture and uses advised against

Relevant identified uses	Flux cored electrode.
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# Details of the supplier of the safety data sheet

Registered company name	Weldwell New Zealand
Address	64 Thames Street, Napier, New Zealand
Telephone	+64 6 834-1600
Fax	+64 6 835-4568
Website	www.weldwell.co.nz
Email	Not Available

# Emergency telephone number

Association / Organisation	Weldwell New Zealand
Emergency telephone numbers	+64 6 834-1600
Other emergency telephone numbers	0800 CHEMCALL (0800 243 622) 24 Hours

# **SECTION 2 HAZARDS IDENTIFICATION**

### Classification of the substance or mixture

# HAZARDOUS CHEMICAL. NON-DANGEROUS GOODS. According to the WHS Regulations and the ADG Code.

#### CHEMWATCH HAZARD RATINGS

	Min	Max	
Flammability	0		
Toxicity	0		0 = Minimum
Body Contact	1 📕	1	1 = Low
Reactivity	1	1	2 = Moderate 3 = High
Chronic	3		4 = Extreme

Poisons Schedule	Not Applicable
Classification <sup>[1]</sup>	Carcinogenicity Category 1A
Legend:	1. Classified by Chemwatch; 2. Classification drawn from HCIS; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI

### Hazard statement(s)

 H350
 May cause cancer.

 Precautionary statement(s)
 Prevention

 P201
 Obtain special instructions before use.

 P281
 Use personal protective equipment as required.

# Precautionary statement(s) Response

P308+P313	IF exposed or concerned: Get medical advice/attention.	

# 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
Not Available		carbon steel tube, containing
7439-89-6	10	iron
13463-67-7	<6	titanium dioxide
7439-96-5	<6	manganese
92704-41-1	<6	kaolin, calcined
7440-21-3	1	silicon
Not Available	0.7	silica crystalline - quartz
Not Available	<0.7	kaolin
Not Available	<0.7	calcium fluoride
Not Available		In use product generates
Not Available	>60	welding fumes
Not Available		iron oxide fume
7439-96-5.		manganese fume
16984-48-8		fluoride fume
69012-64-2		silica welding fumes
Not Available		action of arc on air may generate
10028-15-6		ozone
Mixture		nitrogen oxides

# SECTION 4 FIRST AID MEASURES

# Description of first aid measures

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If this product comes in contact with the eyes: Wash • out immediately with fresh running water.

Eye Contact

Finite complete irrigation of the eye by keeping eyelids apart and away from eye and moving the eyelids by

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	<ul> <li>occasionally lifting the upper and lower lids.</li> <li>Seek medical attention without delay; if pain persists or recurs seek medical attention.</li> <li>Removal of contact lenses after an eye injury should only be undertaken by skilled personnel.</li> </ul>
Skin Contact	If skin or hair contact occurs: ▶ Flush skin and hair with running water (and soap if available). ▶ Seek medical attention in event of irritation.
Inhalation	<ul> <li>If fumes or combustion products are inhaled remove from contaminated area.</li> <li>Lay patient down. Keep warm and rested.</li> <li>Prostheses such as false teeth, which may block airway, should be removed, where possible, prior to initiating first aid procedures.</li> <li>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.</li> <li>Transport to hospital, or doctor.</li> </ul>
Ingestion	<ul> <li>Immediately give a glass of water.</li> <li>First aid is not generally required. If in doubt, contact a Poisons Information Centre or a doctor.</li> </ul>

# Indication of any immediate medical attention and special treatment needed

Treat symptomatically.

### **SECTION 5 FIREFIGHTING 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	Avoid contamination with oxidising agents i.e. nitrates, oxidising acids, chlorine bleaches, pool chlorine etc. as ignition may result
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#### Advice for firefighters

Fire Fighting	Alert Fire Brigade and tell them location and nature of hazard. Wear breathing apparatus plus protective gloves in the event of a fire. Prevent, by any means available, spillage from entering drains or water courses. Use fire fighting procedures suitable for surrounding area. <b>DO NOT</b> approach containers suspected to be hot. Cool fire exposed containers with water spray from a protected location. If safe to do so, remove containers from path of fire. Equipment should be thoroughly decontaminated after use.
Fire/Explosion Hazard	Non combustible. Not considered a significant fire risk, however containers may burn. Decomposition may produce toxic fumes of: metal oxides May emit poisonous fumes.
HAZCHEM	Not Applicable

# 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	Clean up all spills immediately. Secure load if safe to do so. Bundle/collect recoverable product. Collect remaining material in containers with covers for disposal.
Major Spills	Clean up all spills immediately. Wear protective clothing, safety glasses, dust mask, gloves. Secure load if safe to do so. Bundle/collect recoverable product. Use dry clean up procedures and avoid generating dust. Vacuum up (consider explosion-proof machines designed to be grounded during storage and use). Water may be used to prevent dusting.

<ul> <li>Collect remaining material in containers with covers for disposal.</li> <li>Flush spill area with water.</li> </ul>	
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Personal Protective Equipment advice is contained in Section 8 of the SDS.

# SECTION 7 HANDLING AND STORAGE

# Precautions for safe handling

Safe handling	<ul> <li>Avoid all personal contact, including inhalation.</li> <li>Wear protective clothing when risk of exposure occurs.</li> <li>Use in a well-ventilated area.</li> <li>Prevent concentration in hollows and sumps.</li> <li>DO NOT enter confined spaces until atmosphere has been checked.</li> <li>DO NOT allow material to contact humans, exposed food or food utensils.</li> <li>Avoid contact with incompatible materials.</li> <li>When handling, DO NOT eat, drink or smoke.</li> <li>Keep containers securely sealed when not in use.</li> <li>Avoid physical damage to containers.</li> <li>Always wash hands with soap and water after handling.</li> <li>Work clothes should be laundered separately. Launder contaminated clothing before re-use.</li> <li>Use good occupational work practice.</li> <li>Observe manufacturer's storage and handling recommendations contained within this SDS.</li> <li>Atmosphere should be regularly checked against established exposure standards to ensure safe working conditions are maintained.</li> </ul>
Other information	Store in original containers. Keep containers securely sealed. Store in a cool, dry, well-ventilated area. Store away from incompatible materials and foodstuff containers. Protect containers against physical damage and check regularly for leaks. Observe manufacturer's storage and handling recommendations contained within this SDS.

# Conditions for safe storage, including any incompatibilities

Suitable container	Polyethylene or polypropylene container. Packing as recommended by manufacturer. Check all containers are clearly labelled and free from leaks.
Storage incompatibility	Avoid reaction with oxidising agents

# SECTION 8 EXPOSURE CONTROLS / PERSONAL PROTECTION

# **Control parameters**

# OCCUPATIONAL EXPOSURE LIMITS (OEL)

# INGREDIENT DATA

Source	Ingredient	Material name	TWA	STEL	Peak	Notes
Australia Exposure Standards	titanium dioxide	Titanium dioxide	10 mg/m3	Not Available	Not Available	<ul> <li>(a) This value is for inhalable dust containing no asbestos and &lt; 1% crystalline silica.</li> </ul>
Australia Exposure Standards	manganese	Manganese, fume (as Mn)	1 mg/m3	3 mg/m3	Not Available	Not Available
Australia Exposure Standards	silicon	Silicon	10 mg/m3	Not Available	Not Available	<ul> <li>(a) This value is for inhalable dust containing no asbestos and &lt; 1% crystalline silica.</li> </ul>
Australia Exposure Standards	manganese fume	Manganese, fume (as Mn)	1 mg/m3	3 mg/m3	Not Available	Not Available
Australia Exposure Standards	ozone	Ozone	Not Available	Not Available	0.1 ppm / 0.2 mg/m3	Not Available

### EMERGENCY LIMITS

Ingredient	Material name	TEEL-1	TEEL-2	TEEL-3
iron	Iron	3.2 mg/m3	35 mg/m3	150 mg/m3
titanium dioxide	Titanium oxide; (Titanium dioxide)	30 mg/m3	330 mg/m3	2,000 mg/m3

manganese	Manganese	3 mg/	m3	5 mg/m3	1,800 mg/m3	
silicon	Silicon	45 mg	g/m3	100 mg/m3	630 mg/m3	
manganese fume	Manganese	3 mg/	m3	5 mg/m3	1,800 mg/m3	
fluoride fume	Fluorides (as F)	7.5 m	g/m3	83 mg/m3	500 mg/m3	
silica welding fumes	Silica, amorphous fume	45 mg	g/m3	500 mg/m3	3,000 mg/m3	
ozone	Ozone	0.24 p	opm	1 ppm	10 ppm	
				•	·	
Ingredient	Original IDLH		Revised ID	LH		
iron	500 mg/m3	500 mg/m3		Not Available		
titanium dioxide	5,000 mg/m3	5,000 mg/m3		Available		
manganese	500 mg/m3		Not Available			
kaolin, calcined	Not Available		Not Availabl	Not Available		
silicon	Not Available		Not Available			
manganese fume	500 mg/m3	500 mg/m3		Not Available		
fluoride fume	Not Available	Not Available		Available Not Available		
silica welding fumes	Not Available	Not Available		Not Available		
ozone	5 ppm	5 ppm		Not Available		
nitrogen oxides	Not Available		Not Availabl	e		

#### MATERIAL DATA

# Exposure controls

	Engineering controls are used to remove a hazard or place a barrier betw engineering controls can be highly effective in protecting workers and will 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 Enclosure and/or isolation of emission source which keeps a selected ha ventilation that strategically "adds" and "removes" air in the work environ contaminant if designed properly. The design of a ventilation system musi- contaminant in use. Employers may need to use multiple types of controls to prevent employee Local exhaust ventilation usually required. If risk of overexposure exis to obtain adequate protection. Supplied-air type respirator may be required essential to ensure adequate protection.	I typically be independent of v s is done to reduce the risk. zard "physically" away from the ment. Ventilation can remove st match the particular process see overexposure. ts, wear approved respirator	worker interactions to he worker and or dilute an air s and chemical or
	An approved self contained breathing apparatus (SCBA) may be required	d in some situations.	
	Provide adequate ventilation in warehouse or closed storage area. Air co	•	
	varying "escape" velocities which, in turn, determine the "capture velocitie remove the contaminant.	es or mesh circulating all requ	uneu lo eneclively
	Type of Contaminant:		Air Speed:
ppropriate engineering		0.25-0.5 m/s	
	solvent, vapours, degreasing etc., evaporating from tank (in still air).		(50-100 f/min.)
oriate engineering controls	solvent, vapours, degreasing etc., evaporating from tank (in still air). aerosols, fumes from pouring operations, intermittent container filling, lo transfers, welding, spray drift, plating acid fumes, pickling (released at lo active generation)		
	aerosols, fumes from pouring operations, intermittent container filling, lo transfers, welding, spray drift, plating acid fumes, pickling (released at lo	ow velocity into zone of	(50-100 f/min.) 0.5-1 m/s
	aerosols, fumes from pouring operations, intermittent container filling, lo transfers, welding, spray drift, plating acid fumes, pickling (released at lo active generation) direct spray, spray painting in shallow booths, drum filling, conveyer loa	ow velocity into zone of ding, crusher dusts, gas	(50-100 f/min.) 0.5-1 m/s (100-200 f/min.) 1-2.5 m/s
	aerosols, fumes from pouring operations, intermittent container filling, lo transfers, welding, spray drift, plating acid fumes, pickling (released at lo active generation) direct spray, spray painting in shallow booths, drum filling, conveyer load discharge (active generation into zone of rapid air motion) grinding, abrasive blasting, tumbling, high speed wheel generated dusts	ow velocity into zone of ding, crusher dusts, gas	(50-100 f/min.) 0.5-1 m/s (100-200 f/min.) 1-2.5 m/s (200-500 f/min.) 2.5-10 m/s
	aerosols, fumes from pouring operations, intermittent container filling, lo transfers, welding, spray drift, plating acid fumes, pickling (released at lo active generation) direct spray, spray painting in shallow booths, drum filling, conveyer load discharge (active generation into zone of rapid air motion) grinding, abrasive blasting, tumbling, high speed wheel generated dusts velocity into zone of very high rapid air motion).	ow velocity into zone of ding, crusher dusts, gas	(50-100 f/min.) 0.5-1 m/s (100-200 f/min.) 1-2.5 m/s (200-500 f/min.) 2.5-10 m/s (500-2000 f/min
	aerosols, fumes from pouring operations, intermittent container filling, lo transfers, welding, spray drift, plating acid fumes, pickling (released at la active generation) direct spray, spray painting in shallow booths, drum filling, conveyer load discharge (active generation into zone of rapid air motion) grinding, abrasive blasting, tumbling, high speed wheel generated dusts velocity into zone of very high rapid air motion). Within each range the appropriate value depends on:	ow velocity into zone of ding, crusher dusts, gas s (released at high initial	(50-100 f/min.) 0.5-1 m/s (100-200 f/min.) 1-2.5 m/s (200-500 f/min.) 2.5-10 m/s (500-2000 f/min
	aerosols, fumes from pouring operations, intermittent container filling, lo transfers, welding, spray drift, plating acid fumes, pickling (released at lo active generation) direct spray, spray painting in shallow booths, drum filling, conveyer load discharge (active generation into zone of rapid air motion) grinding, abrasive blasting, tumbling, high speed wheel generated dusts velocity into zone of very high rapid air motion). Within each range the appropriate value depends on: Lower end of the range	bw velocity into zone of ding, crusher dusts, gas s (released at high initial Upper end of the rar	(50-100 f/min.) 0.5-1 m/s (100-200 f/min.) 1-2.5 m/s (200-500 f/min.) 2.5-10 m/s (500-2000 f/min
	aerosols, fumes from pouring operations, intermittent container filling, lo transfers, welding, spray drift, plating acid fumes, pickling (released at la active generation) direct spray, spray painting in shallow booths, drum filling, conveyer load discharge (active generation into zone of rapid air motion) grinding, abrasive blasting, tumbling, high speed wheel generated dusts velocity into zone of very high rapid air motion). Within each range the appropriate value depends on: Lower end of the range 1: Room air currents minimal or favourable to capture	bw velocity into zone of ding, crusher dusts, gas s (released at high initial Upper end of the rar 1: Disturbing room a	(50-100 f/min.) 0.5-1 m/s (100-200 f/min.) 1-2.5 m/s (200-500 f/min.) 2.5-10 m/s (500-2000 f/min nge ir currents nigh toxicity

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Other protection	Overalls. P.V.C. apron. Barrier cream. Skin cleansing cream. Eye wash unit.
Body protection	See Other protection below
Hands/feet protection	Wear chemical protective gloves, e.g. PVC. Wear safety footwear or safety gumboots, e.g. Rubber
Skin protection	See Hand protection below
Eye and face protection	Safety glasses with side shields Chemical goggles. 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	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.
	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

### **Respiratory protection**

Type NO Filter of sufficient capacity. (AS/NZS 1716 & 1715, EN 143:2000 & 149:2001, ANSI Z88 or national equivalent)

Where the concentration of gas/particulates in the breathing zone, approaches or exceeds the "Exposure Standard" (or ES), respiratory protection is required.

Degree of protection varies with both face-piece and Class of filter; the nature of protection varies with Type of filter.

Required Minimum Protection Factor	Half-Face Respirator	Full-Face Respirator	Powered Air Respirator
up to 10 x ES	NO-AUS	-	NO-PAPR-AUS / Class 1
up to 50 x ES	-	NO-AUS / Class 1	-
up to 100 x ES	-	NO-2	NO-PAPR-2 ^

#### ^ - Full-face

A(All classes) = Organic vapours, B AUS or B1 = Acid gasses, B2 = Acid gas or hydrogen cyanide(HCN), B3 = Acid gas or hydrogen cyanide(HCN), E = Sulfur dioxide(SO2), G = Agricultural chemicals, K = Ammonia(NH3), Hg = Mercury, NO = Oxides of nitrogen, MB = Methyl bromide, AX = Low boiling point organic compounds(below 65 degC)

### SECTION 9 PHYSICAL AND CHEMICAL PROPERTIES

#### Information on basic physical and chemical properties

Appearance	Shine silver electrode; insoluble in water.		
Physical state	Manufactured	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 Applicable
pH (as supplied)	Not Applicable	Decomposition temperature	Not Available
Melting point / freezing point (°C)	Not Available	Viscosity (cSt)	Not Applicable
Initial boiling point and boiling range (°C)	Not Applicable	Molecular weight (g/mol)	Not Applicable
Flash point (°C)	Not Applicable	Taste	Not Available

Evaporation rate	Not Applicable	Explosive properties	Not Available
Flammability	Not Applicable	Oxidising properties	Not Available
Upper Explosive Limit (%)	Not Applicable	Surface Tension (dyn/cm or mN/m)	Not Applicable
Lower Explosive Limit (%)	Not Applicable	Volatile Component (%vol)	Not Applicable
Vapour pressure (kPa)	Not Applicable	Gas group	Not Available
Solubility in water	Immiscible	pH as a solution (1%)	Not Applicable
Vapour density (Air = 1)	Not Applicable	VOC g/L	Not Available

# SECTION 10 STABILITY AND REACTIVITY

Reactivity	See section 7
Chemical stability	Product is considered stable and hazardous polymerisation will not occur.
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	Fumes evolved during welding operations may be irritating to the upper-respiratory tract and may be harmful if inhaled.
Ingestion	The material has <b>NOT</b> 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. The material may still be damaging to the health of the individual, following ingestion, especially where pre-existing organ (e.g liver, kidney) damage is evident. Present definitions of harmful or toxic substances are generally based on doses producing mortality rather than those producing morbidity (disease, ill-health). Gastrointestinal tract discomfort may produce nausea and vomiting. In an occupational setting however, ingestion of insignificant quantities is not thought to be cause for concern.
Skin Contact	Skin contact does <b>not</b> normally present a hazard, though it is always possible that occasionally individuals may be found who react to substances usually regarded as inert.
Eye	Fumes from welding/brazing operations may be irritating to the eyes.
Chronic	Limited evidence suggests that repeated or long-term occupational exposure may produce cumulative health effects involving organs or biochemical systems. Principal route of exposure is inhalation of welding fumes from electrodes and workpiece. Reaction products arising from electrode core and flux appear as welding fume depending on welding conditions, relative volatilities of metal oxides and any coatings on the workpiece. Studies of lung cancer among welders indicate that they may experience a 30-40% increased risk compared to the general population. Since smoking and exposure to other cancer-causing agents, such as absetsos fibre, may influence these results, it is not clear whether welding, in fact, represents a significant lung cancer risk. Whilst mild steel welding represents little risk, the stainless steel welder, exposed to chromium and nickel fume, may be at risk and it is this factor which may account for the overall increase in lung cancer incidence among welders. Cold isolated electrodes are relatively harmless. Welding fume with high levels of ferrous materials may lead to particle deposition in the lungs (siderosis) after long exposure. This clears up when exposure stops. Chronic exposure to iron dusts may lead to eye disorders. Chronic excessive iron exposure has been associated with haemosiderosis and consequent possible damage to the liver and pancreas. Haemosiderin is golden-brown insoluble protein produced by phagocytic digestion of haematin (an iron-based pigment). Haemosiderin is found in most tissues, especially in the liver, in the form of granules. Other sites of haedosids of the skin - heart failure may eventually occur. Such exposure may also produce conjunctivitis, choroiditis, retinitis (both inflammatory conditions involving the eye) and siderosis of tissues if iron remains in these tissues. Siderosis is a form of pneumoconicsis produced by iron dusts. Siderosis also includes discoloration of organs, excess circulating iron and degeneration of the tenina, lens and uvea as a re

[K. Schmidt, New Scientist, No. 1919 pp.11-12, 2nd April, 1994]

Metallic dusts generated by the industrial process give rise to a number of potential health problems. The larger particles, above 5 micron, are nose and throat irritants. Smaller particles however, may cause lung deterioration. Particles of less than 1.5 micron can be trapped in the lungs and, dependent on the nature of the particle, may give rise to further serious health consequences.

Metals are widely distributed in the environment and are not biodegradable. Biologically, many metals are essential to living systems and are involved in a variety of cellular, physiological, and structural functions. They often are cofactors of enzymes, and play a role in transcriptional control, muscle contraction, nerve transmission, blood clotting, and oxygen transport and delivery. Although all metals are potentially toxic at some level, some are highly toxic at relatively low levels. Moreover, in some cases the same metal can be essential at low levels and toxic at higher levels, or it may be toxic via one route of entry but not another. Toxic effects of some metals are associated with disruption of functions of essential metals. Metals may have a range of effects, including cancer, neurotoxicity, immunotoxicity, cardiotoxicity, reproductive toxicity, teratogenicity, and genotoxicity. Biological half lives of metals vary greatly, from hours to years. Furthermore, the half life of a given metal varies in different tissues. Lead has a half life of 14 days in soft tissues and 20 years in bone.

In considering how to evaluate the toxicity of metals of potential concern, a number of aspects of metal toxicity should be kept in mind:

Different species vary in their responses to different metals; in some cases, humans are more sensitive than rodents. Thus, there is a need for broad-based testing of metals;

- > The route of exposure may affect the dose and site where the metal concentrates, and thus the observed toxic effects;
- Metal-metal interactions can reduce or enhance toxicity; biotransformation can reduce or enhance toxicity;
- It is difficult to predict the toxicity of one metal based on the adverse effects of another; in trying to evaluate the toxicity of one particular metal compound, predictions based on similar compounds of the same metal may be valid.

Ozone is suspected to produce lung cancer in laboratory animals; no reports of this effect have been documented in exposed human populations.

Other welding process exposures can arise from radiant energy UV flash burns, thermal burns or electric shock The welding arc emits ultraviolet radiation at wavelengths that have the potential to produce skin tumours in animals and in over-exposed individuals, however, no confirmatory studies of this effect in welders have been reported.

	тохісіту	IRRITATION
WIA Austfil 71T	Not Available	Not Available
	тохісіту	IRRITATION
iron	Oral (rat) LD50: 750 mg/kg <sup>[2]</sup>	Not Available
	тохісіту	IRRITATION
	dermal (hamster) LD50: >=10000 mg/kg <sup>[2]</sup>	Eye: no adverse effect observed (not irritating) <sup>[1]</sup>
titanium dioxide	Oral (rat) LD50: >2000 mg/kg <sup>[1]</sup>	Skin (human): 0.3 mg /3D (int)-mild *
		Skin: no adverse effect observed (not irritating) <sup>[1]</sup>
	тохісітү	IRRITATION
	Oral (rat) LD50: >2000 mg/kg <sup>[1]</sup>	Eye (rabbit): 500 mg/24h - mild
manganese		Eye: no adverse effect observed (not irritating) <sup>[1]</sup>
		Skin (rabbit): 500 mg/24h - mild
		Skin: no adverse effect observed (not irritating) <sup>[1]</sup>
	тохісіту	IRRITATION
kaolin, calcined	dermal (rat) LD50: >5000 mg/kg <sup>[1]</sup>	Not Available
	Oral (rat) LD50: >2000 mg/kg <sup>[1]</sup>	
	тохісіту	IRRITATION
silicon	Oral (rat) LD50: >50-300 mg/kg <sup>[1]</sup>	Eye: no adverse effect observed (not irritating) <sup>[1]</sup>
		Skin: no adverse effect observed (not irritating) <sup>[1]</sup>
	тохісіту	IRRITATION
	Oral (rat) LD50: >2000 mg/kg <sup>[1]</sup>	Eye (rabbit) 500mg/24H Mild
manganese fume		Eye: no adverse effect observed (not irritating) <sup>[1]</sup>
		Skin (rabbit) 500mg/24H Mild
		Skin: no adverse effect observed (not irritating) <sup>[1]</sup>

	TOXICITY	IRRITATION
fluoride fume	Not Available	Not Available
	ΤΟΧΙΟΙΤΥ	IRRITATION
silica welding fumes	>5000 mg/kg <sup>[1]</sup>	Eye: no adverse effect observed (not irritating) <sup>[1]</sup>
	Oral (rat) LD50: 3160 mg/kg <sup>[2]</sup>	Skin: no adverse effect observed (not irritating) <sup>[1]</sup>
	ΤΟΧΙΟΙΤΥ	IRITATION
ozone	Inhalation (rat) LC50: 0.001 mg/l/44H <sup>[2]</sup>	Eye: adverse effect observed (irreversible damage) <sup>[1]</sup>
		Skin: adverse effect observed (corrosive) <sup>[1]</sup>
	ΤΟΧΙGITY	IRRITATION
nitrogen oxides	Not Available	Not Available
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	

The material may produce moderate eye irritation leading to inflammation. Repeated or prolonged exposure to irritants may produce conjunctivitis.

For titanium dioxide:

TITANIUM DIOXIDE

Humans can be exposed to titanium dioxide via inhalation, ingestion or dermal contact. In human lungs, the clearance kinetics of titanium dioxide is poorly characterized relative to that in experimental animals. (General particle characteristics and host factors that are considered to affect deposition and retention patterns of inhaled, poorly soluble particles such as titanium dioxide are summarized in the monograph on carbon black.) With regard to inhaled titanium dioxide, human data are mainly available from case reports that showed deposits of titanium dioxide in lung tissue as well as in lymph nodes. A single clinical study of oral ingestion of fine titanium dioxide showed particle size-dependent absorption by the gastrointestinal tract and large interindividual variations in blood levels of titanium dioxide. Studies on the application of sunscreens containing ultrafine titanium dioxide to healthy skin of human volunteers revealed that titanium dioxide particles only penetrate into the outermost layers of the stratum corneum, suggesting that healthy skin is an effective barrier to titanium dioxide. There are no studies on penetration of titanium dioxide in compromised skin. Respiratory effects that have been observed among groups of titanium dioxide-exposed workers include decline in lung function, pleural disease with plaques and pleural thickening, and mild fibrotic changes. However, the workers in these studies were also exposed to asbestos and/or silica.

No data were available on genotoxic effects in titanium dioxide-exposed humans.

Many data on deposition, retention and clearance of titanium dioxide in experimental animals are available for the inhalation route. Titanium dioxide inhalation studies showed differences — both for normalized pulmonary burden (deposited mass per dry lung, mass per body weight) and clearance kinetics — among rodent species including rats of different size, age and strain. Clearance of titanium dioxide is also affected by pre-exposure to gaseous pollutants or co-exposure to cytotoxic aerosols. Differences in dose rate or clearance kinetics and the appearance of focal areas of high particle burden have been implicated in the higher toxic and inflammatory lung responses to intratracheally instilled vs inhaled titanium dioxide particles. Experimental studies with titanium dioxide have demonstrated that rodents experience dose-dependent impairment of alveolar macrophage-mediated clearance. Hamsters have the most efficient clearance of inhaled titanium dioxide. Ultrafine primary particles of titanium dioxide are more slowly cleared than their fine counterparts.

Titanium dioxide causes varying degrees of inflammation and associated pulmonary effects including lung epithelial cell injury, cholesterol granulomas and fibrosis. Rodents experience stronger pulmonary effects after exposure to ultrafine titanium dioxide particles compared with fine particles on a mass basis. These differences are related to lung burden in terms of particle surface area, and are considered to result from impaired phagocytosis and sequestration of ultrafine particles into the interstitium.

Fine titanium dioxide particles show minimal cytotoxicity to and inflammatory/pro-fibrotic mediator release from primary human alveolar macrophages in vitro compared with other particles. Ultrafine titanium dioxide particles inhibit phagocytosis of alveolar macrophages in vitro at mass dose concentrations at which this effect does not occur with fine titanium dioxide. In-vitro studies with fine and ultrafine titanium dioxide and purified DNA show induction of DNA damage that is suggestive of the generation of reactive oxygen species by both particle types. This effect is stronger for ultrafine than for fine titanium oxide, and is markedly enhanced by exposure to simulated sunlight/ultraviolet light. **Animal carcinogenicity data** 

Pigmentary and ultrafine titanium dioxide were tested for carcinogenicity by oral administration in mice and rats, by inhalation in rats and female mice, by intratracheal administration in hamsters and female rats and mice, by subcutaneous injection in rats and by intraperitoneal administration in male mice and female rats.

In one inhalation study, the incidence of benign and malignant lung tumours was increased in female rats. In another inhalation study, the incidences of lung adenomas were increased in the high-dose groups of male and female rats. Cystic keratinizing lesions that were diagnosed as squamous-cell carcinomas but re-evaluated as non-neoplastic pulmonary keratinizing cysts were also observed in the high-dose groups of female rats. Two inhalation studies in rats and one in female mice were negative.

Intratracheally instilled female rats showed an increased incidence of both benign and malignant lung tumours following treatment with two types of titanium dioxide. Tumour incidence was not increased in intratracheally instilled hamsters and female mice.

In-vivo studies have shown enhanced micronucleus formation in bone marrow and peripheral blood lymphocytes of

	intraperitoneally instilled mice. Increased Hprt mutations were seen in lung epithelial cells isolated from titanium dioxide- instilled rats. In another study, no enhanced oxidative DNA damage was observed in lung tissues of rats that were intratracheally instilled with titanium dioxide. The results of most in-vitro genotoxicity studies with titanium dioxide were negative.
	WARNING: This substance has been classified by the IARC as Group 2B: Possibly Carcinogenic to Humans. * IUCLID
SILICON	Intraperitoneal injection of silicon produced only minor local trauma and foreign body reaction. Parenterally administered elemental silica is considered biologically inert. Dogs and rats fed 800 mg silicon/kg/day (as the dioxide) for 1 month showed no clinical signs or histological changes. The compound was largely eliminated in the faeces. Normal human cerebral cortex tissue contains about 3.8 ug/g silicon
SILICA WELDING FUMES	For silica amorphous: When experimental animals inhale synthetic amorphous silica (SAS) dust, it dissolves in the lung fluid and is rapidly eliminated. If swallowed, the vast majority of SAS is excreted in the faeces and there is little accumulation in the body. Following absorption across the gut, SAS is eliminated via urine without modification in animals and humans. SAS is not expected to be broken down (metabolised) in mammals. After ingestion, there is limited accumulation of SAS in body tissues and rapid elimination occurs. Intestinal absorption has not been calculated, but appears to be insignificant in animals and humans. SASs injected subcutaneously are subjected to rapid dissolution and removal. There is no indication of metabolism of SAS in animals on humans based on chemical structure and available data. In contrast to crystalline silica, SAS is soluble in physiological media and the soluble chemical species that are formed are eliminated via the urinary tract without modification. Both the mammalian and environmental toxicology of SASs are significantly influenced by the physical and chemical properties, particularly those of solubility and particle size. SAS has no acute intrinsic toxicity by inhalation. Adverse effects, including suffocation, that have been reported were caused by the presence of high numbers of respirable particles generated to meet the required test atmosphere. These results are not representative of exposure to commercial SASs and should not be used for human risk assessment. Though repeated exposure of the skin may cause dryness and cracking, SAS is not a skin or eye irritant, and it is not a sensitier. Repeated-dose, subchronic and chronic inhalation toxicity studies have been conducted with SAS in a number of species, at altorne concentrations ranging from 0.5 mg/m3 to 150 mg/m3. Lowest-observed adverse effect levels (NOAELs) were between 0.5 and 10 mg/m3. The difference in values may be explained by different particle size, and therefore the number of par
OZONE	NOTE: Ozone aggravates chronic obstructive pulmonary diseases. Ozone is suspected also of increasing the risk of acute and chronic respiratory disease, mutagenesis and foetotoxicity. In animals short-term exposure to ambient concentrations of less than 1 ppm results in reduced capacity to kill intrapulmonary organisms and allows purulent bacteria to proliferate [Ellenhorn etal].
nitrogen oxides	Data for nitrogen dioxide: Substance has been investigated as a mutagen and reproductive effector. NOTE: Interstitial edema, epithelial proliferation and, in high concentrations, fibrosis and emphysema develop after repeated exposure.
TITANIUM DIOXIDE& MANGANESE	The material may cause skin irritation after prolonged or repeated exposure and may produce a contact dermatitis (nonallergic). This form of dermatitis is often characterised by skin redness (erythema) and swelling epidermis. Histologically there may be intercellular oedema of the spongy layer (spongiosis) and intracellular oedema of the epidermis.
MANGANESE & SILICON	The material may be irritating to the eye, with prolonged contact causing inflammation. Repeated or prolonged exposure to irritants may produce conjunctivitis.
KAOLIN, CALCINED & SILICON	No significant acute toxicological data identified in literature search.
SILICON & OZONE & nitrogen oxides	Asthma-like symptoms may continue for months or even years after exposure to the material ceases. This may be due to a non-allergenic condition known as reactive airways dysfunction syndrome (RADS) which can occur following exposure to high levels of highly irritating compound. Key criteria for the diagnosis of RADS include the absence of preceding

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respiratory disease, in a non-atopic individual, with abrupt onset of persistent asthma-like symptoms within minutes to hours of a documented exposure to the irritant. A reversible airflow pattern, on spirometry, with the presence of moderate to severe bronchial hyperreactivity on methacholine challenge testing and the lack of minimal lymphocytic inflammation, without eosinophilia, have also been included in the criteria for diagnosis of RADS. RADS (or asthma) following an irritating inhalation is an infrequent disorder with rates related to the concentration of and duration of exposure to the irritating substance. Industrial bronchitis, on the other hand, is a disorder that occurs as result of exposure due to high concentrations of irritating substance (often particulate in nature) and is completely reversible after exposure ceases. The disorder is characterised by dyspnea, cough and mucus production.

×	Carcinogenicity	×
×	Reproductivity	×
×	STOT - Single Exposure	×
×	STOT - Repeated Exposure	×
×	Aspiration Hazard	×
:	× × ×	X     Reproductivity       X     STOT - Single Exposure       X     STOT - Repeated Exposure

Legend:

Data either not available or does not fill the criteria for classification
 Data available to make classification

# SECTION 12 ECOLOGICAL INFORMATION

#### Toxicity

	ENDPOINT	TEST DURATION (HR)	SPECIES	VALUE	SOURCE
WIA Austfil 71T	Not Available	Not Available	Not Available	Not Available	Not Available
	ENDPOINT	TEST DURATION (HR)	SPECIES	VALUE	SOURCE
	LC50	96	Fish	0.05mg/L	2
	EC50	48	Crustacea	5.11mg/L	2
iron	EC50	96	Algae or other aquatic plants	3.7mg/L	4
	BCF	24	Crustacea	0.0000002mg/L	4
	NOEC	504	Fish	0.52mg/L	2
	ENDPOINT	TEST DURATION (HR)	SPECIES	VALUE	SOURCE
	LC50	96	Fish	>1-mg/L	2
titanium dioxide	EC50	48	Crustacea	>1-mg/L	2
	EC50	72	Algae or other aquatic plants	Algae or other aquatic plants 5.83mg/L	
	NOEC	336	Fish	0.089mg/L	4
	ENDPOINT	TEST DURATION (HR)	SPECIES VALUE		SOURCE
	LC50	96	Fish	>3.6mg/L	2
	EC50	48	Crustacea	>1.6mg/L	2
manganese	EC50	72	Algae or other aquatic plants 2.8mg/L		2
	BCFD	37	Algae or other aquatic plants 2.2mg/L		4
	NOEC	48	Crustacea	1.6mg/L	2
	ENDPOINT	TEST DURATION (HR)	SPECIES	VALUE	SOURCI
	LC50	96	Fish >100mg/L		2
	EC50	48	Crustacea >100mg/L		2
kaolin, calcined	EC50	72	Algae or other aquatic plants	Algae or other aquatic plants 2-500mg/L	
	EC10	72	Algae or other aquatic plants 33mg/L		2
	NOEC	504	Crustacea	1-mg/L	2
	ENDPOINT	TEST DURATION (HR)	SPECIES	VALUE	SOURCI
- 11	EC50	48	Crustacea	ca.35.4mg/L	2
silicon	EC50	72	Algae or other aquatic plants	>100mg/L	2
	NOEC	72	Algae or other aquatic plants	ca.3.2mg/L	2

	ENDPOINT	TEST DURATION (HR)	SPECIES	VALUE	SOURCE
	LC50	96	Fish	>3.6mg/L	2
	EC50	48	Crustacea	Crustacea >1.6mg/L	
manganese fume	EC50	72	Algae or other aquatic plants	Algae or other aquatic plants 2.8mg/L	
	BCFD	37	Algae or other aquatic plants	2.2mg/L	4
	NOEC	48	Crustacea	1.6mg/L	2
	ENDPOINT	TEST DURATION (HR)	SPECIES	VALUE	SOURCE
	LC50	96	Fish	63.215mg/L	3
fluoride fume	EC50	48	Crustacea	36.2mg/L	5
	EC50	96	Algae or other aquatic plants	347.670mg/L	3
	NOEC	504	Crustacea	14mg/L	5
	ENDPOINT	TEST DURATION (HR)	SPECIES	VALUE	SOURCE
	LC50	96	Fish	>100mg/L	2
silica welding fumes	EC50	72	Algae or other aquatic plants	4-200mg/L	2
	NOEC	504	Crustacea	100mg/L	2
	ENDPOINT	TEST DURATION (HR)	SPECIES	VALUE	SOURCI
ozone	LC50	96	Fish	0.0093mg/L	2
	NOEC	2160	Fish	0.002mg/L	5
	ENDPOINT	TEST DURATION (HR)	SPECIES	VALUE	SOURCE
nitrogen oxides	Not Available	Not Available	Not Available	Not Available	Not Available
Legend:	Toxicity 3. EP Data 5. ECE1	NWIN Suite V3.12 (QSAR) - Aqua	pe ECHA Registered Substances - Ecotoxic tic Toxicity Data (Estimated) 4. US EPA, Eco Data 6. NITE (Japan) - Bioconcentration Dat	otox database - Aquatio	

# DO NOT discharge into sewer or waterways.

# Persistence and degradability

Ingredient	Persistence: Water/Soil	Persistence: Air
titanium dioxide	HIGH	HIGH
fluoride fume	LOW	LOW

### **Bioaccumulative potential**

Ingredient	Bioaccumulation	
titanium dioxide	LOW (BCF = 10)	
fluoride fume	LOW (LogKOW = 0.2259)	

# Mobility in soil

Ingredient	Mobility
titanium dioxide	LOW (KOC = 23.74)
fluoride fume	LOW (KOC = 14.3)

### SECTION 13 DISPOSAL CONSIDERATIONS

#### Waste treatment methods

<ul> <li>Containers may still present a chemical hazar</li> </ul>	d/ danger when empty.
---	-----------------------

Product / Packaging disposal

- Return to supplier for reuse/ recycling if possible.
   Otherwise:
- If container can not be cleaned sufficiently well to ensure that residuals do not remain or if the container cannot be used to store the same product, then puncture containers, to prevent re-use, and bury at an authorised landfill.

+ Where possible retain label warnings and SDS and observe all notices pertaining to the product.

# SECTION 14 TRANSPORT INFORMATION

#### Labels Required

Marine Pollutant	NO
HAZCHEM	Not Applicable

# Land transport (ADG): NOT REGULATED FOR TRANSPORT OF DANGEROUS GOODS

### Air transport (ICAO-IATA / DGR): NOT REGULATED FOR TRANSPORT OF DANGEROUS GOODS

# Sea transport (IMDG-Code / GGVSee): NOT REGULATED FOR TRANSPORT OF DANGEROUS GOODS

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

Not Applicable

### **SECTION 15 REGULATORY INFORMATION**

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

#### IRON(7439-89-6) IS FOUND ON THE FOLLOWING REGULATORY LISTS

Australia Inventory of Chemical Substances (AICS)

#### TITANIUM DIOXIDE(13463-67-7) IS FOUND ON THE FOLLOWING REGULATORY LISTS

Australia Exposure Standards	IMO MARPOL (Annex II) - List of Noxious Liquid Substances Carried in
Australia Inventory of Chemical Substances (AICS)	Bulk
GESAMP/EHS Composite List - GESAMP Hazard Profiles	International Agency for Research on Cancer (IARC) - Agents Classified
IMO IBC Code Chapter 17: Summary of minimum requirements	by the IARC Monographs
	International WHO List of Proposed Occupational Exposure Limit (OEL)
	Values for Manufactured Nanomaterials (MNMS)
MANGANESE(7439-96-5) IS FOUND ON THE FOLLOWING REGULATORY LIS	STS
Australia Dangerous Goods Code (ADG Code) - Dangerous Goods List	Australia Inventory of Chemical Substances (AICS)
Australia Dangerous Goods Code (ADG Code) - List of Emergency Action	International Air Transport Association (IATA) Dangerous Goods Regulations
Codes	International Maritime Dangerous Goods Requirements (IMDG Code)
Australia Exposure Standards	United Nations Recommendations on the Transport of Dangerous Goods
Australia Hazardous Chemical Information System (HCIS) - Hazardous Chemicals	Model Regulations
KAOLIN, CALCINED(92704-41-1) IS FOUND ON THE FOLLOWING REGULAT	ORY LISTS
Australia Inventory of Chemical Substances (AICS)	
SILICON(7440-21-3) IS FOUND ON THE FOLLOWING REGULATORY LISTS	
Australia Exposure Standards	Australia Inventory of Chemical Substances (AICS)
MANGANESE FUME(7439-96-5.) IS FOUND ON THE FOLLOWING REGULAT	DRY LISTS
Australia Exposure Standards	Australia Inventory of Chemical Substances (AICS)
Australia Hazardous Chemical Information System (HCIS) - Hazardous	
Chemicals	
FLUORIDE FUME(16984-48-8) IS FOUND ON THE FOLLOWING REGULATOR	Y LISTS
International Agency for Research on Cancer (IARC) - Agents Classified	
by the IARC Monographs	
	ULATORY LISTS
SILICA WELDING FUMES(69012-64-2) IS FOUND ON THE FOLLOWING REGI	
Australia Hazardous Chemical Information System (HCIS) - Hazardous	Australia Inventory of Chemical Substances (AICS)
Chemicals	

OZONE(10028-15-6) IS FOUND ON THE FOLLOWING REGULATORY LISTS

Australia Dangerous Goods Code (ADG Code) - Dangerous Goods List	International Air Transport Association (IATA) Dangerous Goods Regulations	
Australia Dangerous Goods Code (ADG Code) - List of Emergency Action	International Maritime Dangerous Goods Requirements (IMDG Code)	
Codes	United Nations Recommendations on the Transport of Dangerous Goods	
Australia Dangerous Goods Code (ADG Code) - Packing Instruction -	Model Regulations	
Compressed Gases		
Australia Exposure Standards		

# NITROGEN OXIDES(MIXTURE) IS FOUND ON THE FOLLOWING REGULATORY LISTS

Australia Dangerous Goods Code (ADG Code) - Dangerous Goods List	International Air Transport Association (IATA) Dangerous Goods Regulations
Australia Dangerous Goods Code (ADG Code) - List of Emergency Action	- Prohibited List Passenger and Cargo Aircraft
Codes	International Maritime Dangerous Goods Requirements (IMDG Code)
Australia Dangerous Goods Code (ADG Code) - Packing Instruction -	United Nations Recommendations on the Transport of Dangerous Goods
Liquefied and Dissolved Gases	Model Regulations
International Air Transport Association (IATA) Dangerous Goods Regulations	

### **National Inventory Status**

National Inventory	Status
Australia - AICS	No (fluoride fume; ozone; nitrogen oxides)
Canada - DSL	No (ozone; nitrogen oxides)
Canada - NDSL	No (manganese fume; fluoride fume; silica welding fumes; manganese; silicon; nitrogen oxides; kaolin, calcined; iron)
China - IECSC	No (nitrogen oxides)
Europe - EINEC / ELINCS / NLP	No (fluoride fume; nitrogen oxides)
Japan - ENCS	No (manganese fume; fluoride fume; ozone; manganese; silicon; nitrogen oxides; kaolin, calcined; iron)
Korea - KECI	No (fluoride fume; nitrogen oxides)
New Zealand - NZloC	No (nitrogen oxides)
Philippines - PICCS	No (ozone; nitrogen oxides)
USA - TSCA	No (fluoride fume; nitrogen oxides)
Taiwan - TCSI	No (nitrogen oxides)
Mexico - INSQ	No (silica welding fumes; nitrogen oxides)
Vietnam - NCI	No (nitrogen oxides)
Russia - ARIPS	No (fluoride fume; silica welding fumes; nitrogen oxides)
Thailand - TECI	No (silica welding fumes; nitrogen oxides; kaolin, calcined; iron)
Legend:	Yes = All CAS declared ingredients are on the inventory No = 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**

Revision Date	27/06/2017
Initial Date	Not Available

### Other information

# Ingredients with multiple cas numbers

Name	CAS No
titanium dioxide	13463-67-7, 1317-70-0, 1317-80-2, 12188-41-9, 1309-63-3, 100292-32-8, 101239-53-6, 116788-85-3, 12000-59-8, 12701-76-7, 12767-65-6, 12789-63-8, 1344-29-2, 185323-71-1, 185828-91-5, 188357-76-8, 188357-79-1, 195740-11-5, 221548-98-7, 224963-00-2, 246178-32-5, 252962-41-7, 37230-92-5, 37230-94-7, 37230-95-8, 37230-96-9, 39320-58-6, 39360-64-0, 39379-02-7, 416845-43-7, 494848-07-6, 494848-23-6, 494851-77-3, 494851-98-8, 55068-84-3, 55068-85-4, 552316-51-5, 62338-64-1, 767341-00-4, 97929-50-5, 98084-96-9
kaolin, calcined	92704-41-1, 39388-40-4, 66402-68-4
silicon	7440-21-3, 152284-21-4, 157383-37-4, 160371-18-6, 17375-03-0, 71536-23-7, 72516-01-9, 72516-02-0, 72516-03-1, 90337-93-2

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Version No: 3.1.1.1	Vortex 71T	Print Date: 20/06/2019

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

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

#### **Definitions and abbreviations**

PC – TWA: Permissible Concentration-Time Weighted Average PC – STEL: Permissible Concentration-Short Term Exposure Limit IARC: International Agency for Research on Cancer ACGIH: American Conference of Governmental Industrial Hygienists STEL: Short Term Exposure Limit TEEL: Temporary Emergency Exposure Limit<sub>o</sub> 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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