

# **Apollo Scientific**

Part Number: **OR930278** Version No: **1.1** Safety Data Sheet (Conforms to Annex II of REACH (1907/2006) - Regulation 2020/878) Chemwatch Hazard Alert Code: 3

Issue Date: **16/05/2022** Print Date: **03/08/2023** S.REACH.GBR.EN

### SECTION 1 Identification of the substance / mixture and of the company / undertaking

### **1.1. Product Identifier**

Product name	tert-Butyl-dimethyl-(tributylstannylmethoxy)silane		
Chemical Name	methyl(2-methyl-2- propanyl)[(tributylstannyl)methoxy]silane		
Synonyms	Not Available		
Proper shipping name	ORGANOTIN COMPOUND, LIQUID, N.O.S.		
Chemical formula	Not Available		
Other means of identification	Not Available		
CAS number	123061-64-3*		

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

Relevant identified uses	Not Available	
Uses advised against	No specific uses advised against are identified.	

### 1.3. Details of the manufacturer or supplier of the safety data sheet

Registered company name	Apollo Scientific		
Address	hitefield Road, Bredbury SK62QR United Kingdom		
Telephone	4060505		
Fax	0161 406 0506		
Website	http://www.apolloscientific.co.uk/		
Email	sales@apolloscientific.co.uk		

### 1.4. Emergency telephone number

Association / Organisation	Not Available
Emergency telephone numbers	Not Available
Other emergency telephone numbers	Not Available

### **SECTION 2 Hazards identification**

### 2.1. Classification of the substance or mixture

Classification according to regulation (EC) No 1272/2008 [CLP] and amendments <sup>[1]</sup>

H312 - Acute Toxicity (Dermal) Category 4, H332 - Acute Toxicity (Inhalation) Category 4, H335 - Specific Target Organ Toxicity -Single Exposure (Respiratory Tract Irritation) Category 3, H315 - Skin Corrosion/Irritation Category 2, H319 - Serious Eye Damage/Eye Irritation Category 2, H301 - Acute Toxicity (Oral) Category 3

Legend: 1. Classified by Chemwatch; 2. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI

## 2.2. Label elements

Hazard pictogram(s)	
Signal word	Danger

### Hazard statement(s)

H312	larmful in contact with skin.	
H332	Harmful if inhaled.	
H335	May cause respiratory irritation.	
H315	Causes skin irritation.	
H319	Causes serious eye irritation.	
H301	Toxic if swallowed.	

# Supplementary statement(s)

Not Applicable

### Precautionary statement(s) Prevention

P264	Wash all exposed external body areas thoroughly after handling.	
P270	Do not eat, drink or smoke when using this product.	
P271	Use only outdoors or in a well-ventilated area.	
P261	Avoid breathing mist/vapours/spray.	
P280	P280 Wear protective gloves, protective clothing, eye protection and face protection.	

#### Precautionary statement(s) Response

P301+P310	IF SWALLOWED: Immediately call a POISON CENTER/doctor/physician/first aider.		
P330	Rinse mouth.		
P305+P351+P338	IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.		
P312	Call a POISON CENTER/doctor/physician/first aider/if you feel unwell.		
P337+P313	If eye irritation persists: Get medical advice/attention.		
P302+P352	IF ON SKIN: Wash with plenty of water.		
P304+P340	IF INHALED: Remove person to fresh air and keep comfortable for breathing.		
P332+P313	If skin irritation occurs: Get medical advice/attention.		
P362+P364	Take off contaminated clothing and wash it before reuse.		

### Precautionary statement(s) Storage

P405	Store locked up.	
P403+P233         Store in a well-ventilated place. Keep container tightly closed.		

### Precautionary statement(s) Disposal

Dispose of contents/container to authorised hazardous or special waste collection point in accordance with any local regulation.

### 2.3. Other hazards

Inhalation and/or ingestion may produce health damage\*.

P501

REACH - Art.57-59: The mixture does not contain Substances of Very High Concern (SVHC) at the SDS print date.

### **SECTION 3 Composition / information on ingredients**

1. CAS No 2.EC No 3.Index No 4.REACH No	%[weight]	Name	Classification according to regulation (EC) No 1272/2008 [CLP] and amendments	SCL / M-Factor	Nanoform Particle Characteristics
Not Available	100	tert-Butyl-dimethyl- (tributylstannylmethoxy)silane	Not Applicable	Not Applicable	Not Available

Legend: 1. Classified by Chemwatch; 2. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 3. Classification drawn from C&L; \* EU IOELVs available; [e] Substance identified as having endocrine disrupting properties

#### 3.2.Mixtures

See 'Information on ingredients' in section 3.1

### **SECTION 4 First aid measures**

#### 4.1. Description of first aid measures

Eye Contact	<ul> <li>If this product comes in contact with the eyes:</li> <li>Wash out immediately with fresh running water.</li> <li>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.</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	<ul> <li>If skin or hair contact occurs:</li> <li>Flush skin and hair with running water (and soap if available).</li> <li>Seek medical attention in event of irritation.</li> </ul>
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>If swallowed do NOT induce vomiting.</li> <li>If vomiting occurs, lean patient forward or place on left side (head-down position, if possible) to maintain open airway and prevent aspiration.</li> <li>Observe the patient carefully.</li> <li>Never give liquid to a person showing signs of being sleepy or with reduced awareness; i.e. becoming unconscious.</li> <li>Give water to rinse out mouth, then provide liquid slowly and as much as casualty can comfortably drink.</li> <li>Seek medical advice.</li> </ul>

#### 4.2 Most important symptoms and effects, both acute and delayed

See Section 11

#### 4.3. Indication of any immediate medical attention and special treatment needed

Scanty animal data indicate that BAL may be useful against dialkyl but not trialkyl organotin compounds. D-penicillamine is thought to be inactive. GOSSELIN, SMITH & HODGE: Clinical Toxicology of Commercial Products, 5th Ed

Dimercaprol is suggested to be an effective antidote for dialkyltin poisoning and has been reported to prevent the accumulation of alpha-keto acids produced by dialkyltin compounds. It does not however appear to protect rats from the general toxic effects of triethyltin compounds. This may be due to the fact that dialkyltin compounds, at least up to dihexyl derivatives, react readily with sulfhydryl groups and trialkyltin compounds do not.

Surgical decompression was considered to be the only treatment that offered any benefit in human cases of cerebral oedema caused by trialkyl compounds. Tin and Organotin Compounds: A Preliminary Review.

ENVIRONMENTAL HEALTH CRITERIA: World Health Organization Geneva 1980.

### **SECTION 5 Firefighting measures**

#### 5.1. Extinguishing media

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

#### 5.2. Special hazards arising from the substrate or mixture

Fire Incompatibility None known.

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

#### **SECTION 6 Accidental release measures**

### 6.1. Personal precautions, protective equipment and emergency procedures

See section 8

### 6.2. Environmental precautions

See section 12

# 6.3. Methods and material for containment and cleaning up

Minor Spills	<ul> <li>Clean up all spills immediately.</li> <li>Avoid breathing vapours and contact with skin and eyes.</li> <li>Control personal contact with the substance, by using protective equipment.</li> <li>Contain and absorb spill with sand, earth, inert material or vermiculite.</li> <li>Wipe up.</li> <li>Place in a suitable, labelled container for waste disposal.</li> </ul>
Major Spills	<ul> <li>Clear area of personnel and move upwind.</li> <li>Alert Fire Brigade and tell them location and nature of hazard.</li> <li>Wear breathing apparatus plus protective gloves.</li> <li>Prevent, by any means available, spillage from entering drains or water course.</li> <li>Stop leak if safe to do so.</li> <li>Contain spill with sand, earth or vermiculite.</li> <li>Collect recoverable product into labelled containers for recycling.</li> <li>Neutralise/decontaminate residue (see Section 13 for specific agent).</li> <li>Collect solid residues and seal in labelled drums for disposal.</li> <li>Wash area and prevent runoff into drains.</li> <li>After clean up operations, decontaminate and launder all protective clothing and equipment before storing and re-using.</li> <li>If contamination of drains or waterways occurs, advise emergency services.</li> </ul>

### 6.4. Reference to other sections

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

### **SECTION 7 Handling and storage**

### 7.1. 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>
Fire and explosion protection	See section 5

Other information	<ul> <li>Store in original containers.</li> <li>Keep containers securely sealed.</li> <li>Store in a cool, dry, well-ventilated area.</li> <li>Store away from incompatible materials and foodstuff containers.</li> <li>Protect containers against physical damage and check regularly for leaks.</li> <li>Observe manufacturer's storage and handling recommendations contained within this SDS.</li> </ul>
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# 7.2. Conditions for safe storage, including any incompatibilities

Suitable container	<ul> <li>Lined metal can, lined metal pail/ can.</li> <li>Plastic pail.</li> <li>Polyliner drum.</li> <li>Packing as recommended by manufacturer.</li> <li>Check all containers are clearly labelled and free from leaks.</li> <li>For low viscosity materials</li> <li>Drums and jerricans must be of the non-removable head type.</li> <li>Where a can is to be used as an inner package, the can must have a screwed enclosure.</li> <li>For materials with a viscosity of at least 2680 cSt. (23 deg. C) and solids (between 15 C deg. and 40 deg C.):</li> <li>Removable head packaging;</li> <li>Cans with friction closures and</li> <li>low pressure tubes and cartridges may be used.</li> <li>Where combination packages are used, and the inner packages are of glass, there must be sufficient inert cushioning material in contact with inner and outer packages *.</li> <li>In addition, where inner packagings are glass and contain liquids of packing group I and II there must be sufficient inert absorbent to absorb any spillage *.</li> <li>* unless the outer packaging is a close fitting moulded plastic box and the substances are not incompatible with the plastic.</li> </ul>
Storage incompatibility	<ul> <li>Avoid strong acids, bases.</li> <li>Store at 2-8°C</li> <li>Moisture sensitive</li> <li>Store under argon</li> </ul>
Hazard categories in accordance with Regulation (EC) No 1272/2008	Not Available
Qualifying quantity (tonnes) of dangerous substances as referred to in Article 3(10) for the application of	Not Available

### 7.3. Specific end use(s)

See section 1.2

# **SECTION 8 Exposure controls / personal protection**

### 8.1. Control parameters

Ingredient	DNELs Exposure Pattern Worker	PNECs Compartment
Not Available	Not Available	Not Available

\* Values for General Population

### Occupational Exposure Limits (OEL)

# INGREDIENT DATA

Source	Ingredient	Material name	TWA	STEL	Peak	Notes
Not Available						

Not Applicable

#### Emergency Limits

Ingredient	TEEL-1	TEEL-2	TEEL-3

Ingredient	TEEL-1	TEEL-2		TEEL-3
tert-Butyl-dimethyl- (tributylstannylmethoxy)silane	Not Available	Not Available		Not Available
Ingredient	Original IDLH		Revised IDLH	
tert-Butyl-dimethyl- (tributylstannylmethoxy)silane	Not Available		Not Available	

# 8.2. Exposure controls

8.2.1. Appropriate       Engineering controls are used to renove a hazard or place a barrier between the worker and the hazard. Wild-designed migrowide this high level of protections.       Wild-designed migrowide the high level of protections are will typically be independent of worker introducts to be independent of worker and vertilation that is trabatigically fields and "removed ail in the work environment. Vertilation can remove or diffuse and vertilation that is trabatigically fields and "removed ail in the work environment. Vertilation can remove or diffuse and vertilation that is trabatigically fields and "removed ail in the work environment. Vertilation can remove or diffuse and constantion of the designed properit. The designed migrowide in special encourses and chemical or constantion of the designed properit. The designed migrowide is special encourses and chemical or constantions. Content the is essential to encourse adequate proteints.         Local on-hause ventilation usual (SCBA) may be required in special encourses and chemical or constantions.       Air Special set on the antipate proteints.         Provide adequates proteints.       Air Special Constantions.       Air Special Set on the antipate proteints.         Provide adequates proteints.       Air Special Constantions.       Air Special Set on the antipate proteints.         Provide adequates proteints.       Air Special Set on the antipate proteints.       Air Special Set on the antipate proteints.         Provide adequates proteints.       Air Special Set on the antipate proteints.       Air Special Set on the antipate antints.       Air Special Set on the						
8.2.1. Appropriate engineering controls       solvent, Vapours, degleasing etc., evaporating from tank (in suit air).       (50-100 fmin.)         accosis, furnes from pouring operations, intermittent container filling, low speed conveyer transfers, welding, spray quint, plating acid furnes, picking (released at low velocity into zone of active generation)       0.5-1 m/S (100-200 fmin.)         discharge (active generation into zone of rapid air motion).       interesting active generation into zone of rapid air motion).         ginding, 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 fmin.)         Within each range the appropriate value depends on: Lower end of the range       Upper end of the range       2.5-10 m/S (500-2000 fmin.)         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 fails rapidly with distance away from the opening of a simple extraction pipe. Velocity generally decreases with the square of distance from the extraction point. Other mechanical considerations, producing performance defloits 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.         8.2.2. Individual protection measures, such as parsonal protective equipment       • Safety glasses with side shields. • Contact lenese may poes a special hazard;		engineering controls can be highly effective in protecting wor provide this high level of protection. The basic types of engineering controls are: Process controls which involve changing the way a job activi Enclosure and/or isolation of emission source which keeps a that strategically "adds" and "removes" air in the work enviro designed properly. The design of a ventilation system must n Employers may need to use multiple types of controls to pre- Local exhaust ventilation usually required. If risk of overexpor obtain adequate protection. Supplied-air type respirator may ensure adequate protection. An approved self contained breathing apparatus (SCBA) ma Provide adequate ventilation in warehouse or closed storage "escape" velocities which, in turn, determine the "capture vel contaminant.	rkers and will typically be independent of work ity or process is done to reduce the risk. a selected hazard "physically" away from the work nment. Ventilation can remove or dilute an ai natch the particular process and chemical or went employee overexposure. Issure exists, wear approved respirator. Correct be required in special circumstances. Correct y be required in some situations. a area. Air contaminants generated in the work	xer interactions to worker and ventilation r contaminant if contaminant in use. It fit is essential to t fit is essential to kplace possess varying worker and ventilation		
8.2.1. Appropriate engineering controls       acrossis, furnes from pouring operations, intermittent container filling, low speed conveyer transfers. (100-200 fmin.)         acrossis, furnes from pouring operations, intermittent container filling, low speed conveyer transfers. (100-200 fmin.)       0.5.1 m/s (100-200 fmin.)         direct spray, spray paining in shallow booths, drum filling, conveyer loading, crusher dusts, gas discharge (active generation into zone of rapid air motion).       12.5 m/s (200-500 fmin.)         grinding, abrasive blashing, tumbling, high speed wheel generated dusts (released at high initial velocity into zone of very high rapid air motion).       2.5-10 m/s (200-2000 fmin.)         Within each range the appropriate value depends on: Lower end of the range       Upper end of the range       0.5-11 m/s (200-2000 fmin.)         3: Intermittent, low production.       3: High production, heavy use (200-100 m/s), developed at mass in motion       4: Small hood-local control only         Simple theory shows that air velocity fails rapidly with distance away from the opening of a simple extraction pipe. Velocity generately decreases with the square of distance from the comainingting source. The air velocity at the extraction fain for example, should be a minimum of 1-2 m/s (200-400 fmin) for extraction of solvents generated in a tank 2 meters distant for the extraction paint (fmin 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.         8.2.2.1. Individual protection measures, such as personal protective equipment       Saf		solvent, vapours, degreasing etc., evaporating from tank (i	n still air).			
engineering controls       Use 1 m/26 generation)         engineering controls       useding, spray drift, plating acid fumes, pickling (released at low velocity into zone of active generation)       Use 1 m/26 (102-200 fr/min.)         discharge (active generation into zone of rapid air motion)       iffer spray, spray paining 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 fr/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 fr/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: Sarall hood-local control only       1: metros and the extraction point (nimple cases). Therefore the air speed at the extraction point should be adjusted, accordingly, after reference to distance from the extraction point (nimple cases). Therefore the air speed at the extraction point should be adjusted, accordingly, after reference to distance from the extraction systems are installed or used.         8.2.2. Individual protection measures, such as personal protection equipment       Safety glasses with side shields.       Chemical goggles, IASNZ2 1337.1. EN166 or national equivalent] </td <th></th> <td></td> <td>,</td> <td>(50-100 f/min.)</td>			,	(50-100 f/min.)		
discharge (active generation into zone of rapid air motion)       (200-500 f/min.)         grinding, abrasive blasting, turbling, high speed wheel generated dusts (released at high initial velocity into zone of very high rapid air motion).       (200-500 f/min.)         Within each range the appropriate value depends on:       Lower end of the range       (200-2000 f/min.)         1. Room air currents minimal or favourable to capture       1: Disturbing room air currents       (200-2000 f/min.)         2. Contaminants of low toxicity or of nuisance value only.       2: Contaminants of high toxicity       (3)         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 rapidy with distance away from the opening of a simple extraction pipe. Velocity generally decreases with the square of distance from the extraction of solvents generated in a tank 2 meters distant from the extraction point. Other mechanical considerations, producing performance deficits within the extraction apparatus, make it essential that theoretical air velocities are multiplied by factors of 10 or more when extraction systems are installed or used.         8.2.2. Individual protective equipment       • Safety glasses with side 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 readiv avelable or task. This should include a reviewof lens absorption and adsorption for the clas		welding, spray drift, plating acid fumes, pickling (released a				
S2.2. Individual protection measures, such as personal protective equipment       Safety glasses with side shields.         Eye and face protection       Safety glasses with side shields.         Chemical gogles. [AS/N2S 13.1, EN166 or national equivalent]         Contact lenses may pose a special hazard; soft contact lenses on use and adsorption for the class of chemicals in use and an account of injury experience. Medical and first-aid persone) should be transition for the class of chemicals in use and an account of injury experience. Medical and first-aid persone) should be removed at the first-aid persone) should be created for each workplace on task. This should be created in their endes on the contact lenses on as practicable. Lens should be created endes on as practicable. Lens should be removed at the first-aid persone) should be removed in their removed and account of injury experience. Medical and first-aid persone) should be removed and discrete form the catact on point in the extraction point. Other mechanical considerations, producing performance definits within the extraction apparatus. make it essential that theoretical air velocities are multiplied by factors of 10 or more when extraction systems are initialled or used.         8.2.2. Individual protection measures, such as personal protective.       Safety glasses with side shields.         Chemical gogles. [AS/N2S 1337.1, EN166 or national equivalent]       Contact lenses may pees a special hazard; soft contact lenses may based and concentrate irritants. A writen 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						
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 point should be adjusted, accordingly, after reference to distance from the contaminating source. The air velocity at the extraction point should be adjusted, accordingly, after reference to distance from the extraction apparatus, make it essential that theoretical air velocities are multiplied by factors of 10 or more when extraction apparatus, make it essential that theoretical air velocities are multiplied by factors of 10 or more when extraction systems are installed or used.         8.2.2. Individual protection measures, such as personal protective       • Safety glasses with side shields.         • Contact lenses may pose a special hazard; soft contact lenses may absorb and concentrate iritants. A written policy document, describing the wearing of lenses or restrictions on use, should be created for each workplace or task. This should bis removed and first-aid personnel should be trained in their removal and suitable equipment should						
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 pain thould be adjusted, accordingly, after reference to distance from the contaminating source. The air velocity at the extraction fan, for example, should be a minimum of 1-2 m/s (200-400 fmin) for extraction of solvents generated in a tank 2 meters distant from the extraction point. Other mechanical considerations, producing performance deficits within the extraction apparatus, make it essential that theoretical air velocities are multiplied by factors of 10 or more when extraction systems are installed or used.         8.2.2. Individual protection generated in a generated in a 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.         8.2.2. Individual protection generate in a generate in a tank 2 meters distant from the extraction point. Cher 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		Within each range the appropriate value depends on:				
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 a dijusted, 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 t/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.         8.2.2. Individual protection genupment       • Safety glasses with side shields.         • Chemical goggles, IAS/NZS 1337.1, EN166 or national equivalent]       • Safety glasses with a 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 removal and suitable equipment should be readily available. In the removal and suitable equipment only after workers have washed hands thoroughly. [CDC NIOSH Current Intelligence Bulletin 59].		Lower end of the range	Upper end of the range			
3: Intermittent, low production.       3: High production, heavy use         4: Large hood or large air mass in motion       4: Small hood-local control only         Simple theory shows that air velocity falls rapidly with distance away from the opening of a simple extraction pipe. Velocity generally decreases with the square of distance from the extraction point (in simple cases). Therefore the air speed at the extraction point should be adjusted, accordingly, after reference to distance from the contaminating source. The air velocity at the extraction point for 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.         8.2.2. Individual protection measures, such as personal protective equipment       • Safety glasses with side 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 readity available. In the extraction experience hear special hazard; soft contact lenses as cont as practicable. Lense 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].		1: Room air currents minimal or favourable to capture	1: Disturbing room air currents			
4: Large hood or large air mass in motion       4: Small hood-local control only         Simple theory shows that air velocity falls rapidly with distance away from the opening of a simple extraction pipe. Velocity generally decreases with the square of distance from the extraction point (in simple cases). Therefore the air speed at the extraction point should be adjusted, accordingly, after reference to distance from the contaminating source. The air velocity at the extraction 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.         8.2.2. Individual protection measures, such as personal protective equipment       • Safety glasses with side shields.         • Chemical goggles. [AS/NZS 1337.1, EN166 or national equivalent]       • 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 readily available. In the event of chemical exposure, begin eye irrigation inmediately and ermove contact lens as son as practicable. Lens should be readily and albe removed at the first signs of eye redness or irritation - lens should be readily and leneword in a clean environment only after workers have washed hands thoroughly. [CDC NIOSH Current Intelligence Bulletin 59].		2: Contaminants of low toxicity or of nuisance value only.	2: Contaminants of high toxicity			
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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.         8.2.2. Individual protection measures, such as personal protective equipment       Image: Comparison of the extraction point of the extraction considerations, producing performance deficits within the extraction systems are installed or used.         8.2.2. Individual protection measures, such as personal protective equipment       Image: Comparison of the extraction point of the extraction point of 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.         8.2.2. Individual protection measures, such as personal protective equipment       Image: Comparison of the extraction point of the extraction point of 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.         B.2.2. Individual protection measures, such as personal protective equipment       Image: Comparison of the extraction point of the extraction point of the extraction approximate the extraction approximat		4: Large hood or large air mass in motion	4: Small hood-local control only			
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<ul> <li>Chemical goggles. [AS/NZS 1337.1, EN166 or national equivalent]</li> <li>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].</li> </ul>	measures, such as personal protective					
Skin protection See Hand protection below	Eye and face protection	<ul> <li>Chemical goggles. [AS/NZS 1337.1, EN166 or national e</li> <li>Contact lenses may pose a special hazard; soft contact l document, describing the wearing of lenses or restriction include a review of lens absorption and adsorption for the Medical and first-aid personnel should be trained in their event of chemical exposure, begin eye irrigation immedia be removed at the first signs of eye redness or irritation</li> </ul>	enses may absorb and concentrate irritants. Is on use, should be created for each workplate e class of chemicals in use and an account of removal and suitable equipment should be re- ately and remove contact lens as soon as pra- lens should be removed in a clean environm	ce or task. This should injury experience. adily available. In the cticable. Lens should		
	Skin protection	See Hand protection below				

Hands/feet protection	The selection of suitable gloves does not only depend on the material, but also on further marks of quality which vary from manufacturer. Where the chemical is a preparation of several substances, the resistance of the glove material can not be calculated in advance and has therefore to be checked prior to the application. The exact break through time for substances has to be obtained from the manufacturer of the protective gloves and has to be observed when making a final choice. Personal hygiene is a key element of effective hand care. Gloves must only be worn on clean hands. After using gloves, hands should be washed and dried thoroughly. Application of a non-perfumed moisturiser is recommended. Suitability and durability of glove type is dependent on usage. Important factors in the selection of gloves include:  • frequency and duration of contact,  • glove thickness and  • Some glove polymer types are less affected by movement and this should be taken into account when considering gloves for  iong-term use.  • Contaminated gloves should be replaced.  As defined in ASTM F-739-96 in any application, gloves are rated as:  • Excellent when breakthrough time < 20 min  • Fair when breakthrough time < 20 min  • Fair when breakthrough time < 20 min  • Foor when brea
Body protection	See Other protection below
Other protection	<ul> <li>Overalls.</li> <li>Eyewash unit.</li> <li>Barrier cream.</li> <li>Skin cleansing cream.</li> </ul>

### 8.2.3. Environmental exposure controls

See section 12

# **SECTION 9 Physical and chemical properties**

# 9.1. Information on basic physical and chemical properties

Appearance	Not Available		
Physical state	Liquid	Relative density (Water = 1)	Not Available
Odour	Not Available	Partition coefficient n-octanol / water	Not Available
Odour threshold	Not Available	Auto-ignition temperature (°C)	Not Available
pH (as supplied)	Not Available	Decomposition temperature (°C)	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	Not Available	pH as a solution (1%)	Not Available
Vapour density (Air = 1)	Not Available	VOC g/L	Not Available
Nanoform Solubility	Not Available	Nanoform Particle Characteristics	Not Available
Particle Size	Not Available		

### 9.2. Other information

Not Available

# **SECTION 10 Stability and reactivity**

10.1.Reactivity	See section 7.2
10.2. Chemical stability	<ul> <li>Unstable in the presence of incompatible materials.</li> <li>Product is considered stable.</li> <li>Hazardous polymerisation will not occur.</li> </ul>
10.3. Possibility of hazardous reactions	See section 7.2
10.4. Conditions to avoid	See section 7.2
10.5. Incompatible materials	See section 7.2
10.6. Hazardous decomposition products	See section 5.3

### **SECTION 11 Toxicological information**

# 11.1. Information on hazard classes as defined in Regulation (EC) No 1272/2008 Information on toxicological effects

tert-Butyl-dimethy (tributylstannylmethoxy)silan		IRRITATION Not Available	
Chronic	using animal models); nevertheless exposure by all Tributyltins and dibutyltins affect hormone function a immune system, making one fall ill easily and freque	produce chronic effects adverse to the health (as classified by EC Dir routes should be minimised as a matter of course. Ind reproduction. They also damage the nerves and suppress the boo ntly. They may also cause cancers. Several organotin compounds al are classified as persistent, bioaccumulative and toxic.	dy s
Eye	This material can cause eye irritation and damage in	some persons.	
Skin Contact	using animal models). Nevertheless, good hygiene p be used in an occupational setting. Open cuts, abraded or irritated skin should not be ex Entry into the blood-stream, through, for example, cu	h effects or skin irritation following contact (as classified by EC Direc ractice requires that exposure be kept to a minimum and that suitabl posed to this material its, abrasions or lesions, may produce systemic injury with harmful ef ensure that any external damage is suitably protected.	e glove
Ingestion	Accidental ingestion of the material may be damagin Subchronic exposures to mono-, di- and tri- and tetra nervous, immune and renal systems, the liver and bi	a-substituted organotin compounds may elicit toxic response in the ca	entral
Inhaled	inhalation of vapours, fumes or aerosols, especially distress.	tation (as classified by EC Directives using animal models). Neverthe for prolonged periods, may produce respiratory discomfort and occas herated by the material during the course of normal handling, may be esembles that found by other means of exposure.	sionally

Unless otherwise specified data extracted from RTECS - Register of Toxic Effect of chemical Substances

arcinogenic	icity 🗙
Reproductiv	ivity 🗙
ngle Exposi	sure 🗸
ated Exposi	sure ×
iration Haza	zard 🗙

Legend: X − Data either not available or does not fill the criteria for classification → − Data available to make classification

#### 11.2 Information on other hazards

#### **11.2.1. Endocrine disrupting properties**

No evidence of endocrine disrupting properties were found in the current literature.

#### 11.2.2. Other information

See Section 11.1

#### **SECTION 12 Ecological information**

#### 12.1. Toxicity

tent Dutul dimethod	Endpoint	Test Duration (hr)	Species	Value	Source
tert-Butyl-dimethyl- (tributylstannylmethoxy)silane	Not Available	Not Available	Not Available	Not Available	Not Available
	4. US EPA, Ecot	. IUCLID Toxicity Data 2. Europe ECHA Re ox database - Aquatic Toxicity Data 5. ECE Data 7. METI (Japan) - Bioconcentration D	TOC Aquatic Hazard Assessment Data 6. I		-

Organotin compounds are characterized by a Sn4+ ion with between one to four organic ligands attached. They are classified according to the type of organic ligand, with butyltins, octyltins och phenyltins being the most common. A large number of organtin substances are used in society, with some, such as the butyltins, being recognised as environmental pollutants.

Ecotoxicity increases dramatically in the order methylbutyltin (MBT, RSn) < dibutyltin (DBT, R2Sn) < tributyltin (TBT, R3Sn) for certain endpoints. Organotin degrades by UV irradiation or by biological or chemical cleavage, where the tin-carbon bond is broken. This can occur rapidly in water, however adhesion to suspended particles deposited as sedimentation is the key removal process. Organotin compounds are rapidly degraded by light at surfaces, and hence are not released to the air from various surfaces by any significant amounts. The speciation of organotin compounds is pH-dependent, with the cationic form being the primary form at low pHs, and he neutral hydroxide compounds becoming the predominant form with increased pH. In the environmentally relevant pH range (pH 5–9), the neutral hydroxide compounds (i.e., R3SnOH, R2Sn(OH)2, and Rsn(OH)3) predominate. High concentrations of chloride favor the formation of chloro species. The pKa values for trimethyltin, tributyltin, and triphenyltin cations are approximately 6.60, 6.81, 6.25, and 5.2, respectively. Organotin compounds degrate much slower in sediments than in water, with half-lives estimated to be several years. Chemical and/or biological degradration may occur by dealkylation or methylation, which may result in volatile tin compounds. At ambient temperatures, the solubilities of organotin compounds range from 0.0001 to about 50 mg/L. Organotin compounds may be transferred from water to aquatic organisms, and for tributylin and triphenyltin, the more significant route is through the uptake of water rather than what is being channelled through the food chain. BCF values for all organotin compounds show that organotin compounds tends to accumulate in the livers of organisms, with the tributyltin compounds showing the highest BCFs..

The use of tributyltin (TBT) in ship antifouling paints has caused significant harm to the marine environment worldwide. Female molluscs are masculinized by TBT at levels as low as ca 1 ng/l, preventing them from reproducing. Other organotin substances such as dibutyltin (DBT), dioctyltin and monobutyltin (MBT) are commercially used in other applications. Most industrial organotin chemicals (OTCs) are composed of an organotin cation and one or several ligands, and most of these chemicals are reconverted to the organotin cation compounds in natural waters. The cation may form dissolved complexes with e.g. chloride in seawater. Therefore, their environmental partitioning properties such as Kd and Kh depend in part on the balancing anion in the environment.

Organotins are moderately hydrophobic and associate strongly to particles in natural waters. This increases with increasing number of alkyl groups and increased chain length. In harbour sediments MBT exhibits the weakest affinity to particles, and TBT the strongest (MBT < DBT < TBT). In various soils, however, the reverse pattern was observed. In organic soils, log Kd exceeded 4.0, whereas adsorption was less strong in mineral soils. In contrast to hydrophobic pollutants such as PCBs or PAHs (that partition to lipids in organic matter), OTCs are adhere to the functional groups of organic matter, e.g. phenolic and carboxylic groups. Due to their cationic nature, long-range atmospheric transport has not been considered as important. However, TBT forms highly volatile chloride species in seawater and thus has the potential for long-range atmospheric transportation. As organotins are progressively dealkylated in nature, subsequent dealkylation in the atmosphere, may convert TBT to DBT and MBT. This is important to consider when monitoring data are evaluated, since the occurrence of, e.g., DBT may be due to direct release of DBT or to release of TBT that is subsequently dealkylated. Half-lives in soils and sediments usually range from one to a few years, but may be longer under reducing conditions, whereas half-lives in natural waters may range from a few days to several weeks.

Organotin compounds have been detected in various marine organisms, from invertebrates to mammals. In fish and marine mammals, TBT and TPT bioaccumulate more strongly in liver than in muscle. Bioaccumulation is often stronger in bivalves than in fish, a consequence of lower metabolic capacity in bivalves. Trisubstituted OTCs are more strongly bioaccumulated than the less lipophilic disubstituted OTCs. Because TBT is dealkylated in many organisms, DBT may be present in organisms but may not necessarily be the organotin substance that was assimilated. Most studies do not suggest that TBT is biomagnified in aquatic food-chains. However, TPT appears to be biomagnified fairly strongly in the aquatic food chain. The trisubstituted substances, TPT and in particular TBT, are widely considered the most toxic organotin substances. TBT has been demonstrated to cause masculinization (imposex) of female marine molluscs, as well as

fish, which consequently compromises their ability to reproduce. While DBT and MBT do not cause imposex, both TBT and DBT have negative effects on the reproductive system of mammals. In line with these facts, TBT and TPT were given the highest category in a European review of endocrine disrupting chemicals: "Evidence for endocrine disruption in living organisms". TBT was also classified as "Evidence of potential to cause endocrine disruption in humans". **DO NOT** discharge into sewer or waterways.

#### 12.2. Persistence and degradability

Ingredient	Persistence: Water/Soil	Persistence: Air
	No Data available for all ingredients	No Data available for all ingredients

#### 12.3. Bioaccumulative potential

Ingredient	Bioaccumulation
	No Data available for all ingredients

### 12.4. Mobility in soil

Ingredient	Mobility
	No Data available for all ingredients

### 12.5. Results of PBT and vPvB assessment

	Р	В	т	
Relevant available data	Not Available	Not Available	Not Ava	ailable
PBT	×	×	×	
vPvB	×	×	×	
				I
PBT Criteria fulfilled?				No
vPvB				No

#### 12.6. Endocrine disrupting properties

No evidence of endocrine disrupting properties were found in the current literature.

#### 12.7. Other adverse effects

No evidence of ozone depleting properties were found in the current literature.

### **SECTION 13 Disposal considerations**

### 13.1. Waste treatment methods

Product / Packaging disposal	Legislation addressing waste disposal requirements may differ by country, state and/ or territory. Each user must refer to laws operating in their area. In some areas, certain wastes must be tracked. A Hierarchy of Controls seems to be common - the user should investigate: • Reduction • Reuse • Recycling • Disposal (if all else fails) This material may be recycled if unused, or if it has not been contaminated so as to make it unsuitable for its intended use. If it has been contaminated, it may be possible to reclaim the product by filtration, distillation or some other means. Shelf life considerations should also be applied in making decisions of this type. Note that properties of a material may change in use, and recycling or reuse may not always be appropriate. • DO NOT allow wash water from cleaning or process equipment to enter drains. • It may be necessary to collect all wash water for treatment before disposal. • In all cases disposal to sewer may be subject to local laws and regulations and these should be considered first. • Where in doubt contact the responsible authority.
Waste treatment options	Not Available
Sewage disposal options	Not Available

### **SECTION 14 Transport information**

	6
Marine Pollutant	NO
HAZCHEM	2X

# Land transport (ADR-RID)

14.1. UN number or ID number	2788		
14.2. UN proper shipping name	ORGANOTIN COM	MPOUND, LIQU	JID, N.O.S.
14.3. Transport hazard	Class	6.1	
class(es)	Subsidiary risk	Not Applicab	le
14.4. Packing group	III		
14.5. Environmental hazard	Not Applicable		
	Hazard identifica	tion (Kemler)	60
	Classification co	de	Т3
14.6. Special precautions	Hazard Label		6.1
for user	Special provisions		43 274
	Limited quantity		5 L
	Tunnel Restrictio	on Code	2 (E)

# Air transport (ICAO-IATA / DGR)

14.1. UN number	2788		2788			
14.2. UN proper shipping name	Organotin compound, liquid, n.o.s. *					
	ICAO/IATA Class	6.1				
14.3. Transport hazard class(es)	ICAO / IATA Subrisk	Not Applicable				
01033(03)	ERG Code	6L				
14.4. Packing group	III	III				
14.5. Environmental hazard	Not Applicable					
	Special provisions		A3 A4 A6			
	Cargo Only Packing Ir	nstructions	663			
	Cargo Only Maximum	Qty / Pack	220 L			
14.6. Special precautions for user	Passenger and Cargo	Packing Instructions	655			
	Passenger and Cargo	Maximum Qty / Pack	60 L			
	Passenger and Cargo Limited Quantity Packing Instructions		Y642			
	Passenger and Cargo Limited Maximum Qty / Pack		2 L			

# Sea transport (IMDG-Code / GGVSee)

14.1. UN number	2788	
14.2. UN proper shipping name	ORGANOTIN COMPOUND, LIQUID, N.O.S.	
14.3. Transport hazard class(es)	IMDG Class 6.1	
	IMDG Subrisk Not Applicable	
14.4. Packing group	Ш	

14.5. Environmental hazard	Not Applicable		
14.6. Special precautions for user	EMS Number	F-A, S-A	
	Special provisions	43 223 274	
	Limited Quantities	5 L	

#### Inland waterways transport (ADN)

14.1. UN number	2788	
14.2. UN proper shipping name	ORGANOTIN COMPOL	JND, LIQUID, N.O.S.
14.3. Transport hazard class(es)	6.1 Not Applicable	
14.4. Packing group	111	
14.5. Environmental hazard	Not Applicable	
14.6. Special precautions for user	Classification code	Т3
	Special provisions	43; 274; 802
	Limited quantity	5 L
	Equipment required	PP, EP, TOX, A
	Fire cones number	0

#### 14.7. Maritime transport in bulk according to IMO instruments

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

Not Applicable

#### 14.7.2. Transport in bulk in accordance with MARPOL Annex V and the IMSBC Code

Product name	Group
14.7.3. Transport in bulk i	n accordance with the IGC Code

Product name Ship Type
------------------------

### **SECTION 15 Regulatory information**

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

This safety data sheet is in compliance with the following EU legislation and its adaptations - as far as applicable - : Directives 98/24/EC, - 92/85/EEC, - 94/33/EC, - 2008/98/EC, - 2010/75/EU; Commission Regulation (EU) 2020/878; Regulation (EC) No 1272/2008 as updated through ATPs.

### Information according to 2012/18/EU (Seveso III):

Seveso Category	Not Available

### 15.2. Chemical safety assessment

For further information please look at the Chemical Safety Assessment and Exposure Scenarios prepared by your Supply Chain if available.

#### ECHA SUMMARY

Not Applicable

### **National Inventory Status**

National Inventory	Status
Australia - AIIC / Australia Non-Industrial Use	Not Available
Canada - DSL	Not Available
Canada - NDSL	Not Available
China - IECSC	Not Available

National Inventory	Status
Europe - EINEC / ELINCS / NLP	Not Available
Japan - ENCS	Not Available
Korea - KECI	Not Available
New Zealand - NZIoC	Not Available
Philippines - PICCS	Not Available
USA - TSCA	Not Available
Taiwan - TCSI	Not Available
Mexico - INSQ	Not Available
Vietnam - NCI	Not Available
Russia - FBEPH	Not Available
Legend:	Yes = All CAS declared ingredients are on the inventory No = One or more of the CAS listed ingredients are not on the inventory. These ingredients may be exempt or will require registration.

#### **SECTION 16 Other information**

Revision Date	16/05/2022
Initial Date	16/05/2022

### Full text Risk and Hazard codes

#### Other information

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.

For detailed advice on Personal Protective Equipment, refer to the following EU CEN Standards:

EN 166 Personal eye-protection

EN 340 Protective clothing

EN 374 Protective gloves against chemicals and micro-organisms

EN 13832 Footwear protecting against chemicals

EN 133 Respiratory protective devices

### **Definitions and abbreviations**

PC - TWA: Permissible Concentration-Time Weighted Average PC - STEL: Permissible Concentration-Short Term Exposure Limit IARC: International Agency for Research on Cancer ACGIH: American Conference of Governmental Industrial Hygienists STEL: Short Term Exposure Limit TEEL: Temporary Emergency Exposure Limit. IDLH: Immediately Dangerous to Life or Health Concentrations ES: Exposure Standard 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** AIIC: Australian Inventory of Industrial Chemicals **DSL: Domestic Substances List** NDSL: Non-Domestic Substances List IECSC: Inventory of Existing Chemical Substance in China EINECS: European INventory of Existing Commercial chemical Substances ELINCS: European List of Notified Chemical Substances NLP: No-Longer Polymers ENCS: Existing and New Chemical Substances Inventory

# othoxy/silano

tert-Butyl-dimethyl-(tributylstannylmethoxy)silane

KECI: Korea Existing Chemicals Inventory NZIoC: New Zealand Inventory of Chemicals PICCS: Philippine Inventory of Chemicals and Chemical Substances TSCA: Toxic Substances Control Act TCSI: Taiwan Chemical Substance Inventory INSQ: Inventario Nacional de Sustancias Químicas NCI: National Chemical Inventory FBEPH: Russian Register of Potentially Hazardous Chemical and Biological Substances

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