

# **Apollo Scientific**

Part Number: **OR72396** Version No: **2.2** Safety Data Sheet (Conforms to Annex II of REACH (1907/2006) - Regulation 2020/878)

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

### 1.1. Product Identifier

Product name	9-Borabicyclo[3.3.1]nonane, 0.5M solution in THF			
Synonyms	Not Available			
Proper shipping name	ORGANOMETALLIC SUBSTANCE, LIQUID, WATER-REACTIVE, FLAMMABLE			
Other means of identification Not Available				

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

Relevant identified uses	Epoxides (oxiranes) have long been recognised as valuable building blocks which allow derivatization either by forming C-X bonds (through reactions with alcohols, ammonia, amines, phenolates, etc.) or by forming new C-C bonds (through reactions with cyanide, malonates, allyl silyl reagents, or metal-organic reagents). Cyclic ethers have ring structure where the oxygen has become part of the ring. The term of epoxide indicate three membered cyclic ether (also called oxirane); four membered cyclic ether is called oxetane; five membered cyclic ether, furan (or oxolane); six membered cyclic ether, pyran (also called oxane) respectively. Their unhindered oxygen atom carries two unshared pairs of electrons - a structure which favours the formation of coordination complexes and the solvation of cations. Cyclic ethers are used as important solvents, as chemical intermediate and as monomer for ring-opening polymerization.
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	
Registered company name	
Address	Whitefield Road, Bredbury SK62QR United Kingdom
Telephone	01614060505
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>	H225 - Flammable Liquids Category 2, H261 - Substances and Mixtures which in Contact with Water Emit Flammable Gases Category 2, H302 - Acute Toxicity (Oral) Category 4, H314 - Skin Corrosion/Irritation Category 1B, H318 - Serious Eye Damage/Eye Irritation Category 1, H335 - Specific Target Organ Toxicity - Single Exposure (Respiratory Tract Irritation) Category 3, H336 - Specific Target Organ Toxicity - Single Exposure (Narcotic Effects) Category 3, H351 - Carcinogenicity Category 2
Legend:	1. Classified by Chernwatch; 2. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI

Chemwatch Hazard Alert Code: 4

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S.REACH.GB-NIR.EN

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# 9-Borabicyclo[3.3.1]nonane, 0.5M solution in THF



Signal word Danger

### Hazard statement(s)

H225	Highly flammable liquid and vapour.
H261	In contact with water releases flammable gases.
H302	Harmful if swallowed.
H314	Causes severe skin burns and eye damage.
H335	May cause respiratory irritation.
H336	May cause drowsiness or dizziness.
H351	Suspected of causing cancer.

### Supplementary statement(s)

Not Applicable

## Precautionary statement(s) Prevention

P201	Obtain special instructions before use.
P210	Keep away from heat, hot surfaces, sparks, open flames and other ignition sources. No smoking.
P231+P232	Handle and store contents under inert gas. Protect from moisture.
P260	Do not breathe mist/vapours/spray.
P264	Wash all exposed external body areas thoroughly after handling.
P271	Use only outdoors or in a well-ventilated area.
P280	Wear protective gloves, protective clothing, eye protection and face protection.
P240	Ground and bond container and receiving equipment.
P241	Use explosion-proof electrical/ventilating/lighting/intrinsically safe equipment.
P242	Use non-sparking tools.
P243	Take action to prevent static discharges.
P270	Do not eat, drink or smoke when using this product.
P223	Do not allow contact with water.

## Precautionary statement(s) Response

P301+P330+P331	IF SWALLOWED: Rinse mouth. Do NOT induce vomiting.			
P302+P335+P334	P302+P335+P334 IF ON SKIN: Brush off loose particles from skin. Immerse in cool water [or wrap in wet bandages].			
P303+P361+P353	IF ON SKIN (or hair): Take off immediately all contaminated clothing. Rinse skin with water [or shower].			
P305+P351+P338	IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.			
P308+P313	IF exposed or concerned: Get medical advice/ attention.			
P310	Immediately call a POISON CENTER/doctor/physician/first aider.			
P370+P378	In case of fire: Use alcohol resistant foam or normal protein foam to extinguish.			
P363	Wash contaminated clothing before reuse.			
P301+P312	IF SWALLOWED: Call a POISON CENTER/doctor/physician/first aider if you feel unwell.			
P304+P340	IF INHALED: Remove person to fresh air and keep comfortable for breathing.			

## Precautionary statement(s) Storage

P403+P235         Store in a well-ventilated place. Keep cool.	
P405	Store locked up.
P402+P404	Store in a dry place. Store in a closed container.

### Precautionary statement(s) Disposal

P501 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 skin contact may produce health damage\*.

Cumulative effects may result following exposure\*.

### Limited evidence of a carcinogenic effect\*.

May be harmful to the foetus/ embryo\*.

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

### 3.1.Substances

See 'Composition on ingredients' in Section 3.2

### 3.2.Mixtures

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
1. Not Available 2.Not Available 3.Not Available 4.Not Available	8	9-Borabicyclo[3.3.1]nonane	Pyrophoric Solids Category 1, Substances and Mixtures which in Contact with Water Emit Flammable Gases Category 1, Skin Corrosion/Irritation Category 1B, Serious Eye Damage/Eye Irritation Category 1; H250, H260, H314, H318 <sup>[1]</sup>	Not Available	Not Available
1. 109-99-9 2.203-726-8 3.603-025-00-0 4.Not Available	92	tetrahydrofuran *	Flammable Liquids Category 2, Serious Eye Damage/Eye Irritation Category 2, Carcinogenicity Category 2, Specific Target Organ Toxicity - Single Exposure (Respiratory Tract Irritation) Category 3; H225, H319, H351, H335 <sup>[2]</sup>	STOT SE 3; H335: C ≥ 25 %   Eye Irrit.2; H319: C ≥ 25 %	Not Available
Legend: 1. Classified by Chernwatch; 2. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 3. Classification drawn from C&L * I IOELVs available; [e] Substance identified as having endocrine disrupting properties		n from C&L * EU			

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

4.1. Description of first aid me	asures
Eye Contact	<ul> <li>If this product comes in contact with the eyes:</li> <li>Immediately hold eyelids apart and flush the eye continuously with 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>Continue flushing until advised to stop by the Poisons Information Centre or a doctor, or for at least 15 minutes.</li> <li>Transport to hospital or doctor without delay.</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>Immediately flush body and clothes with large amounts of water, using safety shower if available.</li> <li>Quickly remove all contaminated clothing, including footwear.</li> <li>Wash skin and hair with running water. Continue flushing with water until advised to stop by the Poisons Information Centre.</li> <li>Transport to hospital, or doctor.</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, without delay.</li> </ul>
Ingestion	<ul> <li>For advice, contact a Poisons Information Centre or a doctor at once.</li> <li>Urgent hospital treatment is likely to be needed.</li> <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>Transport to hospital or doctor without delay.</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

Keep drv

Treat symptomatically.

## **SECTION 5 Firefighting measures**

### 5.1. Extinguishing media

### DO NOT USE WATER, CO2 OR FOAM ON SUBSTANCE ITSELF

For SMALL FIRES:

- Dry chemical, soda ash or lime.
- For LARGE FIRES:
- DRY sand, dry chemical, soda ash;
- OR withdraw and allow fire to burn itself out.

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

- Avoid contamination with oxidising agents i.e. nitrates, oxidising acids, chlorine bleaches, pool chlorine etc. as ignition may result
- Fire Incompatibility
- NOTE: May develop pressure in containers; open carefully. Vent periodically.

## 5.3. Advice for firefighters

	Alert Fire Brigade and tell them location and nature of hazard.
	May be violently or explosively reactive.
	Wear full protective clothing plus breathing apparatus.
	Prevent, by any means available, spillage from entering drains or water course.
	<ul> <li>Consider evacuation (or protect in place)</li> </ul>
	DO NOT use water on fires.
	CAUTION: If only water available, use flooding quantities of water or withdraw personnel.
Fire Fighting	DO NOT allow water to enter containers.
File Fighting	DO NOT approach containers suspected to be hot.
	Cool fire exposed containers with flooding quantities of water from a protected location until well after fire is out.
	If safe to do so, remove undamaged containers from path of fire.
	If fire gets out of control withdraw personnel and warn against entry.
	Equipment should be thoroughly decontaminated after use.
	Fight fire from a protected position or use unmanned hose holders or monitor nozzles.
	Withdraw immediately in case of rising sound from venting safety devices or discolouration of tanks.
	ALWAYS stay away from tank ends.
	May ignite on contact with air, moist air or water.
	May react vigorously or explosively on contact with water.
	May decompose explosively when heated or involved in fire.
	May REIGNITE after fire is extinguished.
	Gases generated after contact with water or moist air may be poisonous, corrosive or irritating.
	<ul> <li>Gases generated in fire may be poisonous, corrosive or irritating.</li> </ul>
Fire/Explanian Harard	Containers may explode on heating.
Fire/Explosion Hazard	Runoff may create multiple fire or explosion hazard.
	Combustion products include:
	carbon monoxide (CO)
	carbon dioxide (CO2)
	other pyrolysis products typical of burning organic material.
	WARNING: Long standing in contact with air and light may result in the formation
	of potentially explosive peroxides.

## **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>Material from spill may be contaminated with water resulting in generation of gas which subsequently may pressure closed containers.</li> <li>Hold spill material in vented containers only and plan for prompt disposal</li> <li>Eliminate all ignition sources.</li> <li>Cover with DRY earth, sand or other non-combustible material.</li> <li>Then cover with plastic sheet to minimise spreading and to prevent exposure to rain or other sources of water.</li> <li>Use clean, non-sparking tools to collect absorbed material and place into loosely-covered metal or plastic containers ready for disposal.</li> <li>Wear gloves and safety glasses as appropriate.</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 full protective clothing and breathing apparatus.</li> <li>Prevent, by any means available, spillage from entering drains or water courses.</li> <li>No smoking, naked lights or ignition sources.</li> <li>Increase ventilation.</li> <li>Stop leak if safe to do so.</li> <li>DO NOT USE WATER OR NEUTRALISING AGENTS INDISCRIMINATELY ON LARGE SPILLS.</li> <li>Absorb or cover spill with sand, earth, inert material or vermiculite and cover with white mineral oil.</li> <li>Collect recoverable product into labelled containers for recycling.</li> <li>Collect residues and seal in labelled drums for disposal.</li> <li>Wash spill area with detergent and water.</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 as a result of the above actions, 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 hand	ling
Safe handling	<ul> <li>The substance accumulates peroxides which may become hazardous only if it evaporates or is distilled or otherwise treated to concentrate the peroxides. The substance may concentrate around the container opening for example.</li> <li>Purchases of peroxidisable chemicals should be restricted to ensure that the chemical is used completely before it can become peroxidised.</li> <li>A responsible person should maintain an inventory of peroxidisable chemicals or annotate the general chemical inventory to indicate which chemicals are subject to peroxidation. An expiration date should be determined. The chemical should either be treated to remove peroxides or disposed of before this date.</li> <li>The person or laboratory receiving the chemical should record a receipt date on the bottle. The individual opening the container should add an opening date.</li> </ul>

• Unopened containers received from the supplier should be safe to store for 18 months.• Opened containers should not be stored for more than 12 months.• Avoid all personal contact, including inhalation.• Wear protective clothing when risk of overexposure occurs.• Use in a well-ventilated area.• Avoid contact with moisture.• Avoid contact with incompatible materials.• Avoid contact with incompatible materials.• When handling, DO NOT eat, drink or smoke.• Keep containers securely sealed when not in use.• Avoid physical damage to containers.• Avoid physical damage to container should be re-use• Use good occupational work practice.• Observe manufacturer's storage and handling recommendations contained within this SDS.• Containers, even those that have been emptied, may contain explosive vapours.• Do NOT cut, drill, grind, weld or perform similar operations on or near containers.• Do NOT cut, drill, grind, weld or perform similar operations on or near containers.• Do NOT allow clothing wet with material to stay in contact with skinre and explosion protectionKEEP DRY1 Packages must be protected from water ingress.FOR MINOR QUANTTIES:• Store in an indoor fireproof cabinet or in a room of noncombustible construction and	
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FOR MINOR QUANTITIES:	
Store in an indoor fireproof cabinet or in a room of noncombustible construction and	
provide adequate portable fire-extinguishers in or near the storage area.	
FOR PACKAGE STORAGE:	
Store in original containers in approved flame-proof area.	
No smoking, naked lights, heat or ignition sources.	
DO NOT store in pits, depressions, basements or areas where vapours may be trapped.	
Keep containers securely sealed.	
Other information        Store away from incompatible materials in a cool, dry well ventilated area.	
Protect containers against physical damage and check regularly for leaks.	
<ul> <li>Protect containers from exposure to weather and from organize to incert signal to incert.</li> <li>Protect containers from exposure to weather and from direct sumfaith unless: (a) the packages are of metal or plastic construction; (b) the packages are of metal or plastic construction;</li> </ul>	ie
packages are securely closed are not opened for any purpose while in the area where they are stored; (c) adequate precautions are ta	
ensure that rain water, which might become contaminated by the dangerous goods, is collected and disposed of safely.	CITIO
<ul> <li>Ensure proper stock-control measures are maintained to prevent prolonged storage of dangerous goods.</li> </ul>	
Automatic fire-sprinklers <b>MUST NOT</b> be installed in room or space.	
The room or space must be located at least five metres from the boundaries of the premises and from other buildings unless separated	pv a
wall with a fire resistance of at least four hours.	
Observe manufacturer's storage and handling recommendations contained within this SDS.	-, -

## 7.2. Conditions for safe storage, including any incompatibilities

T.2. Conditions for sale storag	e, including any incompatibilities
Suitable container	For low viscosity materials and solids: Drums and jerricans must be of the non-removable head type. Where a can is to be used as an inner package, the can must have a screwed enclosure. For materials with a viscosity of at least 2680 cSt. (23 deg. C):
Storage incompatibility	<ul> <li>For tetrahydrofuran (THF)</li> <li>Avoid contact with oxygen, air, light and heat</li> <li>Contact with illhium aluminium hydride or with sodium or potassium hydroxide can be hazardous when peroxides are present. THF may polymerise in the presence of cationic initiators such as Lewis acids or strong proton acids.</li> <li>Segregate from lithium aluminium hydride, sodium or potassium hydroxide, cationic initiators such as Lewis acids or strong proton acids.</li> <li>In the absence of inhibitors tetrahydrofuran is subject to auto-oxidation with the formation of 2-tetrahydrofuryl hydroperoxide. When heated this tends to decompose smoothly but if allowed to accumulate over a considerable period it transforms to other peroxide species, such as unstable and explosive polyalkylidene peroxide, which violently decompose.</li> <li>Copper(I) chloride has been recommended to remove trace amounts of peroxide. An attempt to remove peroxides by shaking with solid ferrous sulfate, prior to distillation, did not prevent explosion of the distillation residue. Alkali treatment does not appear to be safe.</li> <li>Peroxides may be destroyed by passage through activated carbon at 20-66 C with contact time in excess of 2 min.</li> <li>Is incompatible with borane, calcium mydride, lithium tetrahydroaluminate, sodium aluminium tetrahydride</li> <li>reads violently with stong oxidisers, bromine, oxygen, magnesium tetrahydroaluminate, metal halides, peroxyacetic acid, potassium hydride is attacks some plastics and costings</li> <li>may accumulate static charges that can result in ignition of its vapours</li> </ul> Storage tanks and other equipment should be absolutely dry and free from air, ammonia, acetylene, hydrogen sulfide, rust and other contaminants. <ul> <li>Avoid strong acids, bases.</li> <li>The unhindred oxygen atom found on cyclic ethers such as the epoxide soxteanes, furans, dioxanes and pyrans, carries two unshared pairs of electrons - a structure which favors the formation of coordination complexes and the</li></ul>

	containing halogens (fluorine, chlorine, bromine, iodine) bonded to the metal typically will generate gaseous hydrohalic acids (HF, HCI, HBr, HI) with water.
Hazard categories in accordance with Regulation (EC) No 1272/2008	P5a: Flammable Liquids, P5b: Flammable Liquids, P5c: Flammable Liquids
Qualifying quantity (tonnes) of dangerous substances as referred to in Article 3(10) for the application of	P5a Lower- / Upper-tier requirements: 10 / 50 P5b Lower- / Upper-tier requirements: 50 / 200 P5c Lower- / Upper-tier requirements: 5 000 / 50 000
. ,	P5c Lower- / Upper-tier requirements: 5 000 / 50 000

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
tetrahydrofuran	Dermal 12.6 mg/kg bw/day (Systemic, Chronic) Inhalation 72.4 mg/m <sup>3</sup> (Systemic, Chronic) Inhalation 150 mg/m <sup>3</sup> (Local, Chronic) Inhalation 96 mg/m <sup>3</sup> (Local, Acute) Inhalation 300 mg/m <sup>3</sup> (Local, Acute) Dermal 1.5 mg/kg bw/day (Systemic, Chronic) * Inhalation 13 mg/m <sup>3</sup> (Systemic, Chronic) * Oral 1.5 mg/kg bw/day (Systemic, Chronic) * Inhalation 75 mg/m <sup>3</sup> (Local, Chronic) * Inhalation 52 mg/m <sup>3</sup> (Systemic, Acute) * Inhalation 150 mg/m <sup>3</sup> (Local, Acute) *	<ul> <li>4.32 mg/L (Water (Fresh))</li> <li>0.432 mg/L (Water - Intermittent release)</li> <li>21.6 mg/L (Water (Marine))</li> <li>23.3 mg/kg sediment dw (Sediment (Fresh Water))</li> <li>2.33 mg/kg sediment dw (Sediment (Marine))</li> <li>2.13 mg/kg soil dw (Soil)</li> <li>4.6 mg/L (STP)</li> <li>67 mg/kg food (Oral)</li> </ul>

\* Values for General Population

# Occupational Exposure Limits (OEL)

INGREDIENT DATA								
Source	Ingredient	Material name	TWA		STEL		Peak	Notes
EU Consolidated List of Indicative Occupational Exposure Limit Values (IOELVs)	tetrahydrofuran	Tetrahydrofuran	50 ppm / 150 mg/m3		300 mg/m3 / 100	) ppm	Not Available	Skin
UK Workplace Exposure Limits (WELs)	tetrahydrofuran	rofuran Tetrahydrofuran 50 ppm / 150 mg/m3			300 mg/m3 / 100 ppm		Not Available	Sk
Emergency Limits								
Ingredient	TEEL-1 TEEL-2		TEEL-3					
tetrahydrofuran	Not Available	Available Not Available		Not Available				
Ingredient	Original IDLH Revised IDLH							
9-Borabicyclo[3.3.1]nonane	Not Available			Not /	Available			
tetrahydrofuran	2,000 ppm			Not /	Available			
Occupational Exposure Banding	I							
Ingredient	Occupational Expos	Occupational Exposure Band Rating			Occupational Exposure Band Limit			
9-Borabicyclo[3.3.1]nonane	С			> 1 to ≤ 10 parts per million (ppm)				
Notes:	Occupational exposure banding is a process of assigning chemicals into specific categories or bands based on a chemical's potency and the adverse health outcomes associated with exposure. The output of this process is an occupational exposure band (OEB), which corresponds to a range of exposure concentrations that are expected to protect worker health.							

# 8.2. Exposure controls

8.2.1. Appropriate engineering controls	Engineering controls are used to remove a hazard or place a barrier between the worker and the hazard. Wele be highly effective in protecting workers and will typically be independent of worker interactions to provide this The basic types of engineering controls are: Process controls which involve changing the way a job activity or process is done to reduce the risk. Enclosure and/or isolation of emission source which keeps a selected hazard "physically" away from the work "adds" and "removes" air in the work environment. Ventilation can remove or dilute an air contaminant if design ventilation system must match the particular process and chemical or contaminant in use. Employers may need to use multiple types of controls to prevent employee overexposure. Local exhaust ventilation usually required. If risk of overexposure exists, wear approved respirator. Correct fit protection. Supplied-air type respirator may be required in special circumstances. Correct fit is essential to en An approved self contained breathing apparatus (SCBA) may be required in some situations. Provide adequate ventilation in warehouse or closed storage area. Air contaminants generated in the workplavelocities which, in turn, determine the "capture velocities" of fresh circulating air required to effectively removed Type of Contaminant: solvent, vapours, degreasing etc., evaporating from tank (in still air).	s high level of protection. ker and ventilation that strategically gned properly. The design of a is essential to obtain adequate isure adequate protection. ace possess varying "escape"
--	---	--

		ransfers, welding, spray	0.5-1 m/s (100-200 f/min.)
	1-2.5 m/s (200-500 f/min.)		
grinding, abrasive blasting, tumbling, high speed wheel ge very high rapid air motion).	2.5-10 m/s (500-2000 f/min.)		
Within each range the appropriate value depends on:			
Lower end of the range	Upper end of the range		
1: Room air currents minimal or favourable to capture	1: Disturbing room air currents		
2: Contaminants of low toxicity or of nuisance value only.	2: Contaminants of high toxicity		
3: Intermittent, low production.	3: High production, heavy use		
4: Large hood or large air mass in motion	4: Small hood-local control only		
with the square of distance from the extraction point (in sim accordingly, after reference to distance from the contaminal 1-2 m/s (200-400 f/min) for extraction of solvents generated producing performance deficits within the extraction appara more when extraction systems are installed or used. Engineering controls are used to remove a hazard or place be highly effective in protecting workers and will typically be The basic types of engineering controls are: Process controls which involve changing the way a job actin Enclosure and/or isolation of emission source which keeps "adds" and "removes" air in the work environment. Ventilatii ventilation system must match the particular process and cl Employers may need to use multiple types of controls to pre Engineering controls are used to remove a hazard or place be highly effective in protecting workers and will typically be The basic types of engineering controls are: Process controls which involve changing the way a job The basic types of engineering controls are: Process controls which involve changing the way a job Enclosure and/or isolation of emission source which kee Ventilation that strategically "adds" and "removes" air in designed properly. The design of a ventilation system not	ple cases). Therefore the air spee ting source. The air velocity at the d in a tank 2 meters distant from th tus, make it essential that theoreti a barrier between the worker and a independent of worker interaction vity or process is done to reduce th a selected hazard "physically" away on can remove or dilute an air con hemical or contaminant in use. event employee overexposure. a barrier between the worker and a independent of worker interaction activity or process is done to reduce the selected hazard "physically" n the work environment. Ventilation must match the particular process	d at the extraction point sho extraction fan, for example, e extraction point. Other me cal air velocities are multipli the hazard. Well-designed hs to provide this high level he risk. ay from the worker and vent taminant if designed proper the hazard. Well-designed hs to provide this high level the hazard. Well-designed hs to provide this high level the risk. " away from the worker and h can remove or dilute an ai	uld be adjusted, should be a minimum of schanical considerations, ed by factors of 10 or engineering controls can of protection. illation that strategically ly. The design of a engineering controls can of protection
<ul> <li>Full face shield may be required for supplementary but</li> <li>Contact lenses may pose a special hazard; soft contact the wearing of lenses or restrictions on use, should be and adsorption for the class of chemicals in use and art their removal and suitable equipment should be readily remove contact lens as soon as practicable. Lens should be and adsorption for the source of the source o</li></ul>	never for primary protection of eye t lenses may absorb and concentr created for each workplace or task account of injury experience. Mer available. In the event of chemica ild be removed at the first signs of	ate irritants. A written policy x. This should include a revi dical and first-aid personnel al exposure, begin eye irriga eye redness or irritation - le	ew of lens absorption should be trained in tion immediately and ns should be removed in
See Hand protection below			
The selection of suitable gloves does not only depend on the manufacturer. Where the chemical is a preparation of sever and has therefore to be checked prior to the application. The exact break through time for substances has to be obta making a final choice. Personal hygiene is a key element of effective hand care. G washed and dried thoroughly. Application of a non-perfume Suitability and durability of glove type is dependent on usag frequency and duration of contact, chemical resistance of glove material, glove thickness and dexterity Select gloves tested to a relevant standard (e.g. Europe EN When prolonged or frequently repeated contact may occuminutes according to EN 374, AS/NZS 2161.10.1 or national	alls outside of boots, to avoid spills ne material, but also on further main ral substances, the resistance of the ained from the manufacturer of the Sloves must only be worn on clean d moisturiser is recommended. ge. Important factors in the selection 4 374, US F739, AS/NZS 2161.1 of r, a glove with a protection class of al equivalent) is recommended.	rks of quality which vary from ne glove material can not be protective gloves and has t hands. After using gloves, on of gloves include: or national equivalent). f 5 or higher (breakthrough	e calculated in advance to be observed when hands should be
	<ul> <li>drift, plating acid fumes, pickling (released at low velocity, direct spray, spray painting in shallow booths, drum filling, generation into zone of rapid air motion)</li> <li>grinding, abrasive blasting, tumbling, high speed wheel ge very high rapid air motion).</li> <li>Within each range the appropriate value depends on: <ul> <li>Lower end of the range</li> <li>1: Room air currents minimal of favourable to capture</li> <li>2: Contaminants of low toxicity or of nuisance value only.</li> <li>3: Intermittent, low production.</li> <li>4: Large hood or large air mass in motion</li> </ul> </li> <li>Simple theory shows that air velocity falls rapidly with distativith the square of distance from the extraction paint (in sim accordingly, after reference to distance from the contamina 1-2 m/s (200-400 f/min) for extraction of solvents generated producing performance deficits within the extraction appart more when extraction systems are installed or used.</li> <li>Engineering controls are used to remove a hazard or place be highly effective in protecting workers and will typically be The basic types of engineering controls are:</li> <li>Process controls which involve changing the way a job acti Enclosure and/or isolation of emission source which keeps adds" and "removes" air in the work environment. Ventilativentilation system must match the particular process and c Employers may need to use multiple types of controls to protecting workers and will typically be The basic types of engineering controls are:</li> <li>Process controls which involve changing the way a job acti Enclosure and/or isolation of emission source which keeps and they required for supplementary but</li> <li>Contact lenses may need to use multiple types of controls to protecting workers and will typically be The basic types of engineering controls are:</li> <li>Process controls which involve changing the way a job acti Enclosure and/or isolation of emission source which keeps and assortypic properly. The design of a ventilation system remover</li></ul>	drift, plating acid fumes, pickling (released at low velocity into zone of active generation)         drift, plating acid fumes, pickling (released at low velocity into zone of radi at motion)         generation into zone of radi at motion)         Within each range the appropriate value depends on:         Lower end of the range       Upper end of the range         1: Room air currents minimal of ravurable 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         Slimple theory shows that air velocity falls rapidly with distance away from the opening of a si with the square of distance from the extraction apperatus, make it essential that theoret more when extraction squeres generated in a tark 2 metes distant from th rootdoring performance deficits within the extraction squeres and with square and 2 meters distant from the contaminating source. The air valcely at the paticular process and chenical concaminants in use.         Process controls which involve changing the way aj ba activity or process is done to redue at the high typically be independent of worker interaction. The basic types of engineering controls are:         Process controls which involve changing the way aj ba activity or process is done to redue at high indycally cancers and chenical concaminants in use.         Employers may need to use multiple types of controls to prevent employee overexpos	direct spray, spray painting in shallow books, drum filling, conveyer loading, crusher dusts, gas discharge (active generation into zone of rapid air motion)           grinding, abrasel batistig, turbing, high speed wheel generated dusts (released at high initial velocity into zone of very high rapid air motion).           With each range the appropriate value depends on:           I. Room air currents minimai of favourable to capital 2. Contaminants of low toxicity or of nuisance value only.         2. Contaminants of high toxicity 3. Infer mitterin, low production.           2. High production, heavy use 4. Large hood or large air mass in motion         3. High production, heavy use 4. Constrainting source and the extraction point (in simple cases). Therefore the air speed at the extraction point, or example, 1.2 mis (200-400 fmin) for extraction of solvents generated in a tank 2 melers distant from the extraction point. Other mit producing performance deficits within the extraction appartus, make it essenial that theoretical air velocities are multipil more when extraction systems are installed or used.           Process controls which involve changing the way alph activity or process is done to reduce the fisk.           Proless controls which involve changing the way alph activity or process is done to reduce the risk.           Engineering controls are used for more value and or place a harrier between the worker and the hazard. Well-designed to highly effective in protecting workers and will pically be independent of worker interactions to provide this high level.           Proless controls which involve changing the way alph activity or process is done to reduce the risk.           Engineering controls are used for merowe a hazard or

· Excellent when breakthrough time > 480 min

· Good when breakthrough time > 20 min

· Fair when breakthrough time < 20 min

 $\cdot$  Poor when glove material degrades

For general applications, gloves with a thickness typically greater than 0.35 mm, are recommended.

It should be emphasised that glove thickness is not necessarily a good predictor of glove resistance to a specific chemical, as the permeation

	<ul> <li>efficiency of the glove will be dependent on the exact composition of the glove material. Therefore, glove selection should also be based on consideration of the task requirements and knowledge of breakthrough times.</li> <li>Glove thickness may also vary depending on the glove manufacturer, the glove type and the glove model. Therefore, the manufacturers technical data should always be taken into account to ensure selection of the most appropriate glove for the task.</li> <li>Note: Depending on the activity being conducted, gloves of varying thickness may be required for specific tasks. For example:</li> <li>Thinner gloves (down to 0.1 mm or less) may be required where a high degree of manual dexterity is needed. However, these gloves are only likely to give short duration protection and would normally be just for single use applications, then disposed of.</li> <li>Thicker gloves (up to 3 mm or more) may be required where there is a mechanical (as well as a chemical) risk i.e. where there is abrasion or puncture potential</li> <li>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.</li> </ul>
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> <li>Some plastic personal protective equipment (PPE) (e.g. gloves, aprons, overshoes) are not recommended as they may produce static electricity.</li> <li>For large scale or continuous use wear tight-weave non-static clothing (no metallic fasteners, cuffs or pockets).</li> <li>Non sparking safety or conductive footwear should be considered. Conductive footwear describes a boot or shoe with a sole made from a conductive compound chemically bound to the bottom components, for permanent control to electrical resistance must range between 0 to 500,000 ohms. Conductive shoes should be stored in lockers close to the room in which they are worn. Personnel who have been issued conductive footwear should not wear them from their place of work to their homes and return.</li> </ul>

### Recommended material(s)

GLOVE SELECTION INDEX

Glove selection is based on a modified presentation of the: "Forsberg Clothing Performance Index".

The effect(s) of the following substance(s) are taken into account in the *computer-generated* selection:

9-Borabicyclo[3.3.1]nonane, 0.5M solution in THF

Material	CPI
PE/EVAL/PE	А
PVA	В
TEFLON	В
BUTYL	С
CPE	С
NEOPRENE	С
VITON/CHLOROBUTYL	С

### **Respiratory protection**

Type A 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 5 x ES	A-AUS / Class 1	-	A-PAPR-AUS / Class 1
up to 25 x ES	Air-line*	A-2	A-PAPR-2
up to 50 x ES	-	A-3	-
50+ x ES	-	Air-line**	-

### ^ - 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 deqC)

- Cartridge respirators should never be used for emergency ingress or in areas of unknown vapour concentrations or oxygen content.
- The wearer must be warned to leave the contaminated area immediately on detecting any odours through the respirator. The odour may indicate that the mask is not functioning properly, that the vapour concentration is too high, or that the mask is not properly fitted. Because of these limitations, only restricted use of cartridge respirators is considered appropriate.
- Cartridge performance is affected by humidity. Cartridges should be changed after 2 hr of continuous use unless it is determined that the humidity is less than 75%, in which case, cartridges can be used for 4 hr. Used cartridges should be discarded daily, regardless of the length of time used

#### B: Satisfactory; may degrade after 4 hours continuous immersion C: Poor to Dangerous Choice for other than short term immersion

\* CPI - Chemwatch Performance Index

NOTE: As a series of factors will influence the actual performance of the glove, a final selection must be based on detailed observation. - \* Where the glove is to be used on a short term, casual or infrequent basis, factors such

as "feel" or convenience (e.g. disposability), may dictate a choice of gloves which might otherwise be unsuitable following long-term or frequent use. A qualified practitioner should be consulted.

#### 8.2.3. Environmental exposure controls

See section 12

A: Best Selection

## **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)	0.884
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)	-105	Viscosity (cSt)	Not Available

Initial boiling point and boiling range (°C)	Not Available	Molecular weight (g/mol)	Not Available
Flash point (°C)	-17	Taste	Not Available
Evaporation rate	Not Available	Explosive properties	Not Available
Flammability	HIGHLY FLAMMABLE.	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>Presence of a stabilising inhibitor prevents/retards peroxide formation.</li> <li>May heat spontaneously</li> <li>Identify and remove sources of ignition and heating.</li> <li>Incompatible material, especially oxidisers, and/or other sources of oxygen may produce unstable product(s).</li> <li>Avoid sources of water contamination (e.g. rain water, moisture, high humidity).</li> <li>Avoid contact with oxygenated solvents/ reagents such as alcohols.</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 toxicological effects

Inhaled	Inhalation of vapours or aerosols (mists, fumes), generated by the material during the course of normal handling, may be harmful. The material can cause respiratory irritation in some persons. The body's response to such irritation can cause further lung damage. Inhalation of vapours may cause drowsiness and dizziness. This may be accompanied by sleepiness, reduced alertness, loss of reflexes, lack of co-ordination, and vertigo. Overexposure to tetrahydrofuran by inhalation may result in irritation of the mucous membrane, and may produce coughing, chest pains, nausea, dizziness, headache and stupor. High concentrations affect the central nervous system.
Ingestion	The material can produce chemical burns within the oral cavity and gastrointestinal tract following ingestion. The material is not thought to produce adverse health effects following ingestion (as classified by EC Directives using animal models). Nevertheless, adverse systemic effects have been produced following exposure of animals by at least one other route and good hygiene practice requires that exposure be kept to a minimum. Ingestion of tetrahydrofuran may not, in itself, produce internal injury, however, contaminating levels of furan, present in certain grades of commercial product, may produce liver and kidney injury exacerbated by the intake of alcoholic beverages. Accidental ingestion of the material may be damaging to the health of the individual.
Skin Contact	The material can produce chemical burns following direct contact with the skin. Skin contact with the material may damage the health of the individual; systemic effects may result following absorption. Open cuts, abraded or irritated skin should not be exposed to this material Entry into the blood-stream, through, for example, cuts, abrasions or lesions, may produce systemic injury with harmful effects. Examine the skin prior to the use of the material and ensure that any external damage is suitably protected. Skin contact with tetrahydrofuran may produce smarting and reddening of the skin and after prolonged exposures; skin inflammation may result because the substance removes skin oils (has a degreasing effect). The material may cause severe inflammation of the skin either following direct contact or after a delay of some time. Repeated exposure can cause contact dermatitis which is characterised by redness, swelling and blistering.
Eye	The material can produce chemical burns to the eye following direct contact. Vapours or mists may be extremely irritating. If applied to the eyes, this material causes severe eye damage. Application of a 20% solution of tetrahydrofuran in water in animals to the eye, produced irritation.

Chronic	Repeated or prolonged exposure to corrosives may result in the erosion (rarely) of the jaw. Bronchial irritation, with cough, and frequent attacks o Long-term exposure to respiratory irritants may result in airways disease There has been concern that this material can cause cancer or mutations Toxic: danger of serious damage to health by prolonged exposure throug This material can cause serious damage if one is exposed to it for long p produce severe defects. Ample evidence from experiments exists that there is a suspicion this ma Substance accumulation, in the human body, may occur and may cause Cyclic ethers can cause cancers, especially of the liver. Repeated exposure to tetrahydrofuran (THF) and related compounds has liver. Animal testing suggests that this group of compounds can cause liv gynaecological disturbance, damage to the adrenal glands and may incre-	f bronchial pneumonia may ensue. , involving difficulty breathing and related whole-body problems. s, but there is not enough data to make an assessment. h inhalation, in contact with skin and if swallowed. eriods. It can be assumed that it contains a substance which can aterial directly reduces fertility. some concern following repeated or long-term occupational exposure. Is been associated with liver inflammation and fatty degeneration of the rer damage, irritation of the skin and airway, metabolic imbalance,
9-Borabicyclo[3.3.1]nonane,	ΤΟΧΙΟΙΤΥ	IRRITATION
0.5M solution in THF	Not Available	Not Available

	ΤΟΧΙΟΙΤΥ	IRRITATION
9-Borabicyclo[3.3.1]nonane	Not Available	Not Available
	ΤΟΧΙΟΙΤΥ	IRRITATION
	dermal (rat) LD50: >2000 mg/kg <sup>[1]</sup>	Eye: adverse effect observed (irritating) <sup>[1]</sup>
tetrahydrofuran	Inhalation(Rat) LC50: 45 mg/l4h <sup>[2]</sup>	Skin: no adverse effect observed (not irritating) <sup>[1]</sup>
	Oral (Rat) LD50: 2816 mg/kg <sup>[2]</sup>	
Legend:	1. Value obtained from Europe ECHA Registered Substances - specified data extracted from RTECS - Register of Toxic Effect	Acute toxicity 2. Value obtained from manufacturer's SDS. Unless otherwise of chemical Substances

TETRAHYDROFURAN	Oral (human) LDLo: 50 mg/kg* [CCINFO]* Nil reported The material may produce severe irritation to the eye of produce conjunctivitis. The material may cause severe skin irritation after pro production of vesicles, scaling and thickening of the sk	causing pronounced inflammation. Re longed or repeated exposure and ma	y produce on contact skin redness, swelling, the
9-Borabicyclo[3.3.1]nonane, 0.5M solution in THF & TETRAHYDROFURAN	Asthma-like symptoms may continue for months or ev known as reactive airways dysfunction syndrome (RAI criteria for diagnosing RADS include the absence of p asthma-like symptoms within minutes to hours of a do airflow pattern on lung function tests, moderate to sew lymphocytic inflammation, without eosinophilia. RADS the concentration of and duration of exposure to the in result of exposure due to high concentrations of irritati	DS) which can occur after exposure to revious airways disease in a non-atop cumented exposure to the irritant. Oth ere bronchial hyperreactivity on meth (or asthma) following an irritating inh ritating substance. On the other hand	b high levels of highly irritating compound. Main bic individual, with sudden onset of persistent ner criteria for diagnosis of RADS include a reversible acholine challenge testing, and the lack of minimal alation is an infrequent disorder with rates related to , industrial bronchitis is a disorder that occurs as a
	disorder is characterized by difficulty breathing, cough	and mucus production.	
Acute Toxicity	disorder is characterized by difficulty breathing, cough	and mucus production.	✓
Acute Toxicity Skin Irritation/Corrosion			
•	<ul> <li>✓</li> </ul>	Carcinogenicity	✓
Skin Irritation/Corrosion	<ul> <li>✓</li> <li>✓</li> <li>✓</li> </ul>	Carcinogenicity Reproductivity	× ×

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

	Endpoint	Test Duration (hr)	Species	Value	Source
9-Borabicyclo[3.3.1]nonane, 0.5M solution in THF	Not Available	Not Available	Not Available	Not Available	Not Available
	Endpoint	Test Duration (hr)	Species	Value	Source

	Endpoint	Test Duration (hr)	Species	Value	Source
tetrahydrofuran	LC50	96h	Fish	1970-2360mg/l	4
	NOEC(ECx)	24h	Fish	>=5mg/l	1
Legend:	Ecotox database	. IUCLID Toxicity Data 2. Europe ECHA Register - Aquatic Toxicity Data 5. ECETOC Aquatic Haz n Data 8. Vendor Data			

For Tetrahydrofuran (THF): Koc: 23 and 18; Henry's Law Constant: 7.1X10-5 atm-m3/mole; Vapor pressure: 162 mm Hg at 25 deg C.

Atmospheric Fate: Tetrahydrofuran exists only as a vapor in the ambient atmosphere. Vapor-phase tetrahydrofuran will be degraded in the atmosphere by reaction with photochemically-produced hydroxyl and nitrate radicals; the half-life for these reactions in air is about 1 and 3 days, respectively. Tetrahydrofuran is moderately reactive in photochemical smog conditions where nitrogen oxides are present; reactions occur in hours under these conditions. Acrolein and formaldehyde have been reported as reaction products.

Terrestrial Fate: Tetrahydrofuran is expected to have very high mobility in soil. Volatilization from moist soil surfaces is expected to be an important fate process. Tetrahydrofuran may also volatilize from dry soil surfaces. Tetrahydrofuran added to surface soil had an abiotic half-life of 5.7 days. Tetrahydrofuran is expected to biodegrade under aerobic conditions but may be resistant to biodegradation in anaerobic environments.

Aquatic Fate: Tetrahydrofuran is slightly persistent in water with a half-life of between 2 to 20 days. If tetrahydrofuran should contact the water table, aquifer or navigable waterway, time is of the essence. It is highly soluble in water and total remediation may not be possible. A comprehensive emergency response or disaster preparedness / recovery plan should be in place prior to use. Volatilization from water surfaces is expected to be an important fate process. Hydrolysis is not expected to occur. Tetrahydrofuran is not expected to adsorb to suspended solids and sediment.

Ecotoxicity: The potential for bioaccumulation of THF in aquatic organisms is considered to be low and it is not expected to bioaccumulate in aquatic organisms. The concentration of tetrahydrofuran in edible fish tissue is expected to be negligible as compared to the levels found in the water from which the fish were taken. Acute short term toxic environmental effects of THF may include the death of animals, birds, fish and death or low growth rate in plants. Acute effects are seen 2 to 4 days after animals or plants are exposed to tetrahydrofuran. Chronic toxic effects include shortened life span, reproductive problems, lowered fertility, and changes in appearance or behavior in exposed animals. These effects have been seen long after the first exposure(s).

# 12.2. Persistence and degradability

Ingredient Per	ersistence: Water/Soil	Persistence: Air
tetrahydrofuran LO	OW	LOW

### 12.3. Bioaccumulative potential

Ingredient	Bioaccumulation
tetrahydrofuran	LOW (LogKOW = 0.46)

#### 12.4. Mobility in soil

Ingredient	Mobility
tetrahydrofuran	LOW (KOC = 4.881)

### 12.5. Results of PBT and vPvB assessment

	Р	В	т
Relevant available data	Not Available	Not Available	Not Available
PBT	×	×	×
vPvB	×	×	×
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	S
Product / Packaging disposal	<ul> <li>Containers may still present a chemical hazard/ danger when empty.</li> <li>Return to supplier for reuse/ recycling if possible.</li> <li>Otherwise: <ul> <li>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.</li> <li>Where possible retain label warnings and SDS and observe all notices pertaining to the product.</li> <li>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.</li> <li>A Hierarchy of Controls seems to be common - the user should investigate: <ul> <li>Reduction</li> <li>Reuse</li> <li>Recycling</li> <li>Disposal (if all else fails)</li> </ul> </li> <li>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. Shell 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.</li> </ul> </li> </ul>

	<ul> <li>Empty containers retain product residues and can be dangerous</li> <li>Dispose of unused product</li> <li>DO NOT expose opened/ empty containers to moisture/ water, heat, flame, sparks, static electricity, or other sources of ignition.</li> <li>They may explode and cause injury or death</li> <li>DO NOT allow wash water from cleaning or process equipment to enter drains.</li> <li>It may be necessary to collect all wash water for treatment before disposal.</li> </ul>
	<ul> <li>In all cases disposal to sever may be subject to local laws and regulations and these should be considered first.</li> <li>Where in doubt contact the responsible authority.</li> <li>Recycle wherever possible or consult manufacturer for recycling options.</li> <li>Consult State Land Waste Authority for disposal.</li> <li>Bury or incinerate residue at an approved site.</li> <li>Recycle containers if possible, or dispose of in an authorised landfill.</li> </ul>
Waste treatment options	Not Available
Sewage disposal options	Not Available

# **SECTION 14 Transport information**

Labels Required	
Marine Pollutant	NO
HAZCHEM	4W

# Land transport (ADR-RID)

14.1. UN number or ID number	3399			
14.2. UN proper shipping name	ORGANOMETALLIC SUBSTANC	ORGANOMETALLIC SUBSTANCE, LIQUID, WATER-REACTIVE, FLAMMABLE		
14.3. Transport hazard class(es)	Class 4.3 Subsidiary risk 3			
14.4. Packing group	II			
14.5. Environmental hazard	Not Applicable			
	Hazard identification (Kemler)	323		
	Classification code	WF1		
14.6. Special precautions for user	Hazard Label	4.3 +3		
	Special provisions	274		
	Limited quantity	500 ml		
	Tunnel Restriction Code	0 (D/E)		

# Air transport (ICAO-IATA / DGR)

14.1. UN number	3399			
14.2. UN proper shipping name		ce, liquid, water-reactive, flammable *		
	ICAO/IATA Class	4.3		
14.3. Transport hazard class(es)	ICAO / IATA Subrisk	3		
01233(03)	ERG Code	4FW		
14.4. Packing group	Ш			
14.5. Environmental hazard	Not Applicable			
	Special provisions		A3 A803	
	Cargo Only Packing Instructions		494	
	Cargo Only Maximum Qty / Pack		5 L	
14.6. Special precautions for user	Passenger and Cargo	Packing Instructions	493	
user	Passenger and Cargo Maximum Qty / Pack		1 L	
	Passenger and Cargo	Passenger and Cargo Limited Quantity Packing Instructions		
	Passenger and Cargo	Limited Maximum Qty / Pack	Forbidden	

### Sea transport (IMDG-Code / GGVSee)

14.1. UN number	3399
14.2. UN proper shipping name	ORGANOMETALLIC SUBSTANCE, LIQUID, WATER-REACTIVE, FLAMMABLE

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### 9-Borabicyclo[3.3.1]nonane, 0.5M solution in THF

14.3. Transport hazard	IMDG Class	4.3
class(es)	IMDG Subrisk	3
14.4. Packing group	П	
14.5. Environmental hazard	Not Applicable	
14.6. Special precautions for user	EMS Number Special provisions Limited Quantities	

#### Inland waterways transport (ADN)

14.1. UN number	3399		
14.2. UN proper shipping name	ORGANOMETALLIC SU	ORGANOMETALLIC SUBSTANCE, LIQUID, WATER REACTIVE, FLAMMABLE	
14.3. Transport hazard class(es)	4.3 3		
14.4. Packing group	П		
14.5. Environmental hazard	Not Applicable		
14.6. Special precautions for user	Classification code Special provisions Limited quantity Equipment required Fire cones number	WF1 274 500 ml PP, EX, A	
	Fire cones number		

### 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
9-Borabicyclo[3.3.1]nonane	Not Available
tetrahydrofuran	Not Available

### 14.7.3. Transport in bulk in accordance with the IGC Code

Product name	Ship Type
9-Borabicyclo[3.3.1]nonane	Not Available
tetrahydrofuran	Not Available

### **SECTION 15 Regulatory information**

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

### 9-Borabicyclo[3.3.1]nonane is found on the following regulatory lists

Not Applicable

# tetrahydrofuran is found on the following regulatory lists

Chemical Footprint Project - Chemicals of High Concern List EU Consolidated List of Indicative Occupational Exposure Limit Values (IOELVs)

EU European Chemicals Agency (ECHA) Community Rolling Action Plan (CoRAP) List of Substances EU REACH Regulation (EC) No 1907/2006 - Annex XVII - Restrictions on the

manufacture, placing on the market and use of certain dangerous substances, mixtures and articles

Europe EC Inventory

European Union - European Inventory of Existing Commercial Chemical Substances (EINECS)

European Union (EU) Regulation (EC) No 1272/2008 on Classification, Labelling and Packaging of Substances and Mixtures - Annex VI

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

International Agency for Research on Cancer (IARC) - Agents Classified by the IARC Monographs - Group 2B: Possibly carcinogenic to humans

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	P5a, P5b, P5c

### 15.2. Chemical safety assessment

No Chemical Safety Assessment has been carried out for this substance/mixture by the supplier.

#### ECHA SUMMARY

Ingredient	CAS number	Index No	ECHA Dossier
tetrahydrofuran	109-99-9	603-025-00-0	Not Available

Harmonisation (C&L Inventory)	Hazard Class and Category Code(s)	Pictograms Signal Word Code(s)	Hazard Statement Code(s)
1	Flam. Liq. 2; Eye Irrit. 2; STOT SE 3	GHS07; GHS02; Dgr	H225; H319; H335
2	Acute Tox. 4; STOT SE 3; Carc. 2; Eye Dam. 1; STOT SE 3; STOT SE 3; Flam. Liq. 1; Acute Tox. 4; Skin Corr. 1B; Acute Tox. 4	GHS08; Dgr; GHS05; GHS01	H302; H335; H336; H351; H318; H370; H312; H224; H314; H332

Harmonisation Code 1 = The most prevalent classification. Harmonisation Code 2 = The most severe classification.

#### **National Inventory Status**

National Inventory	Status
Australia - AIIC / Australia Non-Industrial Use	Yes
Canada - DSL	Yes
Canada - NDSL	No (tetrahydrofuran)
China - IECSC	Yes
Europe - EINEC / ELINCS / NLP	Yes
Japan - ENCS	Yes
Korea - KECI	Yes
New Zealand - NZIoC	Yes
Philippines - PICCS	Yes
USA - TSCA	Yes
Taiwan - TCSI	Yes
Mexico - INSQ	Yes
Vietnam - NCI	Yes
Russia - FBEPH	Yes
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	23/06/2023
Initial Date	23/06/2023

### Full text Risk and Hazard codes

H224	Extremely flammable liquid and vapour.
H250	Catches fire spontaneously if exposed to air.
H260	In contact with water releases flammable gases which may ignite spontaneously.
H312	Harmful in contact with skin.
H318	Causes serious eye damage.
H319	Causes serious eye irritation.
H332	Harmful if inhaled.
H370	Causes damage to organs.

### **SDS Version Summary**

Version	Date of Update	Sections Updated
1.2	23/06/2023	Hazards identification - Classification, Composition / information on ingredients - Ingredients

### 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

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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 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

### Classification and procedure used to derive the classification for mixtures according to Regulation (EC) 1272/2008 [CLP]

Classification according to regulation (EC) No 1272/2008 [CLP] and amendments	Classification Procedure
Flammable Liquids Category 2, H225	Expert judgement
Substances and Mixtures which in Contact with Water Emit Flammable Gases Category 2, H261	Expert judgement
Acute Toxicity (Oral) Category 4, H302	Expert judgement
Skin Corrosion/Irritation Category 1B, H314	Expert judgement
Serious Eye Damage/Eye Irritation Category 1, H318	Minimum classification
Specific Target Organ Toxicity - Single Exposure (Respiratory Tract Irritation) Category 3, H335	Minimum classification
Specific Target Organ Toxicity - Single Exposure (Narcotic Effects) Category 3, H336	Expert judgement
Carcinogenicity Category 2, H351	Calculation method

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