

Benzylmagnesium chloride 1M solution in THF Apollo Scientific

Part Number: OR320068

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: 22/06/2022 Print Date: 31/07/2023 S.REACH.GBR.EN

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

1.1. Product Identifier

Product name	Benzylmagnesium chloride 1M solution in THF							
Chemical Name	Not Applicable							
Synonyms	ot Available							
Proper shipping name	FLAMMABLE LIQUID, CORROSIVE, N.O.S.							
Chemical formula	Not Applicable							
Other means of identification	Not Available							

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	Whitefield Road, Bredbury SK62QR United Kingdom							
Telephone	01614060505							
Fax	0161 406 0506							
Website	http://www.apolloscientific.co.uk/							
Email	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	H250 - Pyrophoric Solids Category 1, H225 - Flammable Liquids Category 2, H335 - Specific Target Organ Toxicity - Single					
regulation (EC) No	Exposure (Respiratory Tract Irritation) Category 3, H315 - Skin Corrosion/Irritation Category 2, H319 - Serious Eye Damage/Eye					
1272/2008 [CLP] and	Irritation Category 2, H228 - Flammable Solids Category 1, H261 - Substances and Mixtures which in Contact with Water Emit					
amendments [1] Flammable Gases Category 2						
Legend:	1. Classified by Chemwatch: 2. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI					

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2.2. Label elements

Hazard pictogram(s)





Signal word

Danger

Hazard statement(s)

H250	Catches fire spontaneously if exposed to air.						
H225	Highly flammable liquid and vapour.						
H335	May cause respiratory irritation.						
H315	Causes skin irritation.						
H319	Causes serious eye irritation.						
H228	Flammable solid.						
H261	In contact with water releases flammable gases.						

Supplementary statement(s)

Not Applicable

Precautionary statement(s) Prevention

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.								
P271	Use only outdoors or in a well-ventilated area.								
P280	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.								
P261	Avoid breathing dust/fumes.								
P222	Do not allow contact with air.								
P223	Do not allow contact with water.								
P264	Wash all exposed external body areas thoroughly after handling.								

Precautionary statement(s) Response

IF ON SKIN: Immerse in cool water or wrap in wet bandages.
IF ON SKIN: Brush off loose particles from skin. Immerse in cool water [or wrap in wet bandages].
In case of fire: Use alcohol resistant foam or normal protein foam to extinguish.
IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.
Call a POISON CENTER/doctor/physician/first aider/if you feel unwell.
If eye irritation persists: Get medical advice/attention.
IF ON SKIN: Wash with plenty of water.
IF ON SKIN (or hair): Take off immediately all contaminated clothing. Rinse skin with water [or shower].
IF INHALED: Remove person to fresh air and keep comfortable for breathing.
If skin irritation occurs: Get medical advice/attention.
Take off contaminated clothing and wash it before reuse.

Precautionary statement(s) Storage

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

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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 ingestion may produce serious health damage*.

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

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		
Not Available	100	Benzylmagnesium chloride 1M solution in THF	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 Regulation (EU) No 1272/2008 - Annex VI; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 3. Classification drawn from Regulation (EU) No 1272/2008 - Annex VI; 3. Classification drawn from Regulation (EU) No 12							

SECTION 4 First aid measures

4.1. Description of first aid measures

Eye Contact	If this product comes in contact with the eyes: Immediately hold eyelids apart and flush the eye continuously with running water. Ensure complete irrigation of the eye by keeping eyelids apart and away from eye and moving the eyelids by occasionally lifting the upper and lower lids. Continue flushing until advised to stop by the Poisons Information Centre or a doctor, or for at least 15 minutes. Transport to hospital or doctor without delay. Removal of contact lenses after an eye injury should only be undertaken by skilled personnel.
Skin Contact	If skin or hair contact occurs: Immediately flush body and clothes with large amounts of water, using safety shower if available. Quickly remove all contaminated clothing, including footwear. Wash skin and hair with running water. Continue flushing with water until advised to stop by the Poisons Information Centre. Transport to hospital, or doctor.
Inhalation	 If fumes or combustion products are inhaled remove from contaminated area. Lay patient down. Keep warm and rested. Prostheses such as false teeth, which may block airway, should be removed, where possible, prior to initiating first aid procedures. Apply artificial respiration if not breathing, preferably with a demand valve resuscitator, bag-valve mask device, or pocket mask as trained. Perform CPR if necessary. Transport to hospital, or doctor, without delay. Inhalation of vapours or aerosols (mists, fumes) may cause lung oedema. Corrosive substances may cause lung damage (e.g. lung oedema, fluid in the lungs). As this reaction may be delayed up to 24 hours after exposure, affected individuals need complete rest (preferably in semi-recumbent posture) and must be kept under medical observation even if no symptoms are (yet) manifested. Before any such manifestation, the administration of a spray containing a dexamethasone derivative or beclomethasone derivative may be considered. This must definitely be left to a doctor or person authorised by him/her. (ICSC13719)
Ingestion	 For advice, contact a Poisons Information Centre or a doctor at once. Urgent hospital treatment is likely to be needed. If swallowed do NOT induce vomiting. If vomiting occurs, lean patient forward or place on left side (head-down position, if possible) to maintain open airway and prevent aspiration. Observe the patient carefully. Never give liquid to a person showing signs of being sleepy or with reduced awareness; i.e. becoming unconscious. Give water to rinse out mouth, then provide liquid slowly and as much as casualty can comfortably drink. Transport to hospital or doctor without delay.

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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

Depending on the degree of exposure, periodic medical examination is indicated. The symptoms of lung oedema often do not manifest until a few hours have passed and they are aggravated by physical effort. Rest and medical observation is therefore essential. Immediate administration of an appropriate spray, by a doctor or a person authorised by him/her should be considered.

(ICSC24419/24421

SECTION 5 Firefighting measures

5.1. Extinguishing media

For SMALL FIRES:

Dry chemical, CO2, water spray or foam.

For LARGE FIRES:

Water-spray, fog or foam.

5.2. Special hazards arising from the substrate or mixture

Fire Incompatibility

Fire Fighting

Avoid contamination with oxidising agents i.e. nitrates, oxidising acids, chlorine bleaches, pool chlorine etc. as ignition may
result

5.3. Advice for firefighters

۰	1	Aler	t Fire	E	3riga	ade	and	tell	th	em	location	and	nature	of	hazard.

- May be violently or explosively reactive.
- Wear full body protective clothing with breathing apparatus.
- ▶ Prevent, by any means available, spillage from entering drains or water course.
- Fight fire from a safe distance, with adequate cover.
- If safe, switch off electrical equipment until vapour fire hazard removed.
- Use water delivered as a fine spray to control the fire and cool adjacent area.
- Avoid spraying water onto liquid pools.
- Do not approach containers suspected to be hot.
- ► Cool fire exposed containers with water spray from a protected location.
- If safe to do so, remove containers from path of fire.

Flammable solid which burns and propagates flame easily, even when partly wetted with water.

- ▶ Any source of ignition, i.e. friction, heat, sparks or flame, may cause fire or explosion.
- May burn fiercely
- ▶ May form explosive mixtures with air.
- May REIGNITE after fire is extinguished.
- Containers may explode on heating.
- Solids may melt and flow when heated or involved in a fire.
- Runoff may pollute waterways.

Fire/Explosion Hazard with

- Avoid generating dust, particularly clouds of dust in a confined or unventilated space as dusts may form an explosive mixture with air. Dust clouds generated by the fine grinding of the solid are a particular hazard; accumulations of fine dust may burn rapidly and fiercely if ignited.
- Dry dust can be charged electrostatically by turbulence, pneumatic transport, pouring, in exhaust ducts and during transport, thereby providing a source of ignition.
- ▶ Decomposition products may be irritating, poisonous or corrosive.

Combustion products include:

carbon dioxide (CO2)

other pyrolysis products typical of burning organic material.

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

 Drains for storage or use areas should have retention basins for pH adjustments and dilution of spills before discharge or disposal of material.

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Check regularly for spills and leaks. Remove all ignition sources. ▶ DO NOT touch or walk through spilled material. Clean up all spills immediately. Avoid contact with skin and eyes. Prevent dust cloud. With clean shovel (preferably non-sparking) place material into clean, dry container and cover loosely. Move containers from spill area. ▶ Control personal contact with the substance, by using protective equipment. Clear area of personnel and move upwind. Alert Fire Brigade and tell them location and nature of hazard. DO NOT touch or walk through spilled material. Wear full protective clothing and breathing apparatus. ▶ May be violently or explosively reactive. ▶ Prevent, by any means available, spillage from entering drains or water course. No smoking, naked lights or ignition sources. Increase ventilation. **Major Spills** Stop leak if safe to do so. Contain or cover with sand, earth or vermiculite. Use only spark-free shovels and explosion proof equipment. Collect recoverable product into labelled containers for recycling. · Collect solid residues and seal in labelled drums for disposal. ▶ Wash area with water and dike for later disposal; prevent runoff into drains. After clean up operations, decontaminate and launder all protective clothing and equipment before storing and re-using.

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

- Avoid all personal contact, including inhalation.
- Wear protective clothing when risk of overexposure occurs.
- ▶ Use in a well-ventilated area.
- Prevent concentration in hollows and sumps.
- DO NOT enter confined spaces until atmosphere has been checked.
- ▶ DO NOT allow material to contact humans, exposed food or food utensils.

If contamination of drains or waterways occurs, advise emergency services.

- Avoid smoking, naked lights or ignition sources.
- When handling, DO NOT eat, drink or smoke. Safe handling
 - Avoid contact with incompatible materials.
 - Keep containers securely sealed when not in use.
 - Avoid physical damage to containers.
 - Always wash hands with soap and water after handling.
 - ▶ Working clothes should be laundered separately. Launder contaminated clothing before re-use.
 - Use good occupational work practice.
 - Observe manufacturer's storage and handling recommendations contained within this SDS.
 - Atmosphere should be regularly checked against established exposure standards to ensure safe working conditions are maintained.

Fire and explosion protection

See section 5

FOR MINOR QUANTITIES:

- ▶ Store in an indoor fireproof cabinet or in a room of noncombustible construction.
- ▶ 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.
 - Protect containers from exposure to weather and from direct sunlight unless: (a) the packages are of metal or plastic construction; (b) the packages are securely closed are not opened for any purpose while in the area where they are stored and (c) adequate precautions are taken to ensure that rain water, which might become contaminated by the dangerous goods, is collected and disposed of safely.
 - Ensure proper stock-control measures are maintained to prevent prolonged storage of dangerous goods.
 - ▶ Observe manufacturer's storage and handling recommendations contained within this SDS.

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7.2. Conditions for safe storage, including any incompatibilities

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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): Removable head packaging and cans with friction closures may be used. Where combination packages are used, there must be sufficient inert absorbent material to absorb completely any leakage that may occur, unless the outer packaging is a close fitting moulded plastic box and the substances are not incompatible with the plastic. All combination packages for Packing group I and II must contain cushioning material.
Storage incompatibility	 Avoid reaction with oxidising agents Moisture sensitive Store under argon
Hazard categories in accordance with Regulation (EC) No 1272/2008	P5a: Flammable Liquids, P5b: Flammable Liquids, P5c: Flammable Liquids, P8: Oxidising Liquid and Solids
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 P8 Lower- / Upper-tier requirements: 50 / 200

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
Benzylmagnesium chloride 1M solution in THF	Not Available	Not Available	Not Available

Ingredient	Original IDLH	Revised IDLH
Benzylmagnesium chloride 1M solution in THF	Not Available	Not Available

8.2. Exposure controls

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

8.2.1. Appropriate engineering controls

The basic types of engineering controls are:

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

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

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

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

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

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

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

Within each range the appropriate value depends on:

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

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

For large scale or continuous use:

- ▶ Spark-free, earthed ventilation system, venting directly to the outside and separate from usual ventilation systems
- ▶ Provide dust collectors with explosion vents

8.2.2. Individual protection measures, such as personal protective equipment









Eye and face protection

- ► Chemical goggles. [AS/NZS 1337.1, EN166 or national equivalent]
- Full face shield may be required for supplementary but never for primary protection of eyes.
- ▶ 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].

Skin protection

See Hand protection below

Hands/feet protection

- ▶ When handling corrosive liquids, wear trousers or overalls outside of boots, to avoid spills entering boots.
- Wear physical protective gloves, e.g. leather.
- Wear safety footwear.

Body protection

See Other protection below

- ▶ Some plastic personal protective equipment (PPE) (e.g. gloves, aprons, overshoes) are not recommended as they may produce static electricity.
- For large scale or continuous use wear tight-weave non-static clothing (no metallic fasteners, cuffs or pockets).

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 electrically ground the foot an shall dissipate static electricity from the body to reduce the possibility of ignition of volatile compounds. 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.

- Other protection
- Overalls. PVC Apron.

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- ▶ PVC protective suit may be required if exposure severe. ► Eyewash unit.
- Ensure there is ready access to a safety shower.

Respiratory protection

Type -P Filter of sufficient capacity. (AS/NZS 1716 & 1715, EN 143:2000 & 149:2001, ANSI Z88 or national equivalent)

Required Minimum Protection Factor	Half-Face Respirator	Full-Face Respirator	Powered Air Respirator
up to 10 x ES	P1 Air-line*	-	PAPR-P1
up to 50 x ES	Air-line**	P2	PAPR-P2
up to 100 x ES	-	P3	-
		Air-line*	-
100+ x ES	-	Air-line**	PAPR-P3

^{* -} Negative pressure demand ** - Continuous flow

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

8.2.3. Environmental exposure controls

See section 12

SECTION 9 Physical and chemical properties

9.1. Information on basic physical and chemical properties

Physical state Divided Solid Partition coefficient n-octanol / water Odour Not Available Permitting point / freezing point (*C) Initial boiling point and boiling range (*C) Flash point (*C) Not Available Evaporation rate Not Available Flammability Not Available Divided Solid Evaporation rate Not Available Volatile Component (%vol) Not Available Divided Solid Evaporation rate Not Available	Appearance	Not Available		
Odour Not Available Odour threshold Not Available Partition coefficient n-octanol / water Odour threshold Not Available Partition coefficient n-octanol / water Not Available Not Available Not Available Not Available Partition coefficient n-octanol / water (°C) Not Available Flash point (°C) Not Available Flash point (°C) Not Available Evaporation rate Not Available Flammability Not Available Cydidising properties Not Available Upper Explosive Limit (%) Not Available Volatile Component (%vol) Not Available Vapour pressure (kPa) Not Available Partition coefficient n-octanol / water Not Available Not Available Not Available Vapour density (Air = 1) Not Available				
Odour threshold Not Available	Physical state	Divided Solid		Not Available
pH (as supplied) Not Available PH (as supplied) Not Available Relting point / freezing point (°C) Initial boiling point and boiling range (°C) Flash point (°C) Not Available Molecular weight (g/mol) Flash point (°C) Not Available Molecular weight (g/mol) Not Available Flammability Not Available Explosive properties Not Available Flammability Not Available Oxidising properties Not Available Upper Explosive Limit (%) Not Available Volatile Component (%vol) Vapour pressure (kPa) Not Available Phas a solution (1%) Not Available	Odour	Not Available		Not Available
Melting point / freezing point (°C) Initial boiling point and boiling range (°C) Not Available Molecular weight (g/mol) Not Available Flash point (°C) Not Available Evaporation rate Not Available Flammability Not Available Cyaporation rate Not Available Flammability Not Available Surface Tension (dyn/cm or mN/m) Not Applicable Vapour pressure (kPa) Not Available Volatile Component (%vol) Not Available PH as a solution (1%) Not Available	Odour threshold	Not Available		Not Available
Initial boiling point and boiling range (°C) Initial boiling range (°C) Flash point (°C) Not Available Flash point (°C) Not Available Evaporation rate Not Available Flammability Not Available Cyapour pressure (kPa) Solubility in water Viscosity (cSt) Not Available Molecular weight (g/mol) Not Available Explosive properties Not Available Explosive properties Not Available Oxidising properties Not Available Surface Tension (dyn/cm or mN/m) Not Applicable Volatile Component (%vol) Not Available Gas group Not Available Vapour pressure (kPa) Not Available PH as a solution (1%) Not Available VOC g/L Not Available	pH (as supplied)	Not Available	•	Not Available
Not Available Not Available Not Available Not Available	· · · · · · · · · · · · · · · · · · ·	Not Available	Viscosity (cSt)	Not Available
Evaporation rate Not Available Flammability Not Available Oxidising properties Not Available Upper Explosive Limit (%) Not Available Volatile Component (%vol) Vapour pressure (kPa) Solubility in water Vapour density (Air = 1) Not Available	• .	Not Available	Molecular weight (g/mol)	Not Available
Flammability Not Available Oxidising properties Not Available Surface Tension (dyn/cm or mN/m) Not Applicable Lower Explosive Limit (%) Not Available Volatile Component (%vol) Vapour pressure (kPa) Not Available Solubility in water Vapour density (Air = 1) Not Available	Flash point (°C)	Not Available	Taste	Not Available
Upper Explosive Limit (%) Not Available Surface Tension (dyn/cm or mN/m) Lower Explosive Limit (%) Not Available Volatile Component (%vol) Not Available Vapour pressure (kPa) Not Available Solubility in water Vapour density (Air = 1) Not Available	Evaporation rate	Not Available	Explosive properties	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 Nanoform Solubility Not Available Not Available Not Available Not Available Not Available Not Available Not Available Not Available	Flammability	Not Available	Oxidising properties	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 Not Available Not Available	Upper Explosive Limit (%)	Not Available		Not Applicable
Solubility in water Not Available pH as a solution (1%) Not Available Vapour density (Air = 1) Not Available VOC g/L Nanoform Solubility Not Available Not Available Not Available Not Available Not Available	Lower Explosive Limit (%)	Not Available	Volatile Component (%vol)	Not Available
Vapour density (Air = 1) Not Available VOC g/L Not Available Nanoform Solubility Not Available Nanoform Particle Characteristics Not Available	Vapour pressure (kPa)	Not Available	Gas group	Not Available
Nanoform Solubility Not Available Not Available Not Available Not Available	Solubility in water	Not Available	pH as a solution (1%)	Not Available
Nanoform Solubility Not Available Characteristics Not Available	Vapour density (Air = 1)	Not Available	VOC g/L	Not Available
Particle Size Not Available	Nanoform Solubility	Not Available		Not Available
	Particle Size	Not Available		

9.2. Other information

Not Available

SECTION 10 Stability and reactivity

Part Number: **OR320068** Version No: **1.1**

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10.1.Reactivity	See section 7.2
10.2. Chemical stability	 Unstable in the presence of incompatible materials. Product is considered stable. Hazardous polymerisation will not occur.
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

Inhaled	The material can cause respiratory irritation in some persons. The body's response to such irritation can cause further lung damage. The material has NOT been classified by EC Directives or other classification systems as "harmful by inhalation". This is because of the lack of corroborating animal or human evidence. Inhalation of quantities of liquid mist may be extremely hazardous, even lethal due to spasm, extreme irritation of larynx and bronchi, chemical pneumonitis and pulmonary oedema.
Ingestion	The material can produce chemical burns within the oral cavity and gastrointestinal tract following ingestion. The material has NOT been classified by EC Directives or other classification systems as "harmful by ingestion". This is because of the lack of corroborating animal or human evidence.
Skin Contact	The material can produce chemical burns following direct contact with the skin. Skin contact is not thought to have harmful health effects (as classified under EC Directives); the material may still produce health damage following entry through wounds, lesions or abrasions. 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.
Eye	The material can produce chemical burns to the eye following direct contact. Vapours or mists may be extremely irritating.
Chronic	Repeated or prolonged exposure to corrosives may result in the erosion of teeth, inflammatory and ulcerative changes in the mouth and necrosis (rarely) of the jaw. Bronchial irritation, with cough, and frequent attacks of bronchial pneumonia may ensue. Long-term exposure to respiratory irritants may result in airways disease, involving difficulty breathing and related whole-body problems. Substance accumulation, in the human body, may occur and may cause some concern following repeated or long-term occupational exposure.

Benzylmagnesium chloride	TOXICITY	IRRITATION
1M solution in THF	Not Available	Not Available
Legend:	1. Value obtained from Europe ECHA Registered Substances - Acute toxicity 2. Value obtained from manufacturer's SDS. Unless otherwise specified data extracted from RTECS - Register of Toxic Effect of chemical Substances	

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Asthma-like symptoms may continue for months or even years after exposure to the material ends. This may be due to a non-allergic condition known as reactive airways dysfunction syndrome (RADS) which can occur after exposure to high levels of highly irritating compound. Main criteria for diagnosing RADS include the absence of previous airways disease in a non-atopic individual, with sudden onset of persistent asthma-like symptoms within minutes to hours of a documented exposure to the irritant. Other criteria for diagnosis of RADS include a reversible airflow pattern on lung function tests, moderate to severe bronchial hyperreactivity on methacholine challenge testing, and the lack of minimal lymphocytic inflammation, without eosinophilia. RADS (or asthma) following an irritating inhalation is an infrequent disorder with rates related to the concentration of and duration of exposure to the irritating substance. On the other hand, industrial bronchitis is a disorder that occurs as a result of exposure due to high concentrations of irritating substance (often particles) and is completely reversible after exposure ceases. The disorder is characterized by difficulty breathing, cough and mucus production.

Acute Toxicity	×	Carcinogenicity	×
Skin Irritation/Corrosion	~	Reproductivity	×
Serious Eye Damage/Irritation	✓	STOT - Single Exposure	✓
Respiratory or Skin sensitisation	×	STOT - Repeated Exposure	×
Mutagenicity	×	Aspiration Hazard	×

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Legend:

★ - 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

Danmalmannasium ablasida	Endpoint	Test Duration (hr)	Species	Value	Source
Benzylmagnesium chloride 1M solution in THF	Not Available	Not Available	Not Available	Not Available	Not Available
Legend:	4. US EPA, E	n 1. IUCLID Toxicity Data 2. Europe ECHA R icotox database - Aquatic Toxicity Data 5. EC tion Data 7. METI (Japan) - Bioconcentration	ETOC Aquatic Hazard Assessment Data 6. I	•	

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

P	В	Т
Not Available	Not Available	Not Available
x	×	×
×	X	×
		No
		No
	Not Available	Not Available Not Available X

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

► Recycle wherever possible.

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▶ Consult manufacturer for recycling options or consult local or regional waste management authority for disposal if no suitable treatment or disposal facility can be identified. disposal ▶ Treat and neutralise at an approved treatment plant. Treatment should involve: Neutralisation followed by: burial in a land-fill specifically licensed to accept chemical and / or pharmaceutical wastes or Incineration in a licensed apparatus. ▶ Decontaminate empty containers. Observe all label safeguards until containers are cleaned and destroyed. Waste treatment options Not Available Sewage disposal options Not Available

SECTION 14 Transport information

Labels Required



Marine Pollutant HAZCHEM •3WE

Land transport (ADR-RID)

<u> </u>	•		
14.1. UN number or ID number	2924		
14.2. UN proper shipping name	FLAMMABLE LIQUID, CORROSI	VE, N.O.S.	
14.3. Transport hazard class(es) Class 3 Subsidiary risk 8			
14.4. Packing group	II		
14.5. Environmental hazard	Not Applicable		
	Hazard identification (Kemler)	338	
	Classification code	FC	
14.6. Special precautions	Hazard Label	3 +8	
for user	Special provisions	274	
	Limited quantity	1 L	
	Tunnel Restriction Code	2 (D/E)	

Air transport (ICAO-IATA / DGR)

14.1. UN number	2924			
14.2. UN proper shipping name	Flammable liquid, corrosive, n.o.s. *			
	ICAO/IATA Class	3		
14.3. Transport hazard class(es)	ICAO / IATA Subrisk	8		
0.400(00)	ERG Code	3CH		
14.4. Packing group	II			
14.5. Environmental hazard	Not Applicable			
	Special provisions		A3 A803	
	Cargo Only Packing Ir	nstructions	363	
	Cargo Only Maximum	Qty / Pack	5 L	
14.6. Special precautions for user	Passenger and Cargo	Packing Instructions	352	
ioi usci	Passenger and Cargo	Maximum Qty / Pack	1 L	
	Passenger and Cargo	Limited Quantity Packing Instructions	Y340	
	Passenger and Cargo	Limited Maximum Qty / Pack	0.5 L	

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Sea transport (IMDG-Code / GGVSee)

14.1. UN number	2924		
14.2. UN proper shipping name	FLAMMABLE LIQUID	, CORROSIVE, N.O.S.	
14.3. Transport hazard	IMDG Class 3	_	
class(es)	IMDG Subrisk 8		
14.4. Packing group	II		
14.5. Environmental hazard	Not Applicable	Not Applicable	
	EMS Number	F-E, S-C	
14.6. Special precautions for user	Special provisions	274	
101 4001	Limited Quantities	1L	

Inland waterways transport (ADN)

14.1. UN number	2924		
14.2. UN proper shipping name	FLAMMABLE LIQUID, O	CORROSIVE, N.O.S.	
14.3. Transport hazard class(es)	3 8		
14.4. Packing group	II		
14.5. Environmental hazard	Not Applicable	Not Applicable	
	Classification code	FC	
	Special provisions	274	
14.6. Special precautions for user	Limited quantity	1L	
101 4001	Equipment required	PP, EP, EX, A	
	Fire cones number	1	

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 in accordance with the IGC Code

Draduat name	Chin Tuna
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	P5a, P5b, P5c, P8

15.2. Chemical safety assessment

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

ECHA SUMMARY

Not Applicable

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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
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	22/06/2022
Initial Date	22/06/2022

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 Part Number: OR320068 Page 14 of 14 Issue Date: 22/06/2022 Version No: 1.1 Print Date: 31/07/2023

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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

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