



Previous revision: 22/04/2022 Version: 5 Revision: 08/03/2023 Date of printing: 08/03/2023

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

## PRODUCT IDENTIFIER:

**REVESTIDAN FIBER** 

UFI: MH00-C0TV-Q000-1R0J

RELEVANT IDENTIFIED USES OF THE SUBSTANCE OR MIXTURE AND USES ADVISED AGAINST: 1.2

Intended uses (main technical functions): [] Industrial [] Professional [X] Consumers

Waterproof Membrane

Sectors of use:

Professional uses (SU22).

Uses advised against:

This product is not recommended for any use or sector of use (industrial, professional or consumer) other than those previously listed as "Intended or identified uses".

Restrictions on manufacture, placing on market and use, according to Annex XVII of Regulation (EC) No. 1907/2006:

Not restricted

#### **DETAILS OF THE SUPPLIER OF THE SAFETY DATA SHEET:** 1.3

DANOSA - DERIVADOS ASFÁLTICOS NORMALIZADOS, S.A.

Polígono Industrial, Sector 9 - 19290 Fontanar (Guadalajara) ESPAÑA Phone number: 949888210 - Fax: 949 888 223 - www.danosa.com

- E-mail address of the person responsible for the Safety Data Sheet:

info@danosa.com

#### **EMERGENCY TELEPHONE NUMBER:** 1.4

902 422 452 8:30-17:30 h



National Poisons Information Service (NPIS) - In England, Wales or Scotland: dial 111 - In N Ireland: contact your local GP or pharmacist during normal hours.

#### SECTION 2 : HAZARDS IDENTIFICATION

#### **CLASSIFICATION OF THE SUBSTANCE OR MIXTURE:** 2.1

Classification of mixtures is carried out in accordance with the following principles: a) when data (tests) for the classification of mixtures are available, generally is carried out based on these data, b) in the absence of data (tests) for mixtures are generally used interpolation or extrapolation methods of assessing the risk, using the available data for mixtures similarly classified, and c) in the absence of tests and information which would allow to apply interpolation or extrapolation techniques, methods are used to classify risk assessment based on the data of the individual components in the mixture.

Classification in accordance with Regulation (EU) No. 1272/2008~2021/849 (CLP):

WARNING: Skin Sens. 1:H317|Aquatic Chronic 3:H412

Danger class	Classification of the mixture	Cat.	Routes of exposure	Target organs	Effects
Physicochemical: Not classified					
Human health:	🥦 Skin Sens. 1:H317 c)	Cat.1	Skin	Skin	Allergy
Environment:	Aquatic Chronic 3:H412 c)	Cat.3	-	-	-

Full text of hazard statements mentioned is indicated in section 16.

Note: When in section 3 a range of percentages is used, the health and environmental hazards describe the effects of the highest concentration of each component, but below the maximum value.

#### LABEL ELEMENTS: 2.2



This product is labelled with the signal word WARNING in accordance with Regulation (EU) No. 1272/2008~2021/849 (CLP)

### Hazard statements:

H317 May cause an allergic skin reaction.

H412 Harmful to aquatic life with long lasting effects.

### - Precautionary statements:

P280 Wear protective gloves, clothing and eye protection. In case of inadequate ventilation wear respiratory protection.

P362+P364 Take off contaminated clothing and wash it before reuse.

P363 Wash contaminated clothing before reuse.

IF ON SKIN (or hair): Take off immediately all contaminated clothing. Rinse skin with water [or shower]. Wash with P303+P361+P353-

plenty of water and soap.. Call a POISON CENTER or doctor if you feel unwell. P352-P312

Avoid release to the environment. Dispose of contents/container in accordance with local regulations. P273-P501

#### Supplementary statements:

**EUB174** Contains 2-octyl-2H-isothiazol-3-one to protect the film.

#### Substances that contribute to classification:

2-octyl-2H-isothiazol-3-one Other sensitizing components:

1,2-benzisothiazol-3(2H)-one

2-methylisothiazol-3(2H)-one

REACH / ATP13

REACH / ATP15

Skin Sens. 1A, H317: C ≥0,0015 %

Skin Sens. 1A, H317:

C ≥0.0015 %



#### **REVESTIDAN FIBER**



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#### **OTHER HAZARDS:** 2.3

Hazards which do not result in classification but which may contribute to the overall hazards of the mixture:

- Other physicochemical hazards:

No other relevant adverse effects are known.

- Other adverse human health effects:

No other relevant adverse effects are known.

- Other negative environmental effects:

Does not contain substances that fulfil the PBT/vPvB criteria.

Endocrine disrupting properties:

This product contains substances with endocrine disrupting properties identified or under evaluation in a concentration of less than 0.1% by weight:Diuron (ISO)

## SECTION 3: COMPOSITION/INFORMATION ON INGREDIENTS

#### 3.1 SUBSTANCES:

Not applicable (mixture).

#### 3.2 MIXTURES:

This product is a mixture.

Chemical description:

Solution of Calcium carbonate in aqueous media.

### **HAZARDOUS INGREDIENTS:**

Substances taking part in a percentage higher than the exemption limit:

C < 0.01 %



2-methylisothiazol-3(2H)-one

CAS: 2682-20-4, EC: 220-239-6, REACH: 01-2120764690-50 CLP: Danger: Acute Tox. (inh.) 2:H330 | Acute Tox. (skin) 3:H311 | Acute Tox. (oral) 3:H301 | Skin Corr. 1B:H314 | Eye Dam. 1:H318 | Aquatic Acute 1:H400 (M=10) | Aquatic Chronic 1:H410 (M=1) | EUH071 | Skin Sens.

1A:H317

C < 0,0050 %



2-octyl-2H-isothiazol-3-one

CAS: 26530-20-1, EC: 247-761-7, REACH: 01-2120768921-45 CLP: Danger: Acute Tox. (inh.) 2:H330 | Acute Tox. (skin) 3:H311 (ATE=311 mg/kg) | Acute Tox. (oral) 3:H301 (ATE=125 mg/kg) | Skin Corr. 1B:H314 | Eye Dam. 1:H318 | Aquatic Acute 1:H400 (M=100) | Aquatic Chronic 1:H410

(M=100) | EUH071 | Skin Sens. 1A:H317

### Impurities:

Does not contain other components or impurities which will influence the classification of the product.

Stabilizers:

None

# Reference to other sections:

For more information on hazardous ingredients, see sections 8, 11, 12 and 16.

SUBSTANCES OF VERY HIGH CONCERN (SVHC):

List updated by ECHA on 17/01/2023.

Substances SVHC subject to authorisation, included in Annex XIV of Regulation (EC) no. 1907/2006:

None

Substances SVHC candidate to be included in Annex XIV of Regulation (EC) no. 1907/2006:

None

PERSISTENT. BIOACCUMULABLE AND TOXIC PBT. OR VERY PERSISTENT AND VERY BIOACCUMULABLE VPVB SUBSTANCES:

Does not contain substances that fulfil the PBT/vPvB criteria.





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#### SECTION 4: FIRST AID MEASURES

#### 4.1 DESCRIPTION OF FIRST AID MEASURES:



Symptoms may occur after exposure, so that in case of direct exposure to the product, when in doubt, or when symptoms persist, seek medical attention. Never give anything by mouth to an unconscious person. Lifeguards should pay attention to self-protection and use the recommended protective equipment if there is a possibility of exposure. Wear protective gloves when administering first aid.

Route of exposure	Symptoms and effects, acute and delayed	Description of first-aid measures
Inhalation:	It is not expected that symptoms will occur under normal conditions of use.	Remove the patient out of the contaminated area into the fresh air. If breathing is irregular or stops, administer artificial respiration. If the person is unconscious, place in appropriate recovery position. Keep the patient warm and at rest until medical attention arrives.
Skin:	Skin contact causes redness.	Remove immediately contaminated clothing.Wash thoroughly the affected area with plenty of cold or lukewarm water and neutral soap, or use a suitable skin cleanser.
Eyes:	Contact with the eyes may produce slight redness.	Remove contact lenses.Rinse eyes copiously by irrigation with plenty of clean, fresh water, holding the eyelids apart.If irritation persists, consult a physician.
Ingestion:	If swallowed, may cause gastrointestinal disturbances.	If swallowed, seek medical advice immediately and show container or label. Do not induce vomiting, due to the risk of aspiration.Keep the patient at rest.

#### 4.2 MOST IMPORTANT SYMPTOMS AND EFFECTS, BOTH ACUTE AND DELAYED:

The main symptoms and effects are indicated in sections 4.1 and 11.1

INDICATION OF ANY IMMEDIATE MEDICAL ATTENTION AND SPECIAL TREATMENT NEEDED:

Notes to physician:

4.3

Treatment should be directed at the control of symptoms and the clinical condition of the patient...

Antidotes and contraindications:

Specific antidote not known.

## SECTION 5: FIREFIGHTING MEASURES

# 5.1 <u>EXTINGUISHING MEDIA:</u>)

In case of fire in the surroundings, all extinguishing agents are allowed

## 5.2 SPECIAL HAZARDS ARISING FROM THE SUBSTANCE OR MIXTURE:

As consequence of combustion or thermal decomposition, hazardous products may be produced: carbon monoxide, Carbon dioxide, nitrogen oxides, sulfur oxides. Exposure to combustion or decomposition products may be a hazard to health.

# 5.3 ADVICE FOR FIREFIGHTERS:

# Special protective equipment:

Depending on magnitude of fire, heat-proof protective clothing may be required, appropriate independent breathing apparatus, gloves, protective glasses or face masks and boots. If the fire-proof protective equipment is not available or is not being used, combat fire from a sheltered position or from a safe distance. The standard EN469 provides a basic level of protection for chemical incidents.

#### Other recommendations:

Cool with water the tanks, cisterns or containers close to sources of heat or fire.Bear in mind the direction of the wind.Do not allow fire-fighting residue to enter drains, sewers or water courses.





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SECTION	I 6: ACCIDENTAL RELEASE MEASURES
6.1	PERSONAL PRECAUTIONS, PROTECTIVE EQUIPMENT AND EMERGENCY PROCEDURES:
	Avoid direct contact with this product. Avoid breathing vapours. Keep people without protection in opposition to the wind direction.
6.2	ENVIRONMENTAL PRECAUTIONS:
	Avoid contamination of drains, surface or subterranean water and soil.In the case of large scale spills or when the product contaminates lakes, rivers or sewages, inform the appropriate authorities in accordance with local regulations.
6.3	METHODS AND MATERIAL FOR CONTAINMENT AND CLEANING UP:
	Contain and mop up spills with absorbent materials (sawdust, earth, sand, vermiculite, diatomaceous earth, etc). Keep the remains in a closed container.
6.4	REFERENCE TO OTHER SECTIONS:
	For contact information in case of emergency, see section 1. For information on safe handling, see section 7. For exposure controls and personal protection measures, see section 8. For waste disposal, follow the recommendations in section 13.

#### SECTION 7: HANDLING AND STORAGE

### 7.1 PRECAUTIONS FOR SAFE HANDLING:

Comply with the existing legislation on health and safety at work.

- General recommendations:

Avoid any type of leakage or escape. Keep the container tightly closed.

- Recommendations for the prevention of fire and explosion risks:

The product is not liable to ignite, deflagrate or explode, and does not sustain the combustion reaction by oxygen from air in the environment in which it is, so it is not included in the scope of Directive 2014/34/EU concerning equipment and protective systems intended for use in potentially explosive atmospheres.

- Recommendations for the prevention of toxicological risks:

Do not eat, drink or smoke while handling. After handling, wash hands with soap and water. For exposure controls and personal protection measures, see section 8.

- Recommendations for the prevention of environmental contamination:

Avoid any spillage in the environment. Pay special attention to the cleaning water. In the case of accidental spillage, follow the instructions indicated in section 6.

### 7.2 CONDITIONS FOR SAFE STORAGE, INCLUDING ANY INCOMPATIBILITIES:

Forbid the entry to unauthorized persons. Keep out of reach of children. Keep away from sources of heat. If possible, avoid direct contact with sunlight. In order to avoid leakages, the containers, after use, should be closed carefully and placed in a vertical position. For more information, see section 10.

- Class of store:

According to current legislation.

- Maximum storage period:

24 Months.

- Temperature interval:

min:5 °C, max:30 °C (recommended).

- Incompatible materials:

Keep away from reducing agents, oxidizing agents, acids, alkalis.

- Type of packaging:

According to current legislation.

- Limit quantity (Seveso III): Directive 2012/18/EU:

Not applicable (the classification criteria are not met).

### 7.3 SPECIFIC END USE(S):

For the use of this product particular recommendations apart from that already indicated are not available.





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#### SECTION 8: EXPOSURE CONTROLS/PERSONAL PROTECTION

#### 8.1 CONTROL PARAMETERS:

If a product contains ingredients with exposure limits, may be necessary a personnel monitoring, work place or biological, to determine the effectiveness of the ventilation or other control measures and/or the necessity to use respiratory protective equipment. Reference should be made to EN689, EN14042 and EN482 standard concerning methods for assessing the exposure by inhalation to chemical agents, and exposure to chemical and biological agents. Reference should be also made to national guidance documents for methods for the determination of dangerous substances.

OCCUPATIONAL EXPOSURE LIMIT VALUES (WEL)

Not established

#### - BIOLOGICAL LIMIT VALUES:

Biological monitoring can be a very useful complementary technique to air monitoring when air sampling techniques alone may not give a reliable indication of exposure. Biological monitoring is the measurement and assessment of hazardous substances or their metabolites in tissues, secretions, excreta or expired air, or any combination of these, in exposed workers. Measurements reflect absorption of a substance by all routes. Biological monitoring may be particularly useful in circumstances where there is likely to be significant skin absorption and/or gastrointestinal tract uptake following ingestion, where control of exposure depends on respiratory protective equipment, where there is a reasonably well-defined relationship between biological monitoring and effect, or where it gives information on accumulated dose and target organ body burden which is related to toxicity.

This preparation contains the following substances that have established a biological limit value:

- N,N-dimethylformamide (2016): 1°) Biological determinant: N-methylformamide in urine (total N-methylformamide represents the sum of N-methylformamide and N-(hydroxymethyl)-N-methylformamide), BEI: 30 mg/l, Sampling time: end of shift (2). 2°) Biological determinant: N-acetyl-S-(N-methylcarbamoyl)cysteine in urine, BEI: 30 mg/l, Sampling time: end of shift at end of workweek (4).

These indicators accumulate in the body during the work week, therefore the sampling time is critical in relation to previous exposures. (2) When the end of the exposition not coincide with the end of the working day, the sample will be taken as soon as possible after the real exposition ceases. Once the steady state that depends on each biological indicator (weeks, months) has been reached, sampling of these can be done at any time. &The biological determinant is an indicator of exposure to the chemical, but the quantitative interpretation of the measurement is ambiguous. &(CDC: Guidelines for the identification and management of lead exposure in pregnant and lactating women, 2010).

(4) The value refers to the difference of the results of the samples taken at the end and at the beginning of the working day.

#### - DERIVED NO-EFFECT LEVEL (DNEL):

Derived no-effect level (DNEL) is a level of exposure that is considered safe, derived from toxicity data according to specific guidances included in REACH. DNEL values may differ from a occupational exposure limit (OEL) for the same chemical. OEL values may come recommended by a particular company, a government regulatory agency or an organization of experts. Although considered protective of health, the OEL values are derived by a process different of REACH.

- DERIVED NO-EFFECT LEVEL, WORKERS:- Systemic effects, acute and chronic:	DNEL Inhalation mg/m3		DNEL Cutaneous mg/kg bw/d		DNEL Oral mg/kg bw/d	
2-methylisothiazol-3(2H)-one	s/r (a)	s/r (c)	s/r (a)	s/r (c)	- (a)	- (c)
2-octyl-2H-isothiazol-3-one	- (a)	- (c)	- (a)	- (c)	- (a)	- (c)
- DERIVED NO-EFFECT LEVEL, WORKERS:- Local effects, acute and chronic:	DNEL Inhalation mg/m3		DNEL Cutaneous mg/cm2		DNEL Eyes mg/cm2	
2-methylisothiazol-3(2H)-one	0,043 (a)	0,021 (c)	m/r <b>(a)</b>	s/r (c)	a/r <b>(a)</b>	- (c)
2-octyl-2H-isothiazol-3-one	- (a)	- (c)	- (a)	- (c)	- (a)	- (c)

## - Derived no-effect level, general population:

Not applicable (product for professional or industrial use).

- (a) Acute, short-term exposure, (c) Chronic, long-term or repeated exposure.
- (-) DNEL not available (without data of registration REACH).
- s/r DNEL not derived (not identified hazard).
- m/r DNEL not derived (medium hazard).
- a/r DNEL not derived (high hazard).

# - PREDICTED NO-EFFECT CONCENTRATION (PNEC):

	,		
- PREDICTED NO-EFFECT CONCENTRATION,	PNEC Fresh water	PNEC Marine	PNEC Intermittent
AQUATIC ORGANISMS:- Fresh water, marine	mg/l	mg/l	mg/l
water and intermittent release:			
2-methylisothiazol-3(2H)-one	0.00339	0.00339	-
2-octyl-2H-isothiazol-3-one	0.0022	0.00022	0.000122
- WASTEWATER TREATMENT PLANTS (STP)	PNEC STP	PNEC Sediments	PNEC Sediments
AND SEDIMENTS IN FRESH- AND MARINE	mg/l	mg/kg dw/d	mg/kg dw/d
WATER:			
2-methylisothiazol-3(2H)-one	0.23	s/r	s/r
2-octyl-2H-isothiazol-3-one	s/r	0.0475	0.00475
- PREDICTED NO-EFFECT CONCENTRATION,	PNEC Air	PNEC Soil	PNEC Oral
TERRESTRIAL ORGANISMS:- Air, soil and	mg/m3	mg/kg dw/d	mg/kg dw/d
effects for predators and humans:			
2-methylisothiazol-3(2H)-one	s/r	0.047	n/b
2-octyl-2H-isothiazol-3-one	s/r	0.0082	n/b

n/b - PNEC not derived (not bioaccumulative potential).

s/r - PNEC not derived (not identified hazard).





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Provide adequate ventilation. Where reasonably practicable, this should be achieved by the use of local exhaust ventilation and good general extraction.

- Protection of respiratory system:

Avoid the inhalation of vapours.

- Protection of eyes and face:

It is recommended to install water taps, sources or eyewash bottles with clean water close to the working area.

- Protection of hands and skin:

It is recommended to install water taps or sources with clean water close to the working area.Barrier creams may help to protect the exposed areas of the skin.Barrier creams should not be applied once exposure has occurred.

## OCCUPATIONAL EXPOSURE CONTROLS: REGULATION (EU) NO. 2016/425:

As a general measure on prevention and safety in the work place, we recommend the use of a basic personal protection equipment (PPE), with the corresponding marking. For more information on personal protective equipment (storage, use, cleaning, maintenance, type and characteristics of the PPE, protection class, marking, category, CEN norm, etc..), you should consult the informative brochures provided by the manufacturers of PPE

Mask:  Safety goggles:  Face shield:  Gloves:	No.  Safety goggles designed to protect against liquid splashes, with suitable lateral protection (EN166). Clean daily and disinfect at regular intervals in accordance with the instructions of the manufacturer.  No.  Gloves resistant against chemicals (EN374). When repeated or prolonged contact with the product is
Face shield: Gloves:	✓ (EN166).Clean daily and disinfect at regular intervals in accordance with the instructions of the manufacturer.  No.
Gloves:	
	Gloves resistant against chemicals (FN374) When repeated or prolonged contact with the product is
	expected, gloves of protection level 5 or higher should be used, with a breakthrough time of >240 min. When short contact with the product is expected, use gloves with a protection level 2 or higher should be used, with a breakthrough time >30 min. The breakthrough time of the selected glove material should be in accordance with the pretended period of use. There are several factors (for example, temperature), they do in practice the period of use of a protective gloves resistant against chemicals is clearly lower than the established standard EN374. Due to the wide variety of circumstances and possibilities, the instructions/specifications provided by the glove supplier should be taken into account. Use the proper technique of removing gloves (without touching glove's outer surface) to avoid contact of the product with the skin. The gloves should be immediately replaced when any sign of degradation is noted.
Boots:	No.
Apron:	No.
Clothing:	Advisable.

### - Thermal hazards:

Not applicable (the product is handled at room temperature).

# **ENVIRONMENTAL EXPOSURE CONTROLS:**

Avoid any spillage in the environment. Avoid any release into the atmosphere.

- Spills on the soil:

Prevent contamination of soil.

- Spills in water:

Do not allow to escape into drains, sewers or water courses.

-Water Management Act:

This product contains the following substances included in the list of priority substances in the field of water policy under Directive 2000/60/EC~2013/39/EU:

Diuron (ISO).

- Emissions to the atmosphere:

Because of volatility, emissions to the atmosphere while handling and use may result. Avoid any release into the atmosphere.

#### VOC (product ready for use\*):

It is applicable the Directive 2004/42/EC, on the limitation of emissions of volatile compounds due to the use of organic solvents: PAINTS AND VARNISHES (defined in the Directive 2004/42/EC, Annex I.1): Emission subcategory i) One-pack performance coating, water-borne. VOC (product ready for use\*): (REVESTIDAN FIBER Cod. 72000 = 100 in volume): 0,2 g/l (VOC max.140 g/l\* starting from 01.01.2010)

### VOC (industrial installations):

If this product is used in an industrial installation, it must be verified if it is applicable the Directive 2010/75/CE (DL.127/2013, on the limitation of emissions of volatile compounds due to the use of organic solvents in certain activities and installations: Solvents: 1,00 % Weight, VOC (supply): 0,01 % Weight, VOC: 0,01 % C (expressed as carbon), Molecular weight (average): 196,65, Number C atoms (average): 8,13





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Liquid

### SECTION 9: PHYSICAL AND CHEMICAL PROPERTIES

#### INFORMATION ON BASIC PHYSICAL AND CHEMICAL PROPERTIES:

**Appearance** 

Physical state:

Colour: See the colour in the package

Odour: Characteristic

Odour threshold: Not available (mixture).

Change of state

Melting point: Not available (mixture). Initial boiling point: > 100\* °C at 760 mmHg

- Flammability:

Flashpoint: Not flammable Lower/upper flammability or explosive limits: Not available

Autoignition temperature: Not applicable (do not sustain combustion).

Stability

Decomposition temperature: 825,00\* °C

pH-value

pH: 8,5 ± 0,5 at 20°C

Viscosity:

Dynamic viscosity: Not available. Kinematic viscosity: Not available. Viscosity (Krebs-Stormer): 140 ± 5 KU at 20°C

- Solubility(ies):

Solubility in water Miscible

Liposolubility: Not applicable (inorganic product).

Partition coefficient: n-octanol/water: Not applicable (mixture).

Volatility:

Vapour pressure: 17,535\* mmHg at 20°C Vapour pressure: 12,113\* kPa at 50°C Evaporation rate: Not available (lack of data).

**Density** 

1,300 ± 0,05 at 20/4°C Relative density: Relative water

Relative vapour density: Not available.

Particle characteristics

Particle size: Not applicable.

**Explosive properties:** 

Not available.

Oxidizing properties:

Not classified as oxidizing product.

\*Estimated values based on the substances composing the mixture.

#### **OTHER INFORMATION:** 9.2

Information regarding physical hazard classes

No additional information available.

Other security features:

VOC (supply): 0,2 g/l

Nonvolatile: 61,23 \* % Weight 1h. 60°C

The values indicated do not always coincide with product specifications. The data for the product specifications can be found in the corresponding technical data sheet. For additional information concerning physical and chemical properties related to safety and environment, see sections 7 and 12.





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1010101	
SECTIO	N 10: STABILITY AND REACTIVITY
10.1	REACTIVITY:
	- Corrosivity to metals:
	It is not corrosive to metals.
	- Pyrophorical properties:
	It is not pyrophoric.
10.2	CHEMICAL STABILITY:
	Stable under recommended storage and handling conditions.
10.3	POSSIBILITY OF HAZARDOUS REACTIONS:
	Possible dangerous reaction with reducing agents, oxidizing agents, acids, alkalis.
10.4	CONDITIONS TO AVOID:
	- Heat:
	Keep away from sources of heat.
	<u>- Light:</u>
	If possible, avoid direct contact with sunlight.
	<u>- Air:</u>
	The product is not affected by exposure to air, but should not be left the containers open.
	- Pressure:
	Not relevant.
	- Shock:
	The product is not sensitive to shocks, but as a recommendation of a general nature should be avoided bumps and rough handling to avoid dents and breakage of packaging, especially when the product is handled in large quantities, and during loading and download operations.
10.5	INCOMPATIBLE MATERIALS:
	Keep away from reducing agents, oxidizing agents, acids, alkalis.
10.6	HAZARDOUS DECOMPOSITION PRODUCTS:
	As consequence of thermal decomposition, hazardous products may be produced: nitrogen oxides, sulfur oxides.

### SECTION 11: TOXICOLOGICAL INFORMATION

No experimental toxicological data on the preparation is available. The toxicological classification for these mixture has been carried out by using the conventional calculation method of the Regulation (EU) No. 1272/2008~2021/849 (CLP).

#### INFORMATION ON HAZARD CLASSES AS DEFINED IN REGULATION (EC) NO 1272/2008 : 11.1

### **ACUTE TOXICITY:**

Dose and lethal concentrations for individual ingredients:	DL50 (OECD401) mg/kg bw Oral	` '	
2-methylisothiazol-3(2H)-one	148 Rat	• •	•
2-octyl-2H-isothiazol-3-one	125 Rat	311 Rabbit	> 270 Rat
Estimates of acute toxicity (ATE) for individual ingredients:	ATE mg/kg bw Oral	ATE mg/kg bw Cutaneous	· · · –
Estimates of acute toxicity (ATE) for individual ingredients: 2-methylisothiazol-3(2H)-one	ATE mg/kg bw Oral 148	mg/kg bw Cutaneous	mg/m3·4h Inhalation

- (\*) Point estimates of acute toxicity corresponding to the classification category (see GHS/CLP Table 3.1.2). These values are designed to be used in the calculation of the ATE for classification of a mixture based on its components and do not represent test results.
- (-) The components that are assumed to have no acute toxicity at the upper threshold of category 4 for the corresponding exposure route are ignored.

# - No observed adverse effect level

Not available

## - Lowest observed adverse effect level

Not available

### INFORMATION ON LIKELY ROUTES OF EXPOSURE: ACUTE TOXICITY:

Routes of exposure	Acute toxicity	Cat.	Main effects, acute and/or delayed	Criteria
Inhalation: Not classified	ATE > 20000 mg/m3	-	Not classified as a product with acute toxicity if inhaled (based on available data, the classification criteria are not met).	GHS/CLP 3.1.3.6.
Skin: Not classified	ATE > 5000 mg/kg bw	-	Not classified as a product with acute toxicity in contact with skin (based on available data, the classification criteria are not met).	
Eyes: Not classified	Not available.	-	Not classified as a product with acute toxicity by eye contact (lack of data).	GHS/CLP 1.2.5.
Ingestion: Not classified	ATE > 5000 mg/kg bw	-	Not classified as a product with acute toxicity if swallowed (based on available data, the classification criteria are not met).	GHS/CLP 3.1.3.6.

GHS/CLP 3.1.3.6: Classification of mixtures based on ingredients of the mixture (additivity formula).





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### CORROSION / IRRITATION / SENSITISATION :

Danger class	Target organs	Cat.	Main effects, acute and/or delayed	Criteria
<ul> <li>Respiratory corrosion/irritation</li> <li>Not classified</li> </ul>	on: -	-	Not classified as a product corrosive or irritant by inhalation (based on available data the classification criteria are not met).	GHS/CLP ,1.2.6. 3.8.3.4.
- Skin corrosion/irritation: Not classified	-	-	Not classified as a product corrosive or irritant in contact with skin (based on available data, the classification criteria are not met).	GHS/CLP 3.2.3.3.
- Serious eye damage/irritatio Not classified	n: -		Not classified as a product corrosive or irritant in contact with eyes (based on available data, the classification criteria are not met).	GHS/CLP 3.3.3.3.
- Respiratory sensitisation: Not classified	-	-	Not classified as a product sensitising by inhalation (based on available data, the classification criteria are not met).	GHS/CLP 3.4.3.3.
- Skin sensitisation:	Skin	Cat.1	SENSITISING: May cause an allergic skin reaction.	GHS/CLP 3.4.3.3.

GHS/CLP 3.2.3.3: Classification of the mixture when data are available for all components or only for some components. GHS/CLP 3.3.3.3: Classification of the mixture when data are available for all components or only for some components. GHS/CLP 3.4.3.3: Classification of the mixture when data are available for all components or only for some components. GHS/CLP 3.8.3.4: Classification of the mixture when data are available for all components or only for some components.

## - ASPIRATION HAZARD:

Danger class	Target organs	Cat.	Main effects, acute and/or delayed	Criteria
- Aspiration hazard: Not classified	_		,	GHS/CLP 3.10.3.3.

GHS/CLP 3.10.3.3: Classification of the mixture when data are available for all components or only for some components.

### SPECIFIC TARGET ORGANS TOXICITY (STOT): Single exposure (SE) and/or Repeated exposure (RE):

Not classified as a dangerous product for target organs.

GHS/CLP 3.8.3.4: Classification of the mixture when data are available for all components or only for some components.

#### CMR EFFECTS:

- Carcinogenic effects:

It is not considered as a carcinogenic product.

Genotoxicity:

It is not considered as a mutagenic product.

Toxicity for reproduction:

Does not harm fertility. Does not harm the unborn child.

Effects via lactation:

Not classified as a hazardous product for children breast-fed.

#### DELAYED AND IMMEDIATE EFFECTS AS WELL AS CHRONIC EFFECTS FROM SHORT AND LONG-TERM EXPOSURE:

Routes of exposure

Not available.

- Short-term exposure:

Not available.

- Long-term or repeated exposure:

Not available.

## **INTERACTIVE EFFECTS:**

Not available.

# INFORMATION ABOUT TOXICOCINETICS, METABOLISM AND DISTRIBUTION:

Dermal absorption:

Not available.

Basic toxicokinetics:

Not available.

# **ADDITIONAL INFORMATION:**

Not available.





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### 11.2 <u>INFORMATION ON OTHER HAZARDS:</u>

**Endocrine disrupting properties:** 

This product contains substances with endocrine disrupting properties identified or under evaluation in a concentration of less than 0.1% by weight:Diuron (ISO).

Other information:

No additional information available.

### SECTION 12: ECOLOGICAL INFORMATION

No experimental ecotoxicological data on the preparation as such is available. The ecotoxicological classification for these mixture has been carried out by using the conventional calculation method of the Regulation (EU) No. 1272/2008~2021/849 (CLP).

#### 12.1 <u>TOXICITY:</u>

- Acute toxicity in aquatic environment for individual ingredients	CL50 (OECD 203) mg/l·96hours	( )	CE50 (OECD 201) mg/l·72hours
2-methylisothiazol-3(2H)-one	4.8 - Fishes	0.93 - Daphniae	0.072 - Algae
2-octyl-2H-isothiazol-3-one	0.12 - Fishes	0.18 - Daphniae	0.15 - Algae

- No observed effect concentration	NOEC (OECD 210)	( /	NOEC (OECD 201) mg/l · 72 hours
2-methylisothiazol-3(2H)-one	4.9 - Fishes	0.044 - Daphniae	0.038 - Algae
2-octyl-2H-isothiazol-3-one	0.022 - Fishes	0.035 - Daphniae	0.068 - Algae

### - Lowest observed effect concentration

Not available

## **ASSESSMENT OF AQUATIC TOXICITY:**

Aquatic toxicity	Cat.	Main hazards to the aquatic environment	Criteria
- Acute aquatic toxicity: Not classified	-	Not classified as a hazardous product with acute toxicity to aquatic life (based on available data, the classification criteria are not met).	GHS/CLP 4.1.3.5.5.3.
- Chronic aquatic toxicity:	Cat.3	HARMFUL: Harmful to aquatic life with long lasting effects.	GHS/CLP 4.1.3.5.5.4.

CLP 4.1.3.5.5.3: Classification of a mixture for acute hazards, based on summation of classified components.

CLP 4.1.3.5.5.4: Classification of a mixture for chronic (long term) hazards, based on summation of classified components.

### 12.2 PERSISTENCE AND DEGRADABILITY:

### - Biodegradability:

Not available.

Aerobic biodegradation for individual ingredients	COD mgO2/g	%DBO/DQO 5 days 14 days 28 days	Biodegradabilidad
2-methylisothiazol-3(2H)-one		54	Not easy
2-octyl-2H-isothiazol-3-one			Not easy

Note: Biodegradability data correspond to an average of data from various bibliographic sources.

### - Hydrolysis:

Not available.

#### - Photodegradability:

Not available.

## 12.3 BIOACCUMULATIVE POTENTIAL:

Not available.

Bioaccumulation for individual ingredients	logPow	BCF L/kg	Potential
2-methylisothiazol-3(2H)-one	-0.48	3.2 (calculated)	No bioaccumulable
2-octyl-2H-isothiazol-3-one	2.61	19.2 (calculated)	Low

# 12.4 MOBILITY IN SOIL:

Not available

Mobility for individual ingredients	log Poc	Constant of Henry Pa·m3/mol 20°C	Potential
2-methylisothiazol-3(2H)-one	0,44		No bioaccumulable
2-octyl-2H-isothiazol-3-one	2,26	0,036 (calculated)	Low

12.5 RESULTS OF PBT AND VPVB ASSESMENT: (Annex XIII of Regulation (EC) no. 1907/2006:)

Does not contain substances that fulfil the PBT/vPvB criteria.

# 12.6 ENDOCRINE DISRUPTING PROPERTIES:

This product contains substances with endocrine disrupting properties identified or under evaluation in a concentration of less than 0.1% by weight:Diuron (ISO).

12.7 OTHER ADVERSE EFFECTS:





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- Ozone depletion potential:

Not available.

- Photochemical ozone creation potential:

Not available.

- Earth global warming potential:

Not available.

#### SECTION 13: DISPOSAL CONSIDERATIONS

#### WASTE TREATMENT METHODS:Directive 2008/98/EC~Regulation (EU) no. 1357/2014: 13.1

Take all necessary measures to prevent the production of waste whenever possible. Analyse possible methods for revaluation or recycling. Do not discharge into drains or the environment, dispose at an authorised waste collection point. Waste should be handled and disposed in accordance with current local and national regulations. For exposure controls and personal protection measures, see section 8.

Disposal of empty containers:Directive 94/62/EC~2015/720/EU, Decision 2000/532/EC~2014/955/EU:

Emptied containers and packaging should be disposed in accordance with currently local and national regulations. The classification of packaging as hazardous waste will depend on the degree of empting of the same, being the holder of the residue responsible for their classification, in accordance with Chapter 15 01 of Decision 2000/532/EC, and forwarding to the appropriate final destination. With contaminated containers and packaging, adopt the same measures as for the product in itself.

Procedures for neutralising or destroying the product:

Authorised landfill in accordance with local regulations.

OFOTION	A TRANSPORT INFORMATION
	N 14: TRANSPORT INFORMATION
14.1	UN NUMBER OR ID NUMBER:
	Not applicable
14.2	<u>UN PROPER SHIPPING NAME:</u>
	Not applicable
14.3	TRANSPORT HAZARD CLASS(ES):
	Transport by road (ADR 2021) and
	Transport by rail (RID 2021):
	No reglamented
	Transport by sea (IMDG 39-18):
	No reglamented
	Transport by air (ICAO/IATA 2021):
	No reglamented
	Transport by inland waterways (ADN):
	No reglamented
14.4	PACKING GROUP:
	No reglamented
14.5	ENVIRONMENTAL HAZARDS:
	Not applicable.
14.6	SPECIAL PRECAUTIONS FOR USER:
	Ensure that persons transporting the product know what to do in case of accident or spill. Always transport in closed containers that are
	upright and secure.
14.7	MARITIME TRANSPORT IN BULK ACCORDING TO IMO INSTRUMENTS:
	Not applicable.





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#### SECTION 15: REGULATORY INFORMATION

#### 15.1 SAFETY, HEALTH AND ENVIRONMENTAL REGULATIONS/LEGISLATION SPECIFIC FOR THE SUBSTANCE OR MIXTURE

The regulations applicable to this product generally are listed throughout this Safety Data Sheet.

Restrictions on manufacture, placing on market and use:

See section 1.2

Tactile warning of danger:

Not applicable (the classification criteria are not met).

Child safety protection:

Not applicable (the classification criteria are not met).

VOC information on the label:

Contains VOC max. 0,2 g/l for the product ready for use - The limit value 2004/42/EC-IIA cat. i) One-pack performance coating, water-

borne. is VOC max. 140 g/l (2010)

OTHER REGULATIONS:

Control of the risks inherent in major accidents (Seveso III):

See section 7.2

Other local legislations:

The receiver should verify the possible existence of local regulations applicable to the chemical

15.2 CHEMICAL SAFETY ASSESSMENT:

A chemical safety assessment has not been carried out for this mixture.

### SECTION 16: OTHER INFORMATION

#### 16.1 TEXT OF THE PHRASES AND NOTES REFERENCED IN SECTIONS 2 AND/OR 3:

#### Hazard statements according the Regulation (EU) No. 1272/2008~2021/849 (CLP), Annex III:

H301 Toxic if swallowed. H311 Toxic in contact with skin. H314 Causes severe skin burns and eye damage. H317 May cause an allergic skin reaction. H318 Causes serious eye damage. H330 Fatal if inhaled. H400 Very toxic to aquatic life. H410 Very toxic to aquatic life with long lasting effects. H412 Harmful to aquatic life with long lasting effects. EUH071 Corrosive to the respiratory tract.

**EVALUATION OF THE INFORMATION ON THE DANGER OF MIXTURES:** 

See sections 9.1, 11.1 and 12.1.

#### ADVICES ON ANY TRAINING APPROPRIATE FOR WORKERS:

It is recommended for all staff that will handle this product to carry out a basic training in occupational risk and prevention, in order to provide understanding and interpretation of Safety Data Sheets and labelling of products as well.

# MAIN LITERATURE REFERENCES AND SOURCES FOR DATA:

- · European Chemicals Agency: ECHA, http://echa.europa.eu/
- Access to European Union Law, http://eur-lex.europa.eu/
- · European agreement on the international carriage of dangerous goods by road, (ADR 2021).
- · International Maritime Dangerous Goods Code IMDG including Amendment 39-18 (IMO, 2018).

# ABBREVIATIONS AND ACRONYMS:

List of abbreviations and acronyms that can be used (but not necessarily used) in this Safety Data Sheet:

- · REACH: Regulation concerning the Registration, Evaluation, Authorisation and Restriction of Chemicals.
- GHS: Globally Harmonized System of Classification and Labelling of Chemicals of the United Nations.
- CLP: European regularion on Classificatin, Labelling amd Packaging of substances and chemical mixtures.
- · EINECS: European Inventory of Existing Commercial Chemical Substances.
- · ELINCS: European List of Notified Chemical Substances.
- · CAS: Chemical Abstracts Service (Division of the American Chemical Society).
- · UVCB: Substances of Unknown or Variable composition, complex reaction products or biological materials.
- SVHC: Substances of Very High Concern.
- · PBT: Persistent, bioaccumulable and toxic substances.
- · vPvB: Very persistent and very bioaccumulable substances.
- · VOC: Volatile Organic Compounds.
- · DNEL: Derived No-Effect Level (REACH).
- · PNEC: Predicted No-Effect Concentration (REACH).
- · LC50: Lethal concentration, 50 percent.
- · LD50: Lethal dose, 50 percent.
- · UN: United Nations Organisation.
- · ADR: European agreement concerning the international carriage of dangeous goods by road.
- · RID: Regulations concerning the international transport of dangeous goods by rail.
- · IMDG: International Maritime code for Dangerous Goods.
- · IATA: International Air Transport Association.
- · ICAO: International Civil Aviation Organization.

### SAFETY DATA SHEET REGULATIONS:

Safety Data Sheet in accordance with Article 31 of Regulation (EC) No. 1907/2006 (REACH) and Annex of Regulation (EU) No. 2020/878.

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 REVISION:

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# Changes since previous Safety Data Sheet:

Changes that have been introduced with respect to the previous version due to the structural and content adaptation of the Safety Data Sheet to Regulation (EU) No. 2020/878: All sections.





Version: 5 Revision: 08/03/2023 Previous revision: 22/04/2022 Date of printing: 08/03/2023 The information of this Safety Data Sheet, is based on the present state of knowledge and on current UE and national laws, as the users" working conditions are beyond our knowledge and control. The product is not to be used for other purposes than those specified, without first obtaining written handling instruction. It is always the responsibility of the user to take all necessary steps in order to fulfil the demand laid down in the local rules and legislation. The information in this Safety Data Sheet is meant as a description of the safety requirements of the product and it is not to be considered as a guarantee of the product"s properties.