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

- 1.1 Product identifier

- Trade name: RODIBROD BLOCK

- Sds code/version: 0/19

- 1.2 Relevant identified uses of the substance or mixture and uses advised against Ready for use rodenticide (biocidal product-PT14)

- Application of the substance / the mixture Ready for use rodenticide (biocidal product-PT14)

- 1.3 Details of the supplier of the safety data sheet

- Manufacturer/Supplier:

Zapi S.p.A. Via Terza Strada, 12 35026 Conselve (Pd)

Italy

Tel. +39 049 9597737 Fax +39 049 9597735

E-mail address of the competent person responsible for the SDS: techdept@zapi.it

- Further information obtainable from: Tech. dept.
- **1.4 Emergency telephone number:** Zapi customer service: tel. +39 049 9597737 (9:00-12:00/14:00-17:00) National Poisons Information Centre (Ireland), Beaumont Hospital, Dublin: 01-809 2166

SECTION 2: Hazards identification

- 2.1 Classification of the substance or mixture
- Classification according to Regulation (EC) No 1272/2008

STOT RE 2 H373 May cause damage to organs (blood) through prolonged or repeated exposure.

- 2.2 Label elements
- Labelling according to Regulation (EC) No 1272/2008

The product is classified and labelled according to the CLP regulation.

- Hazard pictograms



GHS08

- Signal word Warning
- Hazard-determining components of labelling:

brodifacoum

- Hazard statements

H373 May cause damage to the blood through prolonged or repeated exposure.

- Precautionary statements

P102 Keep out of reach of children.

P301+P310 IF SWALLOWED: Immediately call a POISON CENTER/doctor.

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

P501 Dispose of contents/container in accordance with local regulation.

- 2.3 Other hazards

- Results of PBT and vPvB assessment

- PBT:

56073-10-0 brodifacoum

PBT Brodifacoum fulfils the P, B and T criteria.

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	(coa. c. page .)
- vPvB:	
56073-10-0 brodifacoum	
vPvB Brodifacoum fulfils the vP criterion.	

SECTION 3: Composition/information on ingredients

- 3.2 Mixtures

- **Description:** Mixture of substances listed below with nonhazardous additions.

- Dangerous components:			
CAS: 52-51-7 EINECS: 200-143-0 Index number: 603-085-00-8	bronopol (INN) Eye Dam. 1, H318; Aquatic Acute 1, H400 (M=10); Aquatic Chronic 1, H410 (M=1); Acute Tox. 4, H302; Acute Tox. 4, H312; Skin Irrit. 2, H315; STOT SE 3, H335	_ ≤1%	
CAS: 56073-10-0 EINECS: 259-980-5 Index number: 607-172-00-1	brodifacoum Acute Tox. 1, H300; Acute Tox. 1, H310; Acute Tox. 1, H330; Repr. 1A, H360D; STOT RE 1, H372; Aquatic Acute 1, H400 (M=10); Aquatic Chronic 1, H410 (M=10)	≤1%	
CAS: 107-21-1 EINECS: 203-473-3 Index number: 603-027-00-1	ethanediol STOT RE 2, H373; Acute Tox. 4, H302	≤1%	

- Additional information: For the wording of the listed hazard phrases refer to section 16.

SECTION 4: First aid measures

- 4.1 Description of first aid measures

- General information: Please refer to the instructions below for each specific way of exposure.
- After inhalation: Supply fresh air and to be sure call for a doctor.

After skin contact:

Remove contaminated clothing.

Wash skin with water and then with water and soap.

If needed, seek for medical advice.

- After eye contact:

Rinse eyes with eye-rinse liquid or water, keep eyelids open at least 10 minutes.

If needed, seek medical advice.

- After swallowing:

Rinse mouth carefully with water. Never give anything by mouth to unconscious person. Do not provoke vomiting. If swallowed, seek medical advice immediately and show the product's container or label.

Contact a veterinary surgeon in case of ingestion by a pet.

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

This product contains an anticoagulant substance. If ingested, symptoms, which may be delayed, may include nosebleed and bleeding gums. In severe cases, there may be bruising and blood present in the faeces or urine. Antidote: Vitamin K1 administered by medical/veterinary personnel only.

4.3 Indication of any immediate medical attention and special treatment needed

The primary treatments are the antidote therapy and the clinical assessment. Antidote: Vitamin K1 (phytomenadione). The effectiveness of the treatment should be monitored by measuring the clotting time. Do not interrupt the treatment until the clotting time is back to normality and is stable. Consult a Poison Control Centre.

SECTION 5: Firefighting measures

- 5.1 Extinguishing media

- Suitable extinguishing agents: CO₂, powder or water spray. Fight larger fires with water spray.

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- For safety reasons unsuitable extinguishing agents: To our knowledge, there are no unsuitable equipment.
- 5.2 Special hazards arising from the substance or mixture In case of fire, toxic gases may be generated.
- 5.3 Advice for firefighters Fire-fighters equipment in accordance with EN469 European standards.

- Protective equipment:

Do not inhale explosion gases or combustion gases.

Firefighters equipment in accordance with EN 469 European standards.

- Additional information

Dispose of fire debris and contaminated fire-fighting water in accordance with official regulations.

SECTION 6: Accidental release measures

- 6.1 Personal precautions, protective equipment and emergency procedures

Wear protective equipment. Keep unprotected persons away.

- 6.2 Environmental precautions:

Inform respective authorities in case of seepage into water course or sewage system.

Do not allow to enter sewers/ surface or ground water.

- 6.3 Methods and material for containment and cleaning up:

Pick up mechanically.

After cleaning up, ensure adequate ventilation.

Dispose of the material collected according to regulations.

- 6.4 Reference to other sections

See Section 7 for information on safe handling.

See Section 8 for information on personal protection equipment.

See Section 13 for disposal information.

SECTION 7: Handling and storage

- 7.1 Precautions for safe handling

Wash hands and directly exposed skin after using the product.

Wear appropriate protective gloves.

When using the product, do not eat, drink or smoke.

- Information about fire - and explosion protection:

See Section 6.

See section 5.

-7.2 Conditions for safe storage, including any incompatibilities

- Requirements to be met by storerooms and receptacles:

Store in a dry, cool and well ventilated place. Keep the container closed and away from direct sunlight.

Store in places prevented from the access of children, birds, pets and farm animals.

- Information about storage in one common storage facility:

Place the product away from food, drink and animal feeding stuffs, as well as from utensils or surfaces that have contact with these.

- Further information about storage conditions:

Protect from frost.

Protect from humidity and water.

-7.3 Specific end use(s) This product is a rodenticide bait for rodents' control.

SECTION 8: Exposure controls/personal protection

- Additional information about design of technical facilities: No further data; see item 7.

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8.1 Control parameters (Contd. of page 3)

(Contact Parameters			
- Ingredients with limit values that require monitoring at the workplace:			
107-21-1	ethanediol		
OEL (EU)	Short-term value: 104 mg/m³, 40 ppm Long-term value: 52 mg/m³, 20 ppm Skin		
OEL (IRA)	8-hour reference period: Particulate: 10 mg/m³ Vapour: 20 mg/m³ 15-minute reference period: Vapour: 52 ppm; 104 mg/m³ Notes: Sk, iOELV		
102-71-6 1	102-71-6 triethanolamine		
OEL (IRA)	8-hour reference period: 5 mg/m ³		
57-55-6 propylene glycol			
OEL (IRA)	8-hour reference period: Total (vapour and particulates): 150 ppm; 470 mg/m³ Particulates: 10 mg/m³		

- Regulatory information

OEL (EU): Directives 98/24/EC, 2000/39/EC, 2004/37/EC, 2006/15/EC, 2009/161/EU, 2017/164/EU.

OEL (IRA): 2018 Code of Practice for the Chemical Agents Regulations

	` '	2010 Code of Fractice for the	- 3
- DNE	ELs		
107-	21-1 et	hanediol	
Dern	nal L	ong term - systemic effects.	53 mg/kg bw/d (general population)
			106 mg/kg bw/d (workers)
Inhal	lative L	ong term - local effects	7 mg/m3 (general population)
			35 mg/m3 (workers)
- PNE	Cs		
5607	73-10-0	brodifacoum	
Oral	PNEC	1.28-5 mg/kg bw (bird)	
		1.1-5 mg/kg bw (mammal)	
	PNEC	0.00004 mg/l (aquatic orga	anisms)
		>0.0038 mg/l (microorgani	sms)
	PNEC	>0.88 mg/kg ww (soil)	
107-	21-1 et	hanediol	
	PNEC	10 mg/l (fresh water)	
		10 mg/l (intermittent releases)	
		1 mg/l (marine water) 199.5 mg/l (sewage treatment plant)	
	PNEC 37 mg/kg dw (sediment (fresh water)) 3.7 mg/kg dw (sediment (marine water))		esh water))
	1.53 mg/kg dw (soil)		
- Oth	er exp	osure limit values	
5607	73-10-0	brodifacoum	
Oral	AEL -	short term 3.3-6 mg/kg l	bw (AEL)
	AEL - medium term 6-6.67 mg/		bw (AEL)
	AEL -	long term 3.3-6 mg/kg l	bw (AEL)

- 8.2 Exposure controls

- Personal protective equipment:

- General protective and hygienic measures:

The usual precautionary measures are to be adhered to when handling chemicals.

Keep away from food, drink and animal feed. Wash hands before breaks and at the end of work.

Do not eat, drink, smoke or sniff while working.

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Respiratory protection: Not required during normal use of the product. (Contd. of page 4)

- Protection of hands:



Professional use: wear suitable gloves (EN 374, category III) when handling the product.

The glove material has to be impermeable and resistant to the product/ the substance/ the preparation. Due to missing tests no recommendation to the glove material can be given for the product/ the preparation/ the chemical mixture.

Selection of the glove material on consideration of the penetration times, rates of diffusion and the degradation.

- Material of gloves

The selection of the suitable gloves does not only depend on the material, but also on further marks of quality and varies from manufacturer to manufacturer. As the product is a preparation of several substances, the resistance of the glove material cannot be calculated in advance and has therefore to be checked prior to the application.

- Penetration time of glove material

The exact break-through time has to be found out by the manufacturer of the protective gloves and has to be observed.

- Eye protection: Not required during normal use of the product.
- Limitation and supervision of exposure into the environment See section 6.
- Risk management measures Follow the above-reported directions.

SECTION 9: Physical and chemical pr	SECTION 9: Physical and chemical properties		
- 9.1 Information on basic physical and	l chemical properties		
- General Information			
- Appearance:			
Form:	Solid		
Colour:	Blue		
- Odour:	Characteristic		
- Odour threshold:	No data available.		
- pH-value:	7.2		
- Change in condition			
Melting point/freezing point:	No data available.		
Initial boiling point and boiling rang	e: Not applicable (solid).		
- Flash point:	Not applicable (solid).		
- Flammability (solid, gas):	Not available (the product does not contain any ingredient classified as flammable).		
- Ignition temperature:	No data available.		
- Decomposition temperature:	No data available.		
- Auto-ignition temperature:	Product is not selfigniting.		
- Explosive properties:	Product does not present an explosion hazard.		
- Explosion limits:			
Lower:	No data available.		
Upper:	No data available.		
- Oxidising properties	No data available.		
- Vapour pressure:	Not applicable.		
- Density:	No data available.		

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	(Conta. or page 5)
- Relative density	1.09 g/ml
- Vapour density	Not applicable.
- Evaporation rate	Not applicable.
- Solubility in / Miscibility with	
water:	Insoluble.
- Partition coefficient: n-octanol/water:	No data available.
- Viscosity:	
Dynamic:	Not applicable.
Kinematic:	Not applicable.
- 9.2 Other information	No further relevant information available.

SECTION 10: Stability and reactivity

- 10.1 Reactivity Under standard handling and storing conditions, the product does not show any dangerous reaction.
- 10.2 Chemical stability Stable at room temperature and if used as recommended.
- Thermal decomposition / conditions to be avoided: No decomposition if used according to specifications.
- 10.3 Possibility of hazardous reactions No dangerous reactions known.

- 10.4 Conditions to avoid

Under standard handling and storing conditions, the product does not show any dangerous reaction.

- 10.5 Incompatible materials:

Store only in original container.

Given the lack of information about possible incompatibilities with other substances, it is recommended not to use it in combination with other products.

- 10.6 Hazardous decomposition products:

No dangerous decomposition products known under normal conditions of storage and use.

SECTION 11: Toxicological information

- 11.1 Information on toxicological effects

- Acute toxicity Based on available data, the classification criteria are not met.

Product texticity bacod on available data, the diacombation officina are not mot.			
- LD/LC50	- LD/LC50 values relevant for classification:		
52-51-7 br	52-51-7 bronopol (INN)		
Oral	LD50	305 mg/kg bw (rat) (OECD 401)	
Dermal	LD50	>2,000 mg/kg bw (rat) (OECD 402)	
Inhalative	LC50/4h	>0.588 mg/l (rat)	
56073-10-	56073-10-0 brodifacoum		
Oral	LD50	0.4 mg/kg bw (male rat and mouse)	
Dermal	LD50	3.16 mg/kg bw (rat)	
Inhalative	LC50/4h	3.05 mg/m3 (rat)	
107-21-1 €	107-21-1 ethanediol		
Oral	LD50	7,712 mg/kg bw (rat)	
Dermal	LD50	>3,500 mg/kg bw (mouse)	
Inhalative	LC50/6h	>2.5 mg/l (rat)	

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- Primary irritant effect: (Contd. of page 6)

- Skin corrosion/irritation Based on available data, the classification criteria are not met.

- Serious eye damage/irritation

52-51-7 bronopol (INN)

eye irritation | Corrosive (rabbit; OECD 405).

Based on available data, the classification criteria are not met.

- Respiratory or skin sensitisation

52-51-7 bronopol (INN)

skin sensitisation Not sensitising (guinea pig; OECD 406).

Based on available data, the classification criteria are not met.

- CMR effects (carcinogenity, mutagenicity and toxicity for reproduction)
- Germ cell mutagenicity Based on available data, the classification criteria are not met.
- Carcinogenicity Based on available data, the classification criteria are not met.

- Reproductive toxicity

56073-10-0 brodifacoum

developmental toxicity | Clear developmental toxicity was not observed in rabbits or rats. However, as a precaution, Brodifacoum should be considered teratogenic to humans because it contains the same chemical moiety responsible for the teratogenicity of warfarin, a known human teratogenic agent, and it has the same mode of action that is a known mechanism of teratogenicity in humans.

Based on available data, the classification criteria are not met.

- STOT-single exposure Based on available data, the classification criteria are not met.

- STOT-repeated exposure

56073-10-0 brodifacoum

Oral NOAEL 0.04 mg/kg bw/d (rat)

The study reveals that repeated oral exposure results in toxic effects: prothrombin time prolongation, kaolin-caphalin time prolongation, haemorrhage.

Based on the results of the acute dermal and inhalation toxicity studies and route-to-route extrapolation, it is justified to assume a similar concern for serious damage to health by prolonged exposure through dermal and inhalation routes also.

May cause damage to the blood through prolonged or repeated exposure.

- Aspiration hazard Based on available data, the classification criteria are not met.

SECTION 12: Ecological information

- 12.1 Toxicity

- Aquatic and/or terrestrial toxicity:		
52-51-7 bronopol (INN)		
EC50/72h	0.068 mg/l (Anabaena flos aqua) (OECD 201)	
LC50/96h (dynamic)	3 mg/l (Oncorhynchus mykiss) (OECD 203)	
NOEC/21d	0.06 mg/l (Daphnia magna) (OECD 211)	
NOEC/72h	0.0025 mg/l (Anabaena flos aqua) (OECD 201)	
NOEC/28d	2.61 mg/l (Oncorhynchus mykiss) (OECD 210)	
EC50/48h (static)	1.04 mg/l (Daphnia magna) (OECD 202)	
56073-10-0 brodifacoum		
LC50/14d	(eisenia foetida) >994 mg/kg dry weight >879.6 mg/kg wet weight	
ErC50/72h	0.04 mg/l (Selenastrum capricornutum)	
EC10/3h	>0.058 mg/l (activated sludge) Based on water solubility at pH 7 and T=20°C.	
EC10/6h	>0.0038 mg/l (Pseudomonas putida) Based on water solubility at pH 5.2 and T=20°C.	
LC50/96h	0.042 mg/l (Oncorhynchus mykiss)	(Contd. on page 8)

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LC50 (diet) 0.72		mg/kg food (laughing gull)	(Contd. on page 7)	
NOEC (reproductive toxicity) 0.00		0038 mg/kg food (bird)		
NOEL (reproductive toxicity) 0.000		00385 mg/kg bw/d (bird)		
		mg/kg bw (mallard duck)		
EC50/48h	0.25	mg/l (Daphnia magna)		
107-21-1 ethaned				
EC50/96h	6,50	0-13,000 mg/l (Selenastrum capricornutum)		
LC50/96h	72,8	60 mg/l (Pimephales promelas)	0 mg/l (Pimephales promelas)	
NOEC/7d		80 mg/l (Pimephales promelas)		
EC50/48h	>10	0 mg/l (Daphnia magna)		
- 12.2 Persistenc	_	ability		
52-51-7 bronopol				
biodegradability	CO2 Evolution:	>70% (activated sludge; OECD 301 B).		
	Readily biodeg	s: 63.5% (OECD 314).		
56073-10-0 brodi		adable.		
biodegradability	Not easily biod	egradable		
blodogradability		II probably partition into sewage sludge/sediment due to	its high log Kow and poor	
photolytic half-life	0.083 days. De	grades rapidly by photolysis.		
Hydrolitic half-life	Hydrolitic half-life > 1 year. Stable at pH 5, 7 and 9.			
- 12.3 Bioaccumi	ulative potent	ial		
52-51-7 bronopol	(INN)			
bioconcentration fa	actor	BCF = 3.16 (calculated; EPIWIN).		
		It does not accumulate in organisms.		
		Log Kow = 0.38 (OECD 107)		
56073-10-0 brodi				
E		BCF fish = 35645 (calculated according to TGD eq. 75, t BCF earthworm = 15820 (calculated according to TGD 6.12).	using log Kow = 6.12). ed. 82d, using log Kow =	
octanol-water part	ition coefficient	log Kow = 6.12 (estimated from measured Koc).		
- 12.4 Mobility in				
56073-10-0 brodifacoum				
DT50		157 days.		
		Persistent.		
organic carbon partition coefficient		Koc = 9155 l/kg (pH 7,1-7.6). Immobile in soil.		
soil mobility		Under basic conditions (high pH), Brodifacoum is not	likely to be adsorbed onto	
Soil Mobility		soils or sewage sludge due to the ionisation of th conditions (low pH), Brodifacoum is likely to be adso sludge as the molecule is in its neutral or non-ionised for	e molecule. Under acidic rbed onto soils or sewage	

- **General notes:** Do not allow the product to reach ground water, water course or sewage system.

- 12.5 Results of PBT and vPvB assessment

12.0 1.004.1.0 0.1 2.1 4.1.4 1. 12 4.0000011011.
- PBT:
56073-10-0 brodifacoum
PBT Brodifacoum fulfils the P, B and T criteria.
- vPvB:
56073-10-0 brodifacoum
vPvB Brodifacoum fulfils the vP criterion.
- 12.6 Other adverse effects
56073-10-0 brodifacoum
. The major environmental concern of Brodifacoum is primary and secondary poisoning of non-target animals.

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SECTION 13: Disposal considerations

- 13.1 Waste treatment methods

- Recommendation

Must not be disposed together with household garbage. Do not allow product to reach sewage system. At the end of the treatment, dispose of uneaten bait and the packaging in accordance with EPA requirements for the disposal of hazardous waste. Use of gloves is recommended.

- Uncleaned packaging:
- **Recommendation:** Dispose of in accordance with local requirements.

SECTION 14: Transport information		
- 14.1 UN-Number		
- ADR, ADN, IMDG, IATA	Not applicable	
- 14.2 UN proper shipping name		
- ADR, ADN, IMDG, IATA	Not applicable	
- 14.3 Transport hazard class(es)		
- ADR, ADN, IMDG, IATA		
- Class	Not applicable	
- 14.4 Packing group		
- ADR, IMDG, IATA	Not applicable	
- 14.5 Environmental hazards:	Not applicable.	
- 14.6 Special precautions for user	Not applicable.	
- 14.7 Transport in bulk according to Annex II of		
MARPOL and the IBC Code	Not applicable.	
- UN "Model Regulation":	Not applicable	

SECTION 15: Regulatory information

- 15.1 Safety, health and environmental regulations/legislation specific for the substance or mixture
- Directive 2012/18/EU
- Named dangerous substances ANNEX I None of the ingredients is listed.
- Seveso category This product is not subject to Seveso directive dispositions.
- LIST OF SUBSTANCES SUBJECT TO AUTHORISATION (ANNEX XIV)

The product does not contain any substance included in annex XIV.

- REGULATION (EC) No 1907/2006 ANNEX XVII Restrictions: 30
- National regulations:
- Other regulations, limitations and prohibitive regulations No further information available.
- Holder: Zapi S.p.A. Via Terza Strada, 12 35026 Conselve (Pd) Italia Phone: +39 049 9597737
- N° Authorisation: IE/BPA 70744-005 (for non professional and professional users):

The Competent Authority for Biocides in Ireland, pursuant to the provisions of Regulation (EU) No. 528/2012 of the European Parliament and of the Council concerning the making available on the market and use of biocidal products, as amended by Regulation (EU) No. 334/2014, and European Union (Biocidal Products) Regulations, 2013, grants authorization to make this product available on the market in Ireland.

- Substances of very high concern (SVHC) according to REACH, Article 57 None.
- Regultion (EC) n. 1005/2009: substances that deplete the ozone layer None.
- Regulation (EC) n. 850/2004: persistent organic pollutants None.

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- Substances listed in Regulation (EC) n. 649/2012 (PIC): None.
- -15.2 Chemical safety assessment: A Chemical Safety Assessment has not been carried out for this mixture.

SECTION 16: Other information

This information is based on our present knowledge. However, this shall not constitute a guarantee for any specific product features and shall not establish a legally valid contractual relationship. Any responsibility derived from misuse of the product or in case of violation of current regulations is refused.

- Relevant phrases

H300 Fatal if swallowed.

H302 Harmful if swallowed.

H310 Fatal in contact with skin.

H312 Harmful in contact with skin.

H315 Causes skin irritation.

H318 Causes serious eye damage.

H330 Fatal if inhaled.

H335 May cause respiratory irritation.

H360D May damage the unborn child.

H372 Causes damage to organs through prolonged or repeated exposure.

H373 May cause damage to the blood through prolonged or repeated exposure.

H400 Very toxic to aquatic life.

H410 Very toxic to aquatic life with long lasting effects.

- Classification according to Regulation (EC) No 1272/2008

The classification of the mixture is based on the calculation method stated in annex I of Regulation (CE) n. 1272/2008, using components data.

- Abbreviations and acronyms:

RD50: Respiratory Decrease, 50 percent

LC0: Lethal concentration, 0 percent

NOEC: No Observed Effect Concentration IC50: Inhibitory concentration, 50 percent

NOAEL: No Observed Adverse Effect Level

EC50: Effective concentration, 50 percent

EC10: Effective concentration, 10 percent

AEC: Acceptable Exposure Concentration LL0: Lethal Load, 0 percent

AEL: Acceptable Exposure Limit LL50: Lethal Load, 50 percent

EL0: Effective Load, 0 percent

EL50: Effective Load, 50 percent

ADR: Accord européen sur le transport des marchandises dangereuses par Route (European Agreement concerning the International Carriage of

Dangerous Goods by Road)
IMDG: International Maritime Code for Dangerous Goods

IATA: International Air Transport Association

GHS: Globally Harmonised System of Classification and Labelling of Chemicals

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)

DNEL: Derived No-Effect Level (REACH)

PNEC: Predicted No-Effect Concentration (REACH)

LC50: Lethal concentration, 50 percent

LD50: Lethal dose, 50 percent PBT: Persistent, Bioaccumulative and Toxic

SVHC: Substances of Very High Concern vPvB: very Persistent and very Bioaccumulative

Acute Tox. 1: Acute toxicity – Category 1
Acute Tox. 4: Acute toxicity – Category 4
Skin Irrit. 2: Skin corrosion/irritation – Category 2

Eye Dam. 1: Serious eye damage/eye irritation – Category 1 Repr. 1A: Reproductive toxicity – Category 1A

STOT SE 3: Specific target organ toxicity (single exposure) - Category 3

STOT RE 1: Specific target organ toxicity (repeated exposure) – Category 1 STOT RE 2: Specific target organ toxicity (repeated exposure) – Category 2

Aquatic Acute 1: Hazardous to the aquatic environment - acute aquatic hazard – Category 1

Aquatic Chronic 1: Hazardous to the aquatic environment - long-term aquatic hazard - Category 1

References

- Biocidal Products Committee (BPC) opinion June 2016 on the active substance;
- Assessment Report on the active substance (available at ECHA website)

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- Sources

- 1. The E-Pesticide Manual 2.1 Version (2001)
- The E-Pesticide Manual 2.1 Version (2001)
 Regulation (EC) 1907/2006 and following amendments
 Regulation (EC) 1272/2008 and following amendments
 Regulation (EU) 2015/830
 Regulation (EU) 528/2012
 Regulation (EC) 790/2009 (1st ATP CLP)
 Regulation (EU) 286/2011 (2nd ATP CLP)
 Regulation (EU) 618/2012 (3rd ATP CLP)
 Regulation (EU) 618/2013 (4th ATP CLP)

- 9. Regulation (EU) 487/2013 (4th ATP CLP) 10. Regulation (EU) 944/2013 (5th ATP CLP)

- 11. Regulation (EU) 605/2014 (6th ATP CLP)
 12. Regulation (EU) 2015/1221 (7th ATP CLP)
 13. Regulation (EU) 2016/918 (8th ATP CLP)
 14. Regulation (EU) 2016/1179 (9th ATP CLP)

- 15. Regulation (EU) 2017/776 (10th ATP CLP)
 16. Regulation (EU) 2018/669 (11th ATP CLP)
 17. Directive 2012/18/EU (Seveso III)

- 18. ECHA web site