

Version	Revision Date:	SDS Number:	Date of last issue: 14.08.2024
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SECTION 1: Identification of the substance/mixture and of the company/undertaking

1.1	Product identifier Trade name	:	Diminazene / Phenazone Formulation
1.2	Relevant identified uses of th	ne s	substance or mixture and uses advised against
	Use of the Sub- stance/Mixture	:	Veterinary product
	Recommended restrictions on use	:	Not applicable
1.3	Details of the supplier of the	saf	ety data sheet
	Company	:	MSD Kilsheelan Clonmel Tipperary, IE
	Telephone	:	353-51-601000
	E-mail address of person responsible for the SDS	:	EHSDATASTEWARD@msd.com

1.4 Emergency telephone number

+1-908-423-6000

SECTION 2: Hazards identification

2.1 Classification of the substance or mixture

Classification (REGULATION (EC) No 1272/2008)

Skin irritation, Category 2 Specific target organ toxicity - single exposure, Category 1 Specific target organ toxicity - repeated exposure, Category 1 H315: Causes skin irritation.

H370: Causes damage to organs.

H372: Causes damage to organs through prolonged or repeated exposure.

2.2 Label elements

Labelling (REGULATION (EC) No 1272/2008)

1

•

Hazard pictograms



Signal word

Hazard statements



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			damage to organs. damage to organs through prolonged or re- e.
Preca	utionary statements	P270 Do not e	kin thoroughly after handling. eat, drink or smoke when using this product. otective gloves.
		CENTER/ docto P332 + P313 attention.	IF exposed or concerned: Call a POISON or. If skin irritation occurs: Get medical advice/ Take off contaminated clothing and wash it

Hazardous components which must be listed on the label:

Diminazene

2.3 Other hazards

This substance/mixture contains no components considered to be either persistent, bioaccumulative and toxic (PBT), or very persistent and very bioaccumulative (vPvB) at levels of 0.1% or higher.

Ecological information: The substance/mixture does not contain components considered to have endocrine disrupting properties according to REACH Article 57(f) or Commission Delegated regulation (EU) 2017/2100 or Commission Regulation (EU) 2018/605 at levels of 0.1% or higher.

Toxicological information: The substance/mixture does not contain components considered to have endocrine disrupting properties according to REACH Article 57(f) or Commission Delegated regulation (EU) 2017/2100 or Commission Regulation (EU) 2018/605 at levels of 0.1% or higher.

SECTION 3: Composition/information on ingredients

3.2 Mixtures

Components

Chemical name	CAS-No. EC-No. Index-No. Registration number	Classification	Concentration (% w/w)
Diminazene	536-71-0 208-644-6	Skin Irrit. 2; H315 STOT SE 1; H370 (Brain) STOT RE 1; H372 (Brain)	>= 30 - < 50
Phenazone	60-80-0 200-486-6	Acute Tox. 4; H302	>= 1 - < 10



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			mate	
			Acute oral toxicity: 1.250 mg/kg	

For explanation of abbreviations see section 16.

SECTION 4: First aid measures

4.1 Description of first aid measures

General advice	:	In the case of accident or if you feel unwell, seek medical ad- vice immediately. When symptoms persist or in all cases of doubt seek medical advice.
Protection of first-aiders	:	First Aid responders should pay attention to self-protection, and use the recommended personal protective equipment when the potential for exposure exists (see section 8).
If inhaled	:	If inhaled, remove to fresh air. Get medical attention if symptoms occur.
In case of skin contact	:	In case of contact, immediately flush skin with plenty of water for at least 15 minutes while removing contaminated clothing and shoes. Get medical attention. Wash clothing before reuse. Thoroughly clean shoes before reuse.
In case of eye contact	:	Flush eyes with water as a precaution. Get medical attention if irritation develops and persists.
If swallowed	:	If swallowed, DO NOT induce vomiting unless directed to do so by medical personnel. Get medical attention. Rinse mouth thoroughly with water. Never give anything by mouth to an unconscious person.
4.2 Most important symptoms a	nd e	effects, both acute and delayed
Risks	:	Causes skin irritation. Causes damage to organs. Causes damage to organs through prolonged or repeated exposure.
4.3 Indication of any immediate	med	dical attention and special treatment needed

: Treat symptomatically and supportively. Treatment



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SECTION 5: Firefighting measures

5.1 Extinguishing media

Suitable extinguishing media	:	Water spray Alcohol-resistant foam Carbon dioxide (CO2) Dry chemical
Unsuitable extinguishing media	:	None known.
Special hazards arising from	the	substance or mixture
Specific hazards during fire- fighting	:	Exposure to combustion products may be a hazard to health.
Hazardous combustion prod- ucts	:	Carbon oxides Nitrogen oxides (NOx)
Advice for firefighters		
Special protective equipment for firefighters	:	In the event of fire, wear self-contained breathing apparatus. Use personal protective equipment.
Specific extinguishing meth- ods	:	Use extinguishing measures that are appropriate to local cir- cumstances and the surrounding environment. Use water spray to cool unopened containers. Remove undamaged containers from fire area if it is safe to do so. Evacuate area.
	Unsuitable extinguishing media Special hazards arising from Specific hazards during fire- fighting Hazardous combustion prod- ucts Advice for firefighters Special protective equipment for firefighters Specific extinguishing meth-	media Special hazards arising from the Specific hazards during fire- fighting Hazardous combustion prod- ucts Advice for firefighters Special protective equipment : for firefighters Specific extinguishing meth- :

SECTION 6: Accidental release measures

6.1 Personal precautions, protective equipment and emergency procedures

Personal precautions	:	Use personal protective equipment. Follow safe handling advice (see section 7) and personal pro- tective equipment recommendations (see section 8).
6.2 Environmental precautions		
Environmental precautions	:	Avoid release to the environment. Prevent further leakage or spillage if safe to do so. Prevent spreading over a wide area (e.g. by containment or oil barriers). Retain and dispose of contaminated wash water. Local authorities should be advised if significant spillages cannot be contained.

6.3 Methods and material for containment and cleaning up

Methods for cleaning up	:	Soak up with inert absorbent material.
		For large spills, provide dyking or other appropriate contain-



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		be pumped, stor Clean up remain bent. Local or nationa posal of this man employed in the mine which regu Sections 13 and	aterial from spreading. If dyked material can re recovered material in appropriate container. hing materials from spill with suitable absor- I regulations may apply to releases and dis- terial, as well as those materials and items cleanup of releases. You will need to deter- ulations are applicable. I 15 of this SDS provide information regarding national requirements.

6.4 Reference to other sections

See sections: 7, 8, 11, 12 and 13.

SECTION 7: Handling and storage

7.1 Precautions for safe handling

		0	
	Technical measures	:	See Engineering measures under EXPOSURE CONTROLS/PERSONAL PROTECTION section.
	Local/Total ventilation	:	Use only with adequate ventilation.
	Advice on safe handling	:	Do not get on skin or clothing.
	5		Do not breathe mist or vapours.
			Do not swallow.
			Avoid contact with eyes.
			Wash skin thoroughly after handling.
			Handle in accordance with good industrial hygiene and safety
			practice, based on the results of the workplace exposure as- sessment
			Do not eat, drink or smoke when using this product.
			Take care to prevent spills, waste and minimize release to the environment.
	Hygiene measures	:	If exposure to chemical is likely during typical use, provide eye flushing systems and safety showers close to the working place. When using do not eat, drink or smoke. Wash contami- nated clothing before re-use.
			The effective operation of a facility should include review of engineering controls, proper personal protective equipment, appropriate degowning and decontamination procedures, industrial hygiene monitoring, medical surveillance and the use of administrative controls.
7 2	Conditions for safe storage,	inc	luding any incompatibilities
1.2	.		
	Requirements for storage areas and containers	:	Keep in properly labelled containers. Store locked up. Store in accordance with the particular national regulations.
	Advice on common storage	:	Do not store with the following product types:



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7.3 Specific end use(s)

Specific use(s)

: No data available

SECTION 8: Exposure controls/personal protection

8.1 Control parameters

Occupational Exposure Limits

Components	CAS-No.	Value type (Form of exposure)	Control parameters	Basis
Diminazene	536-71-0	TWA	150 μg/m3 (OEB 2)	Internal

8.2 Exposure controls

Engineering measures

Use appropriate engineering controls and manufacturing technologies to control airborne concentrations (e.g., drip-less quick connections).

All engineering controls should be implemented by facility design and operated in accordance with GMP principles to protect products, workers, and the environment. Laboratory operations do not require special containment.

Personal protective equipment

Eye/face protection	:	Wear safety glasses with side shields or goggles. If the work environment or activity involves dusty conditions, mists or aerosols, wear the appropriate goggles. Wear a faceshield or other full face protection if there is a potential for direct contact to the face with dusts, mists, or aerosols.
Hand protection Material	:	Chemical-resistant gloves
Skin and body protection Respiratory protection	:	Work uniform or laboratory coat. If adequate local exhaust ventilation is not available or expo- sure assessment demonstrates exposures outside the rec- ommended guidelines, use respiratory protection. Equipment should conform to NS EN 143
Filter type	:	Particulates type (P)

SECTION 9: Physical and chemical properties

9.1 Information on basic physical and chemical properties

Physical state	:	liquid
Colour	:	yellow-orange
Odour	:	No data available
Odour Threshold	:	No data available
Melting point/freezing point	:	No data available

according to Regulation (EC) No. 1907/2006, as amended by Commission Regulation (EU) 2020/878



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	Initial b range	oiling point and boiling	:	No data available	
	Flamma	ability (solid, gas)	:	Not applicable	
	Flamma	ability (liquids)	:	No data available	
		explosion limit / Upper bility limit	:	No data available	
		explosion limit / Lower bility limit	:	No data available	
	Flash p	oint	:	No data available	
	Auto-ig	nition temperature	:	No data available	
	Decom	position temperature	:	No data available	
	рН		:	5,0 - 7,0	
	Viscosi Visc	ty :osity, kinematic	:	No data available	
	Solubili Wat	ty(ies) er solubility	:	No data available	
	Partition octanol	n coefficient: n- /water	:	Not applicable	
	Vapour	pressure	:	No data available	
	Relative	e density	:	No data available	
	Density	,	:	No data available	
	Relative	e vapour density	:	No data available	
		characteristics icle size	:	Not applicable	
9.2	Other in	formation			
	Explosi	ves	:	Not explosive	
	Oxidizir	ng properties	:	The substance or	mixture is not classified as oxidizing.
	Evapor	ation rate	:	No data available	

according to Regulation (EC) No. 1907/2006, as amended by Commission Regulation (EU) 2020/878



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Molecular weight		:	No data availal	ble
ECTION	N 10: Stability and	l react	vity	
).1 Reac Not c	tivity lassified as a reactiv	ity haza	ırd.	
	nical stability e under normal conc	ditions.		
).3 Poss	bility of hazardous	s reacti	ons	
	rdous reactions	:		strong oxidizing agents.
	litions to avoid			
Cond	itions to avoid	:	None known.	
).5 Incoi	mpatible materials			
	rials to avoid	:	Oxidizing agen	ts
).6 Haza No ha	rdous decompositi azardous decomposi	ition pro	ducts ducts are known.	
0.6 Haza No ha ECTION	azardous decomposi N 11: Toxicologica mation on hazard o nation on likely route	ition pro al infor	ducts ducts are known. mation	
D.6 Haza No ha ECTION I.1 Infor Inforr expos	azardous decomposi N 11: Toxicologica mation on hazard o nation on likely route sure e toxicity	al infor al infor classes es of :	ducts ducts are known. mation as defined in Re Inhalation Skin contact Ingestion Eye contact	
D.6 Haza No ha ECTION I.1 Infor Inforr expose Acute Not c	azardous decomposi N 11: Toxicologica mation on hazard o nation on likely route sure e toxicity lassified based on a	al infor al infor classes es of :	ducts ducts are known. mation as defined in Re Inhalation Skin contact Ingestion Eye contact	
D.6 Haza No ha ECTION I.1 Infor Inforr expose Acute Not c <u>Prod</u>	azardous decomposi N 11: Toxicologica mation on hazard o nation on likely route sure e toxicity lassified based on a	ition pro al infor classes es of : vailable	ducts ducts are known. mation as defined in Re Inhalation Skin contact Ingestion Eye contact information.	egulation (EC) No 1272/2008
D.6 Haza No ha ECTION I.1 Inforr expose Acute Not c Prod Acute	azardous decomposi N 11: Toxicologica mation on hazard o nation on likely route sure e toxicity lassified based on a <u>uct:</u>	ition pro al infor classes es of : vailable	ducts ducts are known. mation as defined in Re Inhalation Skin contact Ingestion Eye contact information.	egulation (EC) No 1272/2008
D.6 Haza No ha ECTION I.1 Infor Inforr expose Acute Not c Prod Acute	Azardous decomposi A 11: Toxicologica mation on hazard of nation on likely route sure e toxicity lassified based on an <u>uct:</u> e oral toxicity	ition pro al infor classes es of : vailable	ducts ducts are known. mation as defined in Re Inhalation Skin contact Ingestion Eye contact information.	egulation (EC) No 1272/2008
D.6 Haza No ha ECTION I.1 Infor Inforr expose Acute Not c <u>Prod</u> Acute Dimit Acute	azardous decomposi N 11: Toxicologica mation on hazard o nation on likely route sure e toxicity lassified based on a <u>uct:</u> e oral toxicity ponents:	ition pro al infor classes es of : vailable :	ducts ducts are known. mation as defined in Re Inhalation Skin contact Ingestion Eye contact information. Acute toxicity es Method: Calcula	egulation (EC) No 1272/2008
D.6 Haza No ha ECTION I.1 Infor Inforr expose Acute Not c <u>Prod</u> Acute Dimit Acute	azardous decomposi N 11: Toxicologica mation on hazard o nation on likely route sure e toxicity lassified based on a <u>uct:</u> e oral toxicity ponents: nazene: e toxicity (other route	ition pro al infor classes es of : vailable :	ducts ducts are known. mation as defined in Re Inhalation Skin contact Ingestion Eye contact information. Acute toxicity es Method: Calcula LD50 (Rat): 663 Application Rou LD50 (Mouse):	egulation (EC) No 1272/2008 stimate: > 2.000 mg/kg ation method

according to Regulation (EC) No. 1907/2006, as amended by Commission Regulation (EU) 2020/878



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	azone: e oral toxicity	:	LD50 (Cat): 1.250) mg/kg
-	corrosion/irritation			
<u>Com</u>	ponents:			
Dimi	nazene:			
Spec Resu		:	Rabbit Skin irritation	
	bus eye damage/eye irr classified based on availa			
Resp	piratory or skin sensitis	satio	on	
-	sensitisation	able	information.	
-	biratory sensitisation classified based on availa	able	information.	
	n cell mutagenicity classified based on availa	able	information.	
Com	ponents:			
Dimi	nazene:			
Gend	otoxicity in vitro	:	Test system: Saln	bial mutagenesis assay (Ames test) nonella typhimurium icity (Salmonella typhimurium - reverse mu-
			Test Type: Micror Test system: Mou Result: negative	
				o mammalian cell gene mutation test nese hamster cells
Geno	otoxicity in vivo	:	Test Type: Micror Species: Mouse Result: negative	nucleus test
Gern sessi	n cell mutagenicity- As- ment	:	Weight of evidenc	e does not support classification as a germ
Pher	nazone:			
	otoxicity in vitro	:	Test Type: Bacter	ial reverse mutation assay (AMES)



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			Result: negativ	e
Genotoxicity in vivo		:	cytogenetic as Species: Mous Application Ro	e ute: Ingestion) Test Guideline 474
	nogenicity lassified based on avai	ilable	information.	
-	oductive toxicity lassified based on avai	ilahla	information	
	oonents:	liable	iniomation.	
Dimir	nazene:			
	nazene: is on foetal develop-	:	Species: Rat Application Ro General Toxicit Developmental	roductive and developmental toxicity study ute: Oral ty Maternal: LOAEL: 800 mg/kg body weight Toxicity: LOAEL: 800 mg/kg body weight eletal malformations, Embryo-foetal toxicity
Effect		:	Species: Rat Application Ro General Toxicii Developmental Symptoms: Ske Test Type: rep Species: Rat Application Ro General Toxicii	ute: Oral ty Maternal: LOAEL: 800 mg/kg body weight Toxicity: LOAEL: 800 mg/kg body weight eletal malformations, Embryo-foetal toxicity roductive and developmental toxicity study
Effect ment	s on foetal develop-	:	Species: Rat Application Ro General Toxicit Developmental Symptoms: Ske Test Type: rep Species: Rat Application Ro General Toxicit Developmental	ute: Oral ty Maternal: LOAEL: 800 mg/kg body weight Toxicity: LOAEL: 800 mg/kg body weight eletal malformations, Embryo-foetal toxicity roductive and developmental toxicity study ute: Oral ty Maternal: NOAEL: 400 mg/kg body weight Toxicity: NOAEL: 400 mg/kg body weight ave shown reproductive toxicity effects on la-
Effect ment Repro	s on foetal develop-	:	Species: Rat Application Roi General Toxicit Developmental Symptoms: Ski Test Type: rep Species: Rat Application Roi General Toxicit Developmental Experiments ha	ute: Oral ty Maternal: LOAEL: 800 mg/kg body weight Toxicity: LOAEL: 800 mg/kg body weight eletal malformations, Embryo-foetal toxicity roductive and developmental toxicity study ute: Oral ty Maternal: NOAEL: 400 mg/kg body weight Toxicity: NOAEL: 400 mg/kg body weight ave shown reproductive toxicity effects on la-

STOT - single exposure

Causes damage to organs.

according to Regulation (EC) No. 1907/2006, as amended by Commission Regulation (EU) 2020/878



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Com	oonents:			
	nazene:			
Expos Targe	Exposure routes Target Organs Assessment			ice significant health effects in animals at coi 000 mg/kg bw or less.
стот	- repeated exposu	re		
Cause	es damage to organs	through prol	onged or r	epeated exposure.
Com	ponents:			
Dimir	nazene:			
Expos Targe	sure routes et Organs ssment			e to organs through prolonged or repeated
Repe	ated dose toxicity			
<u>Com</u>	oonents:			
Dimir	nazene:			
		: Rat : 63 m : Oral : 3 Mo		
		: Rat : 300 i : Oral : 9 Mo	mg/kg inths	
Expo	EL cation Route sure time et Organs	: Dog : 60 m : Oral : 9 Mo : Brair : Diso	onths n, Testis	
Phen	azone:			
Speci NOAE Applic	es	: Dog : 63 m : Inges : 6 Mo	stion	
۸enir	ation toxicity			

Aspiration toxicity

Not classified based on available information.



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11.2 Information on other hazards

Endocrine disrupting properties

Product:

Assessment

: The substance/mixture does not contain components considered to have endocrine disrupting properties according to REACH Article 57(f) or Commission Delegated regulation (EU) 2017/2100 or Commission Regulation (EU) 2018/605 at levels of 0.1% or higher.

Experience with human exposure

Components:

Diminazene:

Ingestion	: Target Organs: Stomach
-	Symptoms: Vomiting
	Target Organs: Central nervous system
	Symptoms: paralysis
	Target Organs: Immune system
	Symptoms: Fever

SECTION 12: Ecological information

12.1 Toxicity

Components:

Phenazone:

Toxicity to fish	:	LC50 (Oryzias latipes (Japanese medaka)): > 100 mg/l Exposure time: 96 h Method: OECD Test Guideline 203
Toxicity to daphnia and other aquatic invertebrates	:	EC50 (Daphnia magna (Water flea)): >= 1.000 mg/l Exposure time: 48 h Method: OECD Test Guideline 202
Toxicity to algae/aquatic plants	:	ErC50 (Selenastrum capricornutum (green algae)): > 1.000 mg/l Exposure time: 72 h Method: OECD Test Guideline 201
		NOEC (Selenastrum capricornutum (green algae)): 10 mg/l Exposure time: 72 h Method: OECD Test Guideline 201
Toxicity to microorganisms	:	EC50 : 16.900 mg/l Exposure time: 48 h
Toxicity to daphnia and other aquatic invertebrates (Chron- ic toxicity)	:	NOEC: 100 mg/l Exposure time: 21 d Species: Daphnia magna (Water flea)

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		Method: OE0	CD Test Guideline 211		
12.2 Pers	istence and degradabi	lity			
<u>Com</u>	ponents:				
Phen	azone:				
Biodegradability		: Result: Not ir Biodegradati Exposure tim			
12.3 Bioa	ccumulative potential				
<u>Com</u>	ponents:				
Partit	azone: ion coefficient: n- iol/water	: log Pow: 0,38	3		
	i lity in soil ata available				
12.5 Resu	Ilts of PBT and vPvB a	issessment			
<u>Prod</u> Asse	<u>uct:</u> ssment	to be either p	ce/mixture contains no components considered ersistent, bioaccumulative and toxic (PBT), or nt and very bioaccumulative (vPvB) at levels of er.		
12.6 Endo	ocrine disrupting prop	erties			
Prod	uct:				
	ssment	ered to have REACH Artic	ce/mixture does not contain components consid- endocrine disrupting properties according to le 57(f) or Commission Delegated regulation 00 or Commission Regulation (EU) 2018/605 at 6 or higher.		
12.7 Other adverse effects					
No data available					
SECTION 13: Disposal considerations					
13.1 Waste treatment methods					

Product

: Dispose of in accordance with local regulations. According to the European Waste Catalogue, Waste Codes are not product specific, but application specific. Waste codes should be assigned by the user, preferably in discussion with the waste disposal authorities.



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Contaminated packaging		:	 Do not dispose of waste into sewer. Empty containers should be taken to an approved waste handling site for recycling or disposal. If not otherwise specified: Dispose of as unused product. 			
SECTION	N 14: Transport info	rmat	tion			
14.1 UN n	umber or ID number					
ADN		:	Not regulated as	a dangerous good		
ADR		:	Not regulated as	a dangerous good		
RID		:	Not regulated as	a dangerous good		
IMDO	6	:	Not regulated as	a dangerous good		
ΙΑΤΑ		:	Not regulated as	a dangerous good		
14.2 UN p	roper shipping name					
ADN		:	Not regulated as	a dangerous good		
ADR		:	Not regulated as	a dangerous good		
RID		:	Not regulated as	a dangerous good		
IMDO	6	:	Not regulated as	a dangerous good		
ΙΑΤΑ		:	Not regulated as	a dangerous good		
4.3 Tran	sport hazard class(es))				
ADN		:	Not regulated as	a dangerous good		
ADR		:	Not regulated as	a dangerous good		
RID		:	Not regulated as	a dangerous good		
IMDO	5	:	Not regulated as	a dangerous good		
ΙΑΤΑ		:	Not regulated as	a dangerous good		
14.4 Pack	ing group					
ADN		:	Not regulated as	a dangerous good		
ADR		:	Not regulated as	a dangerous good		
RID		:	Not regulated as	a dangerous good		
IMDO	6	:	Not regulated as	a dangerous good		
ΙΑΤΑ	(Cargo)	:	Not regulated as	a dangerous good		
ΙΑΤΑ	(Passenger)	:	Not regulated as	a dangerous good		
	ronmental hazards					
	egulated as a dangerou	-	od			
•	ial precautions for us	er				

Not applicable

14.7 Maritime transport in bulk according to IMO instruments



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Rema	arks	: Not applicable for	product	as supplied.
SECTION	15: Regulatory inf	ormation		
15.1 Safe ture	ty, health and enviror	nmental regulations/leg	jislation	specific for the substance or mix-
the m		e manufacture, placing o in dangerous substances x XVII)		Conditions of restriction for the fol- lowing entries should be considered: Number on list 3
				Substance(s) or mixture(s) are listed here according to their appearance in the regulation, irrespective of their use/purpose or the conditions of the restriction. Please refer to the condi- tions in corresponding Regulation to determine whether an entry is appli- cable to the placing on the market or not.
Conc	ern for Authorisation (A	Substances of Very High Article 59). s subject to authorisation		Not applicable
	ex XIV)		-	Not applicable
· ·	,	ces that deplete the ozo	ne :	Not applicable
	lation (EU) 2019/1021 (recast)	on persistent organic po	ollu- :	Not applicable
ment		12 of the European Parlia erning the export and imp		Not applicable

Seveso III: Directive 2012/18/EU of the European Parliament and of the Council on the control of major-accident hazards involving dangerous substances.

		Quantity 1	Quantity 2
H3	STOT SPECIFIC TARGET	50 t	200 t
	ORGAN TOXICITY –		
	SINGLE EXPOSURE		

Other regulations:

Note the Working Environment Act § 4-1 and § 4-2 on requirements for the employer to protect pregnant employees against discomfort and injury as a result of the work situation and the working environment.

Note the regulation on organization, leadership and participation, chapter 12 on the work of children and young people.

The components of this product are reported in the following inventories:

AICS	:	not determined
DSL	:	not determined
IECSC	:	not determined



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15.2 Chemical safety assessment

A Chemical Safety Assessment has not been carried out.

SECTION 16: Other information

Other information : Items where changes have been made to the previous version are highlighted in the body of this document by two vertical lines.

Full text of H-Statements

		Harmful if swallowed. Causes skin irritation.
H370	:	Causes damage to organs if swallowed.
H372	:	Causes damage to organs through prolonged or repeated exposure if swallowed.

Full text of other abbreviations

Acute Tox. :	Acute toxicity
Skin Irrit. :	Skin irritation
STOT RE :	Specific target organ toxicity - repeated exposure
STOT SE :	Specific target organ toxicity - single exposure

ADN - European Agreement concerning the International Carriage of Dangerous Goods by Inland Waterways; ADR - Agreement concerning the International Carriage of Dangerous Goods by Road; AIIC - Australian Inventory of Industrial Chemicals; ASTM - American Society for the Testing of Materials; bw - Body weight; CLP - Classification Labelling Packaging Regulation; Regulation (EC) No 1272/2008; CMR - Carcinogen, Mutagen or Reproductive Toxicant; DIN - Standard of the German Institute for Standardisation; DSL - Domestic Substances List (Canada); ECHA -European Chemicals Agency; EC-Number - European Community number; ECx - Concentration associated with x% response; ELx - Loading rate associated with x% response; EmS - Emergency Schedule; ENCS - Existing and New Chemical Substances (Japan); ErCx - Concentration associated with x% growth rate response; GHS - Globally Harmonized System; GLP - Good Laboratory Practice; IARC - International Agency for Research on Cancer; IATA - International Air Transport Association; IBC - International Code for the Construction and Equipment of Ships carrying Dangerous Chemicals in Bulk; IC50 - Half maximal inhibitory concentration; ICAO - International Civil Aviation Organization; IECSC - Inventory of Existing Chemical Substances in China; IMDG - International Maritime Dangerous Goods; IMO - International Maritime Organization; ISHL - Industrial Safety and Health Law (Japan); ISO - International Organisation for Standardization; KECI - Korea Existing Chemicals Inventory; LC50 - Lethal Concentration to 50 % of a test population; LD50 - Lethal Dose to 50% of a test population (Median Lethal Dose); MARPOL - International Convention for the Prevention of Pollution from Ships; n.o.s. - Not Otherwise Specified; NO(A)EC - No Observed (Adverse) Effect Concentration; NO(A)EL - No Observed (Adverse) Effect Level; NOELR - No Observable Effect Loading Rate; NZIoC - New Zealand Inventory of Chemicals; OECD - Organization for Economic Co-operation and Development; OPPTS - Office of Chemical Safety and Pollution Prevention; PBT - Persistent, Bioaccumulative and Toxic substance; PICCS - Philippines Inventory of Chemicals and Chemical Substances; (Q)SAR - (Quantitative) Structure Activity Relationship; REACH - Regulation (EC) No 1907/2006 of the European Parliament and of the Council concerning the Registration, Evaluation, Authorisation and Restriction of Chemicals; RID - Regulations concerning the International Carriage of Dangerous Goods by Rail; SADT - Self-Accelerating Decomposition Temperature; SDS - Safety Data Sheet; SVHC - Substance of very high concern; TCSI - Taiwan Chemical Substance Inventory; TECI -Thailand Existing Chemicals Inventory; TSCA - Toxic Substances Control Act (United States); UN



Diminazene / Phenazone Formulation

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- United Nations; UNRTDG - United Nations Recommendations on the Transport of Dangerous Goods; vPvB - Very Persistent and Very Bioaccumulative

Further information

Sources of key data used to compile the Safety Data Sheet		, data from raw material SDSs, OECD results and European Chemicals Agen- eu/
Classification of the mixtur	re:	Classification procedure:
Skin Irrit. 2	H315	Calculation method
STOT SE 1	H370	Calculation method
STOT RE 1	H372	Calculation method

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