according to the Globally Harmonized System



Milbemycin Oxime / Lufenuron Formulation

	Date of last issue: 30.09.2023 Date of first issue: 21.09.2020
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1. PRODUCT AND COMPANY IDENTIFICATION

Product name	:	Milbemycin Oxime / Lufenuron Formulation
Manufacturer or supplier's d	eta	ils
Company	:	MSD
Address	:	Briahnager - Off Pune Nagar Road Wagholi - Pune - India 412 207
Telephone	:	+1-908-740-4000
Emergency telephone number	:	+1-908-423-6000
E-mail address	:	EHSDATASTEWARD@msd.com
Recommended use of the ch	em	ical and restrictions on use
Recommended use Restrictions on use	:	Veterinary product Not applicable

2. HAZARDS IDENTIFICATION

Manufacture, Storage and Import of Hazardous Chemicals Rules 1989

Classification

Not classified as hazardous according to criteria laid down in Part I of Schedule-1.

	001	
GHS Classification		
Skin sensitisation	:	Category 1
Reproductive toxicity	:	Category 1B
Specific target organ toxicity - repeated exposure	:	Category 2 (Central nervous system)
Specific target organ toxicity - repeated exposure (Oral)	:	Category 1 (Central nervous system, Lungs, Liver, Stomach)
Short-term (acute) aquatic hazard	:	Category 1
Long-term (chronic) aquatic hazard	:	Category 1
GHS label elements		

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Haza	rd pictograms		!
Signa	al word	: Danger	\mathbf{v}
Haza	rd statements	H360D May da H372 Causes Lungs, Liver, S sure if swallow H373 May cau through prolon	se an allergic skin reaction. amage the unborn child. damage to organs (Central nervous system, Stomach) through prolonged or repeated expo red. se damage to organs (Central nervous system ged or repeated exposure. ic to aquatic life with long lasting effects.
Preca	autionary statements	P260 Do not b P264 Wash ha P270 Do not e P272 Contami the workplace. P273 Avoid rel	lease to the environment. otective gloves/ protective clothing/ eye protec
		P318 IF expos P333 + P317 I	F ON SKIN: Wash with plenty of water. ed or concerned, get medical advice. f skin irritation or rash occurs: Get medical hel Fake off contaminated clothing and wash it bet pillage.
		Storage: P405 Store loc	ked up.
		Disposal:	of contents/ container to an approved waste

Other hazards which do not result in classification

None known.

3. COMPOSITION/INFORMATION ON INGREDIENTS

Substance / Mixture : Mixture

Components

Chemical name	CAS-No.	Concentration (% w/w)
Lufenuron (ISO)	103055-07-8	>= 30 - < 50
Cellulose	9004-34-6	>= 10 - < 20

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1	Starch				9005-25-8	>= 5 - < 10	
		ycin Oxime			129496-10-2	>= 1 - < 2.5	
4. FI	RST AI	D MEASURES					
	Genera	al advice	:	vice immediatel	ccident or if you feel unwe y. s persist or in all cases of		
	If inhale	ed	:	If inhaled, remo			
	In case	of skin contact	:	Get medical attention. In case of contact, immediately flush skin with soap and plenty of water. Remove 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.		
	If swall	owed	:	Get medical attention if irritation develops and persists. If swallowed, DO NOT induce vomiting. Get medical attention.			
		nportant symptoms ects, both acute and d	:	May damage the unborn child. Causes damage to organs through prolonged or repeated exposure if swallowed. May cause damage to organs through prolonged or repeated			
	Protect	ion of first-aiders	:	exposure. 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).			
	Notes t	o physician	:		atically and supportively.		
5. FI	REFIGI	HTING MEASURES					
	Suitabl	e extinguishing media	:	Water spray Alcohol-resistar Carbon dioxide Dry chemical			
	Unsuita media	able extinguishing	:	None known.			
	Specifi fighting	c hazards during fire-	:	Exposure to cor	nbustion products may be	e a hazard to health.	
	Hazard ucts	lous combustion prod-	:	Carbon oxides Nitrogen oxides Metal oxides	(NOx)		

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			Evacuate area.		
	cial protective equipment refighters	:		e, wear self-contained breathing apparatus. btective equipment.	
6. ACCID	ENTAL RELEASE MEA	SUF	RES		
tive e	onal precautions, protec- equipment and emer- y procedures	:	Follow safe hand	otective equipment. Iling advice (see section 7) and personal pro It recommendations (see section 8).	
Envir	onmental precautions	:	Avoid release to the environment. Prevent further leakage or spillage if safe to do so. Retain and dispose of contaminated wash water. Local authorities should be advised if significant spillages cannot be contained.		
	ods and materials for ainment and cleaning up	:	tainer for disposa Local or national posal of this mat employed in the mine which regu Sections 13 and	cuum up spillage and collect in suitable con- al. regulations may apply to releases and dis- erial, as well as those materials and items cleanup of releases. You will need to deter- lations are applicable. 15 of this SDS provide information regarding ational requirements.	
7. HANDL	ING AND STORAGE				
Tech	nical measures	:		measures under EXPOSURE RSONAL PROTECTION section.	
Loca	I/Total ventilation	:		ation is unavailable, use with local exhaust	
Advid	ce on safe handling	:	Do not get on skin or clothing. Do not breathe dust, fume, gas, mist, vapours or spray. Do not swallow. Avoid contact with eyes. Wash skin thoroughly after handling. Handle in accordance with good industrial hygiene and safet practice, based on the results of the workplace exposure as- sessment Keep container tightly closed. Do not eat, drink or smoke when using this product. Take care to prevent spills, waste and minimize release to th		
Conc	litions for safe storage	:	environment. Keep in properly labelled containers. Store locked up. Keep tightly closed. Store in accordance with the particular national regulations.		
			D		

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8. EXPOSURE CONTROLS/PERSONAL PROTECTION

Components with workplace control parameters

Components	CAS-No.	Value type (Form of exposure)	Control parame- ters / Permissible concentration	Basis
Lufenuron (ISO)	103055-07-8	TWA	60 µg/m3 (OEB 3)	Internal
	Further informa	ation: DSEN		
		Wipe limit	100 µg/100 cm2	Internal
Cellulose	9004-34-6	TWA	10 mg/m3	ACGIH
Starch	9005-25-8	TWA	10 mg/m3	ACGIH
Milbemycin Oxime	129496-10-2	TWA	0.1 mg/m3 (OEB2)	Internal

Engineering measures	All engineering controls should be implemented by facility design and operated in accordance with GMP principles to protect products, workers, and the environment. Containment technologies suitable for controlling compounds are required to control at source and to prevent migration of the compound to uncontrolled areas (e.g., open-face contain- ment devices). Minimize open handling.
Personal protective equipme	nt
Respiratory protection Filter type Hand protection	If adequate local exhaust ventilation is not available or expo- sure assessment demonstrates exposures outside the rec- ommended guidelines, use respiratory protection. Particulates type
Material	Chemical-resistant gloves
Remarks Eye protection	Consider double gloving. 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.
Skin and body protection	 Work uniform or laboratory coat. Additional body garments should be used based upon the task being performed (e.g., sleevelets, apron, gauntlets, disposable suits) to avoid exposed skin surfaces. Use appropriate degowning techniques to remove potentially contaminated clothing.
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. Contaminated work clothing should not be allowed out of the workplace. Wash contaminated clothing before re-use.

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			engineering contr appropriate dego	eration of a facility should include review of rols, proper personal protective equipment, wning and decontamination procedures, e monitoring, medical surveillance and the tive controls.
9. PHYSI	CAL AND CHEMICAL P	ROF	PERTIES	
Арре	earance	:	solid	
Colo	ur	:	brown	
Odo	ur	:	odourless	
Odo	ur Threshold	:	No data available	e
рН		:	No data available	e
Melt	ing point/freezing point	:	No data available	e
Initia rang	I boiling point and boiling e	:	No data available	e
Flas	h point	:	Not applicable	
Evap	poration rate	:	Not applicable	
Flam	nmability (solid, gas)	:	No data available	e
Flam	nmability (liquids)	:	Not applicable	
	er explosion limit / Upper mability limit	:	No data available	e
	er explosion limit / Lower mability limit	:	No data available	e
Vapo	our pressure	:	Not applicable	
Rela	tive vapour density	:	Not applicable	
Rela	tive density	:	No data available	e
Dens	sity	:	No data available	e
	bility(ies) Vater solubility	:	soluble	
	tion coefficient: n-	:	Not applicable	
	nol/water -ignition temperature	:	No data available	e
Deco	omposition temperature	:	No data available	e
Visc	osity			

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Vi	scosity, kinematic	: Not applical	ble	
Explo	sive properties	: Not explosiv	/e	
Oxidiz	zing properties	: The substar	nce or mixture is not classified as oxidizing.	
Moleo	cular weight	: No data ava	ailable	
	le characteristics le size	: No data ava	ailable	
10. STABI	LITY AND REACTIVIT	Y		
Possi tions Cond Incom	nical stability bility of hazardous read itions to avoid npatible materials rdous decomposition	: Stable unde : Can react w : None known : Oxidizing ag		
11. TOXIC	OLOGICAL INFORMA	TION		
Inform	nation on likely routes c sure	f : Skin contact Ingestion Eye contact		
	e toxicity lassified based on avail	able information.		
Prod	uct:			
Acute	e oral toxicity		y estimate: > 5,000 mg/kg culation method	
Acute	inhalation toxicity	Exposure tin Test atmosp	y estimate: > 10 mg/l ne: 4 h here: dust/mist culation method	

Acute dermal toxicity : Acute toxicity estimate: > 5,000 mg/kg Method: Calculation method

Components:

Lufenuron (ISO):		
Acute oral toxicity	: LD50 (Rat): > 2,000 mg/kg	
	LD50 (Mouse): > 2,000 mg/kg	
Acute inhalation toxicit	: LC50 (Rat): 2,350 mg/m3 Test atmosphere: dust/mist	

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Acute	e dermal toxicity	:	LD50 (Rabbit):	> 2,000 mg/kg
I				
Cellu				
Acute	e oral toxicity	:	LD50 (Rat): > 5	5,000 mg/kg
Acute	inhalation toxicity	:	LC50 (Rat): > 5 Exposure time: Test atmosphe	:4 h
Acute	e dermal toxicity	:	LD50 (Rabbit):	> 2,000 mg/kg
Starc	:h:			
Acute	e oral toxicity	:	LD50 (Rat): > 5	5,000 mg/kg
Acute	e dermal toxicity	:	LD50 (Rabbit):	> 2,000 mg/kg
Milbe	emycin Oxime:			
	e oral toxicity	:	LD50 (Rat): 53	2 - 863 mg/kg
			LD50 (Mouse):	722 - 946 mg/kg
Acute	e inhalation toxicity	:	LC50 (Rat): 1,2	
			Exposure time: Test atmosphe	
Acute	e dermal toxicity	:	LD50 (Rat): > 2	
Skin	corrosion/irritation			
Not c	lassified based on ava	ilable	information.	
Com	ponents:			
Lufer	nuron (ISO):			
Speci	ies	:	Rabbit	
Metho		:	Draize Test	
Resu	IT	:	No skin irritatio	'n
Milbe	emycin Oxime:			
Speci		:	Rabbit	
Metho		:	OECD Test Gu	
Resu		:	No skin irritatio	n
	ous eye damage/eye i lassified based on ava			
Com	ponents:			
	nuron (ISO):			
Speci		:	Rabbit	
Metho		:	Draize Test	n
Resu	n	:	No eye irritation	11

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Starc	:h:			
Spec		:	Rabbit	
Resu	It	:	No eye irritation	
Milbe	emycin Oxime:			
Spec Resu	ies It	:	Rabbit No eye irritation	
, toou	i.	•		
Resp	iratory or skin sensi	tisatio	on	
	sensitisation			
	cause an allergic skin		on.	
-	iratory sensitisation lassified based on ava		information	
_	ponents:			
	nuron (ISO):			
Test	Туре	:	Maximisation Te	est
Spec	ies ssment	:	Guinea pig	itisation by skin contact.
Resu		:	Sensitiser	
Starc	:h:			
Test	Туре	:	Maximisation Te	est
	sure routes	:	Skin contact	
Spec Resu		:	Guinea pig negative	
	emycin Oxime: sure routes	:	Skin contact	
Spec	ies		Guinea pig	
Resu	lt	:	negative	
Germ	n cell mutagenicity			
Not c	lassified based on ava	ailable	information.	
<u>Com</u>	ponents:			
	nuron (ISO):			
Geno	otoxicity in vitro	:	Test Type: Ame Result: negative	
			-	
			Test Type: Mous	se Lymphoma iinese hamster cells
			Result: negative	
			Test Type: Cyto	genetic assay
				inese hamster ovary cells

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		Populti pogotivo	
		Result: negative	
		Test Type: DNA damage and repair, unscheduled DNA s thesis in mammalian cells (in vitro) Test system: rat hepatocytes Result: negative	yn-
		Test system: Human lymphocytes Result: negative	
Geno	otoxicity in vivo	: Test Type: Mammalian erythrocyte micronucleus test (in cytogenetic assay) Species: Mouse Result: negative	vivo
		Test Type: Unscheduled DNA synthesis test (UDS) in tes lar cells Species: Rat Result: negative	sticu-
	n cell mutagenicity - essment	: Weight of evidence does not support classification as a g cell mutagen.	erm
Cellu	ulose:		
Geno	otoxicity in vitro	: Test Type: Bacterial reverse mutation assay (AMES) Result: negative	
		Test Type: In vitro mammalian cell gene mutation test Result: negative	
Gene	otoxicity in vivo	 Test Type: Mammalian erythrocyte micronucleus test (in cytogenetic assay) Species: Mouse Application Route: Ingestion Result: negative 	vivo
Star	ch:		
Geno	otoxicity in vitro	: Test Type: Bacterial reverse mutation assay (AMES) Result: negative	
II Milb	emycin Oxime:		
	otoxicity in vitro	: Test Type: Bacterial reverse mutation assay (AMES) Result: negative	
		Test Type: Chromosome aberration test in vitro Result: negative	
Geno	otoxicity in vivo	: Test Type: Mammalian erythrocyte micronucleus test (in cytogenetic assay) Species: Mouse Result: negative	vivo

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	sion Revision Date: SDS Number: 28.09.2024 6365220-00009		Date of last issue: 30.09.2023 Date of first issue: 21.09.2020			
	nogenicity					
Not c	lassified based on ava	allable information.				
Com	ponents:					
Lufer	nuron (ISO):					
Speci Applic Expos Resu	cation Route sure time	: Rat : Ingestion : 18 month(s)				
	nogenicity - Assess-	-	negative Weight of evidence does not support classification as a car- cinogen			
Cellu	lose:					
	cation Route sure time	: Rat : Ingestion : 72 weeks : negative				
May o	oductive toxicity damage the unborn ch	ild.				
May o <u>Com</u>	-	ild.				
May o <u>Com</u> Lufer	damage the unborn ch ponents:	: Test Type: T Species: Ra Application F General Tox Early Embry weight				
May o Com Lufer	damage the unborn ch ponents: nuron (ISO):	 Test Type: T Species: Ra Application F General Tox Early Embry weight Result: Anim Test Type: D Species: Ra Application F General Tox Developmen Symptoms: I 	t Route: Oral icity - Parent: NOAEL: 8.3 mg/kg wet weight onic Development: NOAEL: 20.9 mg/kg body nal testing did not show any effects on fertility. Development t Route: Oral icity Maternal: NOAEL: 500 mg/kg body weigh			
May o <u>Com</u> Lufer Effect	damage the unborn ch ponents: nuron (ISO): ts on fertility	 Test Type: T Species: Rai Application F General Tox Early Embry weight Result: Anim Test Type: D Species: Rai Application F General Tox Developmen Symptoms: I Remarks: No Test Type: F Species: Rai Application F General Tox Embryo-foet 	t Route: Oral icity - Parent: NOAEL: 8.3 mg/kg wet weight onic Development: NOAEL: 20.9 mg/kg body nal testing did not show any effects on fertility. Development t Route: Oral icity Maternal: NOAEL: 500 mg/kg body weigh tal Toxicity: NOAEL: 1,000 mg/kg body weigh No adverse effects o significant adverse effects were reported Fertility/early embryonic development			

Cellulose:

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Effect	s on fertility	Species: Rat	ne-generation reproduction toxicity study oute: Ingestion ve			
Effect ment	s on foetal develop-	Species: Rat	rtility/early embryonic development oute: Ingestion ve			
Milbe	mycin Oxime:					
Effect	s on fertility	: Test Type: One-generation reproduction toxicity study Species: Dog Application Route: Ingestion Result: negative				
Effect ment	s on foetal develop-	Species: Rat	nbryo-foetal development oute: Ingestion ve			
		Species: Rabl	oute: Ingestion			
		Species: Dog	nbryo-foetal development oute: Ingestion ve			

STOT - single exposure

Not classified based on available information.

Components:

Lufenuron (ISO):

Assessment

: The substance or mixture is not classified as specific target organ toxicant, single exposure.

STOT - repeated exposure

Causes damage to organs (Central nervous system, Lungs, Liver, Stomach) through prolonged or repeated exposure if swallowed.

May cause damage to organs (Central nervous system) through prolonged or repeated exposure.

Components:

Lufenuron (ISO):

Exposure routes	:	Oral
Target Organs	:	Central nervous system, Lungs, Liver, Stomach
Exposure routes Target Organs Assessment	:	Shown to produce significant health effects in animals at con- centrations of 10 mg/kg bw or less.

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Expos	mycin Oxime: sure routes	: Ingestion	
	et Organs ssment	: Central nervous : Shown to produce centrations of 1	s system uce significant health effects in animals at con- 0 mg/kg bw or less.
Repe	ated dose toxicity		
Com	oonents:		
Speci NOAE Applic Expos	EL cation Route sure time et Organs	: Rat : 5.34 mg/kg : oral (feed) : 4 Months : Central nervous : central nervous	s system, digestive system system effects
	EL cation Route sure time	: Rat : 1.93 mg/kg : oral (feed) : 2 yr : central nervous	system effects, Convulsions
Expo	EL cation Route sure time et Organs		s system, Liver, Prostate system effects, Convulsions
Expo	EL cation Route sure time et Organs		s system, Liver, Lungs atality, Irregularities
	es	: Rat : >= 9,000 mg/kg : Ingestion : 90 Days	9
	es EL cation Route sure time	: Rat : >= 2,000 mg/kg : Skin contact : 28 Days : OECD Test Gu	
Applio Expos	cation Route sure time	: Skin contact : 28 Days	

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Expos Symp Speci	EL EL cation Route sure time toms es	: Dog	3 mg/kg 15 mg/kg Ingestion 90 Days Liver disorders, Blood disorders				
Applic Expos	LOAEL : Application Route : Exposure time : Symptoms :						
-	ation toxicity assified based on ava	ilable information.					
Expe	Experience with human expos						
Comp	oonents:						
Lufen	uron (ISO):						
Gene	ral Information	: Remarks: May May cause neu	be harmful if swallowed. rotoxic effects.				
Milbe	mycin Oxime:						
Ingest	tion	Vomiting, Trem	ivation, Convulsions, Diarrhoea, Weakness, ors, Coma d on Animal Evidence				
12 ECOL							

12. ECOLOGICAL INFORMATION

Ecotoxicity

Components:

Lufenuron (ISO):

Toxicity to fish		LC50 (Oncorhynchus mykiss (rainbow trout)): > 73,100 μg/l Exposure time: 96 h Method: OECD Test Guideline 203
		LC50 (Oncorhynchus mykiss (rainbow trout)): > 29,000 µg/l Exposure time: 96 h Method: OECD Test Guideline 203
		LC50 (Oncorhynchus mykiss (rainbow trout)): 370 µg/l Exposure time: 96 h Method: OECD Test Guideline 203
Toxicity to daphnia and othe aquatic invertebrates	r:	EC50 (Americamysis): 0.042 μg/l Exposure time: 96 h Method: US-EPA OPPTS 850.1035
		EC50 (Daphnia magna (Water flea)): 0.41 μg/l Exposure time: 48 h

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II				Method: OECD Te	est Guideline 202
	Toxicity plants	to algae/aquatic	:		elis subcapitata (freshwater green alga)): h
				EC50 (Scenedes Exposure time: 72 Method: OECD Te	
	M-Facto icity)	or (Acute aquatic tox-	:	10,000	
	Toxicity icity)	to fish (Chronic tox-	:	NOEC: 80 µg/l Exposure time: 33 Species: Oncorhy Method: OECD Te	nchus mykiss (rainbow trout)
				NOEC: 20 µg/l Exposure time: 35 Species: Oncorhy Method: OECD Te	nchus mykiss (rainbow trout)
á		to daphnia and other invertebrates (Chron- y)	:	NOEC: 8.38 µg/l Exposure time: 21 Species: Daphnia Method: OECD Te	magna (Water flea)
				NOEC: 90 µg/l Exposure time: 21 Species: Daphnia Method: OECD Te	magna (Water flea)
				NOEC: 2 µg/l Exposure time: 21 Species: Chironor Method: OECD Te	nus riparius (harlequin fly)
	M-Facto toxicity)	or (Chronic aquatic	:	10	
11	Cellulos	se:			
	Toxicity		:	Exposure time: 48	pes (Japanese medaka)): > 100 mg/l h on data from similar materials
I	Milbem	ycin Oxime:			
	Toxicity		:	LC50 (Oncorhync Exposure time: 96	hus mykiss (rainbow trout)): 0.16 μg/l i h
		to daphnia and other invertebrates	:	EC50 (Daphnia m Exposure time: 48	agna (Water flea)): 0.03 µg/l ⊧h

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Toxic plant	city to algae/aquatic s	:	EC50: > 87 μg/l Exposure time: 72	? h
M-Fa icity)	actor (Acute aquatic tox-	:	10,000	
aqua	city to daphnia and other tic invertebrates (Chron- kicity)			magna (Water flea)
M-Fa toxic	actor (Chronic aquatic ity)	:	10,000	
Pers	istence and degradabili	ty		
Com	ponents:			
Cellu	ulose:			
Biode	egradability	:	Result: Readily bio	odegradable.
Bioa	ccumulative potential			
<u>Com</u>	ponents:			
Lufe	nuron (ISO):			
Bioa	ccumulation	:	Species: Lepomis Bioconcentration f Method: OECD Te	
	tion coefficient: n- nol/water	:	log Pow: 5.12	
Milb	emycin Oxime:			
Bioa	ccumulation	:	Bioconcentration f	actor (BCF): 440
	tion coefficient: n- nol/water	:	log Pow: 7	
Mob	ility in soil			
<u>Com</u>	ponents:			
Lufe	nuron (ISO):			
	ibution among environ- al compartments	:	log Koc: 5.38 Method: OECD Te	est Guideline 106
	e r adverse effects ata available			

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13. DISPOSAL CONSIDERATIONS

Disposal methods		
Waste from residues	Do not dispose of waste int	
	Dispose of in accordance w	vith local regulations.
Contaminated packaging	Empty containers should be dling site for recycling or dis	e taken to an approved waste han-
	If not otherwise specified: D	bispose of as unused product.

14. TRANSPORT INFORMATION

International Regulations

UNRTDG UN number Proper shipping name	:	UN 3077 ENVIRONMENTALLY HAZARDOUS SUBSTANCE, SOLID, N.O.S.
Class Packing group Labels Environmentally hazardous	:	(Milbemycin Oxime, Lufenuron (ISO)) 9 III 9 yes
IATA-DGR UN/ID No. Proper shipping name	:	UN 3077 Environmentally hazardous substance, solid, n.o.s. (Milbemycin Oxime, Lufenuron (ISO))
Class Packing group Labels Packing instruction (cargo	:	9 III Miscellaneous 956
aircraft) Packing instruction (passen- ger aircraft) Environmentally hazardous	:	956 ves
IMDG-Code UN number Proper shipping name	:	UN 3077 ENVIRONMENTALLY HAZARDOUS SUBSTANCE, SOLID, N.O.S.
Class Packing group Labels EmS Code Marine pollutant		(Milbemycin Oxime, Lufenuron (ISO)) 9 III 9 F-A, S-F yes

Transport in bulk according to IMO instruments

Not applicable for product as supplied.

Special precautions for user

The transport classification(s) provided herein are for informational purposes only, and solely based upon the properties of the unpackaged material as it is described within this Safety Data

according to the Globally Harmonized System



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Sheet. Transportation classifications may vary by mode of transportation, package sizes, and variations in regional or country regulations.

15. REGULATORY INFORMATION

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

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

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

16. OTHER INFORMATION

Revision Date	:	28.09.2024
Further information Sources of key data used to compile the Safety Data Sheet	:	Internal technical data, data from raw material SDSs, OECD eChem Portal search results and European Chemicals Agen- cy, http://echa.europa.eu/

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

Date format	:	dd.mm.yyyy
Full text of other abbreviation	ns	
ACGIH	:	USA. ACGIH Threshold Limit Values (TLV)

ACGIH / TWA : 8-hour, time-weighted average

AIIC - Australian Inventory of Industrial Chemicals; ANTT - National Agency for Transport by Land of Brazil; ASTM - American Society for the Testing of Materials; bw - Body weight; CMR -Carcinogen, Mutagen or Reproductive Toxicant; DIN - Standard of the German Institute for Standardisation; DSL - Domestic Substances List (Canada); 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; ERG - Emergency Response Guide; 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; Nch - Chilean Norm; NO(A)EC - No Observed (Adverse) Effect Concentration; NO(A)EL - No Observed (Adverse) Effect Level; NOELR - No Observable Effect Loading Rate; NOM - Official Mexican Norm; NTP - National Toxicology Program; NZIoC - New Zealand Inventory of Chemicals; OECD - Organization for Economic Co-operation and Develop-

according to the Globally Harmonized System



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ment; 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; SADT - Self-Accelerating Decomposition Temperature; SDS - Safety Data Sheet; TCSI - Taiwan Chemical Substance Inventory; TDG - Transportation of Dangerous Goods; TECI - Thailand Existing Chemicals Inventory; TSCA - Toxic Substances Control Act (United States); UN - United Nations; UNRTDG - United Nations Recommendations on the Transport of Dangerous Goods; vPvB - Very Persistent and Very Bioaccumulative; WHMIS - Workplace Hazardous Materials Information System

The information provided in this Safety Data Sheet is correct to the best of our knowledge, information and belief at the date of its publication. The information is designed only as a guidance for safe handling, use, processing, storage, transportation, disposal and release and shall not be considered a warranty or quality specification of any type. The information provided relates only to the specific material identified at the top of this SDS and may not be valid when the SDS material is used in combination with any other materials or in any process, unless specified in the text. Material users should review the information and recommendations in the specific context of their intended manner of handling, use, processing and storage, including an assessment of the appropriateness of the SDS material in the user's end product, if applicable.

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