according to the OSHA Hazard Communication Standard



Triclabendazole / Abamectin Formulation

Version	Revision Date:	SDS Number:	Date of last issue: 04/06/2024
3.0	07/06/2024	5341825-00013	Date of first issue: 12/05/2019

SECTION 1. IDENTIFICATION

Product name	:	Triclabendazole / Abamectin Formulation			
Manufacturer or supplier's o	deta	ails			
Company name of supplier Address	:	Merck & Co., Inc 126 E. Lincoln Avenue Rahway, New Jersey U.S.A. 07065			
Telephone Emergency telephone E-mail address	:	908-740-4000 1-908-423-6000 EHSDATASTEWARD@merck.com			
Recommended use of the chemical and restrictions on use					
Recommended use Restrictions on use	:	Veterinary product Not applicable			

SECTION 2. HAZARDS IDENTIFICATION

GHS classification in accordance with the OSHA Hazard Communication Standard (29 CFR
1910.1200)

Specific target organ toxicity : Category 2 (Liver, Blood)

- repeated exposure (Oral)

GHS label elements

Hazard pictograms	:	
Signal Word	:	Warning
Hazard Statements	:	H373 May cause damage to organs (Liver, Blood) through pro- longed or repeated exposure if swallowed.
Precautionary Statements	:	Prevention: P260 Do not breathe mist or vapors.
		Response: P314 Get medical attention if you feel unwell.
		Disposal: P501 Dispose of contents and container to an approved waste disposal plant.

Other hazards

None known.

SECTION 3. COMPOSITION/INFORMATION ON INGREDIENTS

Substance / Mixture





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Components

Chemical name	CAS-No.	Concentration (% w/w)
Triclabendazole	68786-66-3	10
Silicon dioxide	7631-86-9	3
Benzyl alcohol	100-51-6	0.5
abamectin (combination of avermec-	71751-41-2	0.02
tin B1a and avermectin B1b) (ISO)		

SECTION 4. FIRST AID MEASURES

General advice	:	In the case of accident or if you feel unwell, seek medical advice immediately. When symptoms persist or in all cases of doubt seek medical advice.
If inhaled	:	If inhaled, remove to fresh air. Get medical attention if symptoms occur.
In case of skin contact	:	Wash with water and soap as a precaution. Get medical attention if symptoms occur.
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. Get medical attention if symptoms occur. Rinse mouth thoroughly with water.
Most important symptoms and effects, both acute and delayed	:	May cause damage to organs through prolonged or repeated exposure if swallowed.
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).
Notes to physician	:	Treat symptomatically and supportively.

SECTION 5. FIRE-FIGHTING MEASURES

Suitable extinguishing media		Alcohol-resistant foam Carbon dioxide (CO2) Dry chemical
Unsuitable extinguishing media	:	None known.
Specific hazards during fire fighting	:	Exposure to combustion products may be a hazard to health.
Hazardous combustion prod- ucts	:	Carbon oxides Nitrogen oxides (NOx) Metal oxides
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.
Special protective equipment for fire-fighters	:	

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SECTION 6. ACCIDENTAL RELEASE MEASURES

Personal precautions, protec- tive equipment and emer- gency procedures	:	Use personal protective equipment. Follow safe handling advice (see section 7) and personal protective equipment recommendations (see section 8).
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.
Methods and materials for containment and cleaning up	:	Soak up with inert absorbent material. For large spills, provide diking or other appropriate containment to keep material from spreading. If diked material can be pumped, store recovered material in appropriate container. Clean up remaining materials from spill with suitable absorbent. Local or national regulations may apply to releases and disposal of this material, as well as those materials and items employed in the cleanup of releases. You will need to determine which regulations are applicable. Sections 13 and 15 of this SDS provide information regarding certain local or national requirements.

SECTION 7. HANDLING AND STORAGE

Technical measures Local/Total ventilation Advice on safe handling	:	See Engineering measures under EXPOSURE CONTROLS/PERSONAL PROTECTION section. Use only with adequate ventilation. Do not breathe mist or vapors. Do not swallow. Avoid contact with eyes. Avoid prolonged or repeated contact with skin. Handle in accordance with good industrial hygiene and safety practice, based on the results of the workplace exposure assessment Take care to prevent spills, waste and minimize release to the environment.
Conditions for safe storage	:	Keep in properly labeled containers. Store in accordance with the particular national regulations.
Materials to avoid	:	Do not store with the following product types: Strong oxidizing agents Gases



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SECTION 8. EXPOSURE CONTROLS/PERSONAL PROTECTION

Ingredients with workplace control parameters

Components	CAS-No.	Value type (Form of exposure)	Control parame- ters / Permissible concentration	Basis
Triclabendazole	68786-66-3	TWA	30 µg/m3 (OEB 3)	Internal
	Further inform	ation: DSEN		
		Wipe limit	100 µg/100 cm2	Internal
Silicon dioxide	7631-86-9	TWA (Dust)	20 Million particles per cubic foot (Silica)	OSHA Z-3
		TWA (Dust)	80 mg/m3 / %SiO2 (Silica)	OSHA Z-3
		TWA	6 mg/m³ (Silica)	NIOSH REL
Benzyl alcohol	100-51-6	TWA	10 ppm	US WEEL
abamectin (combination of avermectin B1a and avermec- tin B1b) (ISO)	71751-41-2	TWA	15 μg/m3 (OEB 3)	Internal
		Wipe limit	150 µg/100 cm ²	Internal

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. 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 containment devices). Minimize open handling.
Personal protective equipment	
Respiratory protection :	General and local exhaust ventilation is recommended to maintain vapor exposures below recommended limits. Where concentrations are above recommended limits or are unknown, appropriate respiratory protection should be worn. Follow OSHA respirator regulations (29 CFR 1910.134) and use NIOSH/MSHA approved respirators. Protection provided by air purifying respirators against exposure to any hazardous chemical is limited. Use a positive pressure air supplied respirator if there is any potential for uncontrolled release, exposure levels are unknown, or any other circumstance where air purifying respirators may not provide adequate protection.
Hand protection	
Material :	Chemical-resistant gloves



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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 a	and body protection	: Work uniform Additional boo task being pe disposable su	or laboratory coat. dy garments should be used based upon the formed (e.g., sleevelets, apron, gauntlets, its) to avoid exposed skin surfaces. ate degowning techniques to remove potentially clothing.			
Hygiene measures		: If exposure to eye flushing s working place When using d Wash contam The effective engineering c appropriate de industrial hygi	chemical is likely during typical use, provide ystems and safety showers close to the			

SECTION 9. PHYSICAL AND CHEMICAL PROPERTIES

Appearance	:	suspension
Color	:	white
Odor	:	No data available
Odor Threshold	:	No data available
рН	:	5.0 - 7.0
Melting point/freezing point	:	< 41 °F / < 5 °C
Initial boiling point and boiling range	:	No data available
Flash point	:	No data available
Evaporation rate	:	No data available
Flammability (solid, gas)	:	Not applicable
Flammability (liquids)	:	No data available
Upper explosion limit / Upper flammability limit	:	No data available
Lower explosion limit / Lower	:	No data available

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	flamma	bility limit			
	Vapor pressure		:	No data available)
	Relative	e vapor density	:	No data available)
	Relative	e density	:	No data available)
	Density	,	:	1,050 - 1,080 g/c	m³ (68 °F / 20 °C)
	Solubili Wat	ty(ies) er solubility	:	soluble	
	Partition octanol	n coefficient: n-	:	Not applicable	
		ition temperature	:	No data available)
	Decom	position temperature	:	No data available)
	Viscosi Visc	ty osity, kinematic	:	No data available	9
	Explosi	ve properties	:	Not explosive	
	Oxidizir	ng properties	:	The substance o	r mixture is not classified as oxidizing.
	Molecu	lar weight	:	No data available)
	Particle Particle	characteristics size	:	Not applicable	

SECTION 10. STABILITY AND REACTIVITY

Reactivity	:	Not classified as a reactivity hazard.
Chemical stability	:	Stable under normal conditions.
Possibility of hazardous reac-	:	Can react with strong oxidizing agents.
tions		
Conditions to avoid	:	None known.
Incompatible materials	:	Oxidizing agents
Hazardous decomposition		No hazardous decomposition products are known.
products		

SECTION 11. TOXICOLOGICAL INFORMATION

Information on likely routes of exposure Inhalation Skin contact Ingestion Eye contact

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rsion	Revision Date: 07/06/2024	SDS Nu 534182	umber: 5-00013	Date of last issue: 04/06/2024 Date of first issue: 12/05/2019
Acute	e toxicity			
Not cl	assified based on ava	ailable infor	mation.	
<u>Produ</u>	<u>uct:</u>			
Acute	dermal toxicity			stimate: > 5,000 mg/kg ation method
Comp	oonents:			
Tricla	bendazole:			
Acute	oral toxicity	: LD5	50 (Mouse):	> 8,000 mg/kg
		LD5	50 (Rabbit):	206 mg/kg
Acute	inhalation toxicity	· 105	50 (Rat): > 0	5 mg/l
		Exp	osure time:	4 h
				e: dust/mist ne substance or mixture has no acute inhala
			toxicity	
Acute	dermal toxicity	: LD5	50 (Rat): > 4	,000 mg/kg
Silico	n dioxide:			
Acute	oral toxicity		50 (Rat): > 5	
		Met	hod: OECD	Test Guideline 401
Acute	inhalation toxicity	: LC5	50 (Rat): > 2	.08 mg/l
	-		osure time:	
				e: dust/mist ne substance or mixture has no acute inhala
			toxicity	
Acute	dermal toxicity	: LD5	50 (Rabbit):	> 5,000 mg/kg
Benzy	yl alcohol:			
-	oral toxicity	: LD5	50 (Rat): 1,6	20 mg/kg
Acute	inhalation toxicity	: LC5	50 (Rat): > 4	.178 mg/l
		Exp	osure time:	4 h
				e: dust/mist Test Guideline 403
		INIEL		
	•			d avermectin B1b) (ISO):
Acute	oral toxicity	: LD5	50 (Rat): 24	mg/kg
		LD5	60 (Mouse):	10 mg/kg
			.o (Monkey) nptoms: Dila	: 24 mg/kg tation of the pupil
Acute	inhalation toxicity	: LC5	50 (Rat): 0.0	23 mg/l

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			Exposure time: 4 Test atmosphere:	
Acu	te dermal toxicity	:	LD50 (Rat): 330 r	ng/kg
			LD50 (Rabbit): 2,	000 mg/kg
	n corrosion/irritation			
	classified based on avail	lable	information.	
	<u>nponents:</u>			
	clabendazole:		Dabbit	
Res	ecies sult	:	Rabbit Mild skin irritation	
Sili	con dioxide:			
	ecies	:	Rabbit	
Met Res	hod	:	OECD Test Guide No skin irritation	eline 404
T(C)	buit	•		
Ber	nzyl alcohol:			
	ecies	:	Rabbit	
Met Res	hod	:	OECD Test Guide No skin irritation	eline 404
1.00	Juit	·		
aba	mectin (combination of	f ave	rmectin B1a and a	avermectin B1b) (ISO):
	ecies	:	Rabbit	
Res	sult	:	No skin irritation	
Ser	ious eye damage/eye ir	ritati	on	
	classified based on avail			
<u>Cor</u>	nponents:			
Tric	clabendazole:			
Spe	ecies	:	Rabbit	
Res	sult	:	No eye irritation	
Sili	con dioxide:			
	ecies	:	Rabbit	
Res	sult hod	:	No eye irritation OECD Test Guide	alian 405
iviet		•		
Ber	nzyl alcohol:			
	ecies	:	Rabbit	
Res	sult hod	:	Irritation to eyes, OECD Test Guide	reversing within 21 days
IVIEI		•		

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abamectin (combination of avermectin B1a and avermectin B1b) (ISO): Species :: Result :: Result :: Mild eye irritation Skin sensitization Skin sensitization Not classified based on available information. Respiratory sensitization Not classified based on available information. Components: Triclabendazole: Result : Result : Not a skin sensitizer. Benzyl alcohol: Test Type : Totals of exposure : Syncies : Method : OCD Test Guideline 406 Result : Routes of exposure : Subtin contact Routes of exposure : Routes of exposure : Matimization Test Routes of exposure : Routes of exposure : Matimization Test Routes of exposure : Routes of exposure : Not classified based on ava	ersion .0	Revision Date: 07/06/2024	-	S Number: 41825-00013	Date of last issue: 04/06/2024 Date of first issue: 12/05/2019
Result : Mild eye irritation Respiratory or skin sensitization Skin sensitization Not classified based on available information. Respiratory sensitization Not classified based on available information. Components: Triclabendazole: Result : Not a skin sensitizer. Benzyl alcohol: Test Type : Maximization Test Routes of exposure : Skin contact Species : Guinea pig Method : OECD Test Guideline 406 Result : negative abamectin (combination of avermectin B1a and avermectin B1b) (ISO): Test Type : Maximization Test Routes of exposure : Skin contact Result : Not a skin sensitizer. Germ cell mutagenicity Not a skin sensitizer. Mot classified based on available information. Components: Triclabendazole: Genotoxicity in vitro Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Result: negative Silicon dioxide: Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Method: OECD Test	abam	ectin (combination	of ave	rmectin B1a an	d avermectin B1b) (ISO):
Result : Mild eye irritation Respiratory or skin sensitization Skin sensitization Not classified based on available information. Respiratory sensitization Not classified based on available information. Components: Triclabendazole: Result : Not a skin sensitizer. Benzyl alcohol: Test Type : Maximization Test Routes of exposure : Skin contact Species : Guinea pig Method : OECD Test Guideline 406 Result : negative abamectin (combination of avermectin B1a and avermectin B1b) (ISO): Test Type : Maximization Test Routes of exposure : Skin contact Result : Not a skin sensitizer. Germ cell mutagenicity Not a skin sensitizer. Mot classified based on available information. Components: Triclabendazole: Genotoxicity in vitro Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Result: negative Silicon dioxide: Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Method: OECD Test	Speci	es	:	Rabbit	
Skin sensitization Not classified based on available information. Respiratory sensitization Not classified based on available information. Components: Triclabendazole: Result : Result : Not a skin sensitizer. Benzyl alcohol: Test Type : Routes of exposure : Skin contact Species : OECD Test Guideline 406 Result : Result : abamectin (combination of avermectin B1a and avermectin B1b) (ISO): Test Type : Maximization Test Routes of exposure : Skin contact Result : Result : Not a skin sensitizer. Germ cell mutagenicity Not classified based on available information. Components: Triclabendazole: Genotoxicity in vitro : Test Type: DNA damage and repair, unscheduled DNA sy thesis in mammalian cells (in vitro) Result: negative Silicon dioxide:			:	Mild eye irritatio	on
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Not classified based on available information. Respiratory sensitization Not classified based on available information. Components: Triclabendazole: Result : Not a skin sensitizer. Benzyl alcohol: Test Type : Maximization Test Routes of exposure : Skin contact Species : Guinea pig Method : OECD Test Guideline 406 Result : negative abamectin (combination of avermectin B1a and avermectin B1b) (ISO): Test Type : Maximization Test Routes of exposure : Skin contact Result : negative abamectin (combination of avermectin B1a and avermectin B1b) (ISO): Test Type : Maximization Test Routes of exposure : Skin contact Result : Not a skin sensitizer. Gern cell mutagenicity Not classified based on available information. Components: Triclabendazole: Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Result: negative Silicon dioxide: Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Method: OECD Test Guideline 471 Result: negative Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Method: OECD Test Guideline 471 Result: negative Genotoxicity in vitro : Test Type: Mutagenicity (in vivo mammalian bone-marrow cytogenetic test, chromosomal analysis) Species: Rat	Resp	iratory or skin sens	itizatio	n	
Not classified based on available information. Respiratory sensitization Not classified based on available information. Components: Triclabendazole: Result : Not a skin sensitizer. Benzyl alcohol: Test Type : Maximization Test Routes of exposure : Skin contact Species : Guinea pig Method : OECD Test Guideline 406 Result : negative abamectin (combination of avermectin B1a and avermectin B1b) (ISO): Test Type : Maximization Test Routes of exposure : Skin contact Result : negative abamectin (combination of avermectin B1a and avermectin B1b) (ISO): Test Type : Maximization Test Routes of exposure : Skin contact Result : Not a skin sensitizer. Gern cell mutagenicity Not classified based on available information. Components: Triclabendazole: Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Result: negative Silicon dioxide: Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Method: OECD Test Guideline 471 Result: negative Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Method: OECD Test Guideline 471 Result: negative Genotoxicity in vitro : Test Type: Mutagenicity (in vivo mammalian bone-marrow cytogenetic test, chromosomal analysis) Species: Rat	Skin	sensitization			
Respiratory sensitization Not classified based on available information. Components: Triclabendazole: Result : Result : Test Type : Maximization Test Routes of exposure : Species : Guinea pig Method : Abamectin (combination of avermectin B1a and avermectin B1b) (ISO): Test Type : Result : negative abamectin (combination of avermectin B1a and avermectin B1b) (ISO): Test Type : Result : Routes of exposure : Skin contact Result : Not classified based on available information. Components: Triclabendazole: Genotoxicity in vitro : Test Type: DNA damage and repair, unscheduled DNA sy thesis in mammalian cells (in vitro) Result: negative Silicon dioxide: : Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Method: OECD Test Guideline 471 Result: negative			ailahle	information	
Not classified based on available information. Components: Triclabendazole: Result : Not a skin sensitizer. Benzyl alcohol: Test Type : Maximization Test Routes of exposure : Species : Guinea pig Method : Abamectin (combination of avermectin B1a and avermectin B1b) (ISO): Test Type : Result : Result : Routes of exposure : Skin contact Result : Result : Routes of exposure : Skin contact Result : Not classified based on available information. Components: Triclabendazole: Genotoxicity in vitro : Test Type: DNA damage and repair, unscheduled DNA sy thesis in mammalian cells (in vitro) Result: negative Silicon dioxide: Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Method: OECD Test Guideline 471 Result: negative					
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Routes of exposure : Skin contact Species : Guinea pig Method : OECD Test Guideline 406 Result : negative abamectin (combination of avermectin B1a and avermectin B1b) (ISO): Test Type : Maximization Test Routes of exposure : Skin contact Result : Not a skin sensitizer. Germ cell mutagenicity Not classified based on available information. Components: Triclabendazole: Genotoxicity in vitro : Test Type: DNA damage and repair, unscheduled DNA sy thesis in mammalian cells (in vitro) Result: negative Silicon dioxide: Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Method: OECD Test Guideline 471 Result: negative Silicon dioxide: : Genotoxicity in vitro : Test Type: Mutagenicity (in vivo mammalian bone-marrow cytogenetic test, chromosomal analysis) Species: Rat	Benzy	yl alcohol:			
Routes of exposure : Skin contact Species : Guinea pig Method : OECD Test Guideline 406 Result : negative abamectin (combination of avermectin B1a and avermectin B1b) (ISO): Test Type : Maximization Test Routes of exposure : Skin contact Result : Not a skin sensitizer. Germ cell mutagenicity Not classified based on available information. Components: Triclabendazole: Genotoxicity in vitro : Test Type: DNA damage and repair, unscheduled DNA sy thesis in mammalian cells (in vitro) Result: negative Silicon dioxide: Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Method: OECD Test Guideline 471 Result: negative Silicon dioxide: : Genotoxicity in vitro : Test Type: Mutagenicity (in vivo mammalian bone-marrow cytogenetic test, chromosomal analysis) Species: Rat	Test 1	- Fvpe	:	Maximization T	est
Species : Guinea pig Method : OECD Test Guideline 406 Result : negative abamectin (combination of avermectin B1a and avermectin B1b) (ISO): Test Type : Maximization Test Routes of exposure : Skin contact Result : Not a skin sensitizer. Germ cell mutagenicity Not a skin sensitizer. Not classified based on available information. Components: Triclabendazole: Genotoxicity in vitro Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Result: negative Silicon dioxide: Genotoxicity in vitro Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Method: OECD Test Guideline 471 Result: negative Silicon dioxide: : Test Type: Bacterial reverse mutation assay (AMES) Method: OECD Test Guideline 471 Result: negative Genotoxicity in vitro : Test Type: Mutagenicity (in vivo mammalian bone-marrow cytogenetic test, chromosomal analysis) Species: Rat					
Method : OECD Test Guideline 406 Result : negative abamectin (combination of avermectin B1a and avermectin B1b) (ISO): Test Type : Maximization Test Routes of exposure : Skin contact Result : Not a skin sensitizer. Germ cell mutagenicity Not a skin sensitizer. Not classified based on available information. Components: Triclabendazole: Genotoxicity in vitro Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Result: negative Silicon dioxide: Genotoxicity in vitro : Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Method: OECD Test Guideline 471 Result: negative Silicon dioxide: Genotoxicity in vitro : Genotoxicity in vitro : Test Type: Mutagenicity (in vivo mammalian bone-marrow cytogenetic test, chromosomal analysis) Species: Rat			:		
abamectin (combination of avermectin B1a and avermectin B1b) (ISO): Test Type : Maximization Test Routes of exposure : Skin contact Result : Not a skin sensitizer. Germ cell mutagenicity Not classified based on available information. Mot classified based on available information. Components: Triclabendazole: Genotoxicity in vitro Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Result: negative Test Type: DNA damage and repair, unscheduled DNA sy thesis in mammalian cells (in vitro) Result: negative Silicon dioxide: Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Method: OECD Test Guideline 471 Result: negative Genotoxicity in vitro : Test Type: Mutagenicity (in vivo mammalian bone-marrow cytogenetic test, chromosomal analysis) Species: Rat			:		ideline 406
Test Type : Maximization Test Routes of exposure : Skin contact Result : Not a skin sensitizer. Germ cell mutagenicity Not a skin sensitizer. Not classified based on available information. Components: Triclabendazole: Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Result: negative Test Type: DNA damage and repair, unscheduled DNA sy thesis in mammalian cells (in vitro) Result: negative Silicon dioxide: Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Method: OECD Test Guideline 471 Result: negative Genotoxicity in vitro : Test Type: Mutagenicity (in vivo mammalian bone-marrow cytogenetic test, chromosomal analysis) Species: Rat	Resul	t	:	negative	
Not classified based on available information. Components: Triclabendazole: Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Result: negative Test Type: DNA damage and repair, unscheduled DNA sy thesis in mammalian cells (in vitro) Result: negative Silicon dioxide: Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Method: OECD Test Guideline 471 Result: negative Genotoxicity in vitro : Test Type: Mutagenicity (in vivo mammalian bone-marrow cytogenetic test, chromosomal analysis) Species: Rat	Route	es of exposure	:	Skin contact	
Not classified based on available information. Components: Triclabendazole: Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Result: negative Test Type: DNA damage and repair, unscheduled DNA sy thesis in mammalian cells (in vitro) Result: negative Silicon dioxide: Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Method: OECD Test Guideline 471 Result: negative Genotoxicity in vitro : Test Type: Mutagenicity (in vivo mammalian bone-marrow cytogenetic test, chromosomal analysis) Species: Rat	Germ	cell mutagenicity			
Components: Triclabendazole: Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Result: negative Test Type: DNA damage and repair, unscheduled DNA sy thesis in mammalian cells (in vitro) Result: negative Silicon dioxide: Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Method: OECD Test Guideline 471 Result: negative Genotoxicity in vivo : Test Type: Mutagenicity (in vivo mammalian bone-marrow cytogenetic test, chromosomal analysis) Species: Rat			ailahle	information	
Triclabendazole: Genotoxicity in vitro: Test Type: Bacterial reverse mutation assay (AMES) Result: negativeSilicon dioxide: Genotoxicity in vitro: Test Type: DNA damage and repair, unscheduled DNA sy thesis in mammalian cells (in vitro) Result: negativeSilicon dioxide: Genotoxicity in vitro: Test Type: Bacterial reverse mutation assay (AMES) Method: OECD Test Guideline 471 Result: negativeGenotoxicity in vivo: Test Type: Mutagenicity (in vivo mammalian bone-marrow cytogenetic test, chromosomal analysis) Species: Rat			anabic	internation.	
Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Result: negative Test Type: DNA damage and repair, unscheduled DNA sy thesis in mammalian cells (in vitro) Result: negative Silicon dioxide: Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Method: OECD Test Guideline 471 Result: negative Genotoxicity in vivo : Test Type: Mutagenicity (in vivo mammalian bone-marrow cytogenetic test, chromosomal analysis) Species: Rat					
Result: negative Test Type: DNA damage and repair, unscheduled DNA sy thesis in mammalian cells (in vitro) Result: negative Silicon dioxide: Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Method: OECD Test Guideline 471 Result: negative Genotoxicity in vivo : : Test Type: Mutagenicity (in vivo mammalian bone-marrow cytogenetic test, chromosomal analysis) Species: Rat					
thesis in mammalian cells (in vitro) Result: negative Silicon dioxide: Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Method: OECD Test Guideline 471 Result: negative Genotoxicity in vivo : Test Type: Mutagenicity (in vivo mammalian bone-marrow cytogenetic test, chromosomal analysis) Species: Rat	Geno	toxicity in vitro	•		
thesis in mammalian cells (in vitro) Result: negative Silicon dioxide: Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Method: OECD Test Guideline 471 Result: negative Genotoxicity in vivo : Test Type: Mutagenicity (in vivo mammalian bone-marrow cytogenetic test, chromosomal analysis) Species: Rat				Test Type: DN	A damage and repair unscheduled DNA sy
Silicon dioxide: Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Method: OECD Test Guideline 471 Result: negative Genotoxicity in vivo : Test Type: Mutagenicity (in vivo mammalian bone-marrow cytogenetic test, chromosomal analysis) Species: Rat					
Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Method: OECD Test Guideline 471 Result: negative Genotoxicity in vivo : Test Type: Mutagenicity (in vivo mammalian bone-marrow cytogenetic test, chromosomal analysis) Species: Rat					
Genotoxicity in vitro : Test Type: Bacterial reverse mutation assay (AMES) Method: OECD Test Guideline 471 Result: negative Genotoxicity in vivo : Test Type: Mutagenicity (in vivo mammalian bone-marrow cytogenetic test, chromosomal analysis) Species: Rat					
Method: OECD Test Guideline 471 Result: negative Genotoxicity in vivo : Test Type: Mutagenicity (in vivo mammalian bone-marrow cytogenetic test, chromosomal analysis) Species: Rat					
Genotoxicity in vivo : Test Type: Mutagenicity (in vivo mammalian bone-marrow cytogenetic test, chromosomal analysis) Species: Rat	Geno	toxicity in vitro	:		
Genotoxicity in vivo : Test Type: Mutagenicity (in vivo mammalian bone-marrow cytogenetic test, chromosomal analysis) Species: Rat					
cytogenetic test, chromosomal analysis) Species: Rat				Result: negativ	e
cytogenetic test, chromosomal analysis) Species: Rat	Canad	tovioitu in vive		Toot Turner Mart	ogonicity (in vivo mommalica base more
Species: Rat	Geno		•		
					a, anomosomai anaiysisj
Application reduct ingestion				•	ute: Indestion



according to the OSHA Hazard Communication Standard

ersion 0	Revision Date: 07/06/2024	SDS Number:Date of last issue: 04/06/20245341825-00013Date of first issue: 12/05/2019
		Result: negative
Benz	yl alcohol:	
	toxicity in vitro	: Test Type: Bacterial reverse mutation assay (AMES) Result: negative
Geno	toxicity in vivo	 Test Type: Mammalian erythrocyte micronucleus test (in vivo cytogenetic assay) Species: Mouse Application Route: Intraperitoneal injection Result: negative
abam	ectin (combination	of avermectin B1a and avermectin B1b) (ISO):
Geno	toxicity in vitro	: Test Type: Bacterial reverse mutation assay (AMES) Result: negative
		Test Type: In vitro mammalian cell gene mutation test Test system: Chinese hamster lung cells Result: negative
		Test Type: Alkaline elution assay Result: negative
Geno	toxicity in vivo	: Test Type: Mutagenicity (in vivo mammalian bone-marrow cytogenetic test, chromosomal analysis) Species: Mouse Application Route: Intraperitoneal injection Result: negative
	nogenicity assified based on av	ailable information
	oonents:	
Tricla	bendazole:	
	cation Route sure time	: Mouse : Oral : 2 Years : negative
Speci		: Rat
	cation Route sure time t	: Oral : 2 Years : negative
Silico	n dioxide:	
	es cation Route sure time	: Rat : Ingestion : 103 weeks
Resul		: negative

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Benzy	/l alcohol:						
Speci	es	: Mouse					
	ation Route	: Ingestion					
Exposure time		: 103 weeks	•				
Metho		: OECD Test G	Guideline 451				
Resul	t	: negative					
abam	ectin (combinatior	n of avermectin B1a a	nd avermectin B1b) (ISO):				
Speci		: Rat					
	ation Route	: Oral					
	sure time	: 105 weeks					
Resul		: negative					
Resul	L	. negative					
Speci	es	: Mouse					
	ation Route	: Oral					
	sure time	: 93 weeks					
Resul		: negative					
IARC			esent at levels greater than or equal to 0.1% is or confirmed human carcinogen by IARC.				
OSHA			of this product present at levels greater than or equal to 0.1% is of regulated carcinogens.				
NTP			esent at levels greater than or equal to 0.1% is ated carcinogen by NTP.				
Repro	identified						
Repro Not cl	identified	as a known or anticipa					
Repro Not cl <u>Comp</u>	identified oductive toxicity assified based on a	as a known or anticipa					
Repro Not cl <u>Comp</u> Tricla	identified oductive toxicity assified based on a <u>oonents:</u> bendazole:	as a known or anticipa vailable information.	ited carcinogen by NTP.				
Repro Not cl <u>Comp</u> Tricla	identified oductive toxicity assified based on a ponents:	as a known or anticipa vailable information.	ated carcinogen by NTP.				
Repro Not cl <u>Comp</u> Tricla	identified oductive toxicity assified based on a <u>oonents:</u> bendazole:	as a known or anticipa vailable information. : Test Type: Fe Application R	ated carcinogen by NTP. ertility/early embryonic development oute: Oral				
Repro Not cl <u>Comp</u> Tricla	identified oductive toxicity assified based on a <u>oonents:</u> bendazole:	as a known or anticipa vailable information. : Test Type: Fe Application R Fertility: NOA	ated carcinogen by NTP.				
Repro Not cl <u>Comp</u> Tricla	identified oductive toxicity assified based on a <u>oonents:</u> bendazole:	as a known or anticipa vailable information. : Test Type: Fe Application R Fertility: NOA Result: No eff	ertility/early embryonic development oute: Oral EL: 50 mg/kg body weight fects on fertility.				
Repro Not cl <u>Comp</u> Tricla	identified oductive toxicity assified based on a <u>oonents:</u> bendazole:	as a known or anticipa vailable information. : Test Type: Fe Application R Fertility: NOA Result: No eff Test Type: Fe	ertility/early embryonic development oute: Oral EL: 50 mg/kg body weight fects on fertility.				
Repro Not cl <u>Comp</u> Tricla	identified oductive toxicity assified based on a <u>oonents:</u> bendazole:	as a known or anticipa vailable information. : Test Type: Fe Application R Fertility: NOA Result: No eff Test Type: Fe Application R	ertility/early embryonic development oute: Oral EL: 50 mg/kg body weight fects on fertility. ertility/early embryonic development oute: Oral				
Repro Not cl <u>Comp</u> Tricla	identified oductive toxicity assified based on a <u>oonents:</u> bendazole:	as a known or anticipa vailable information. : Test Type: Fe Application R Fertility: NOA Result: No eff Test Type: Fe Application R	ertility/early embryonic development oute: Oral EL: 50 mg/kg body weight fects on fertility.				
Repro Not cl <u>Comp</u> Tricla	identified oductive toxicity assified based on a <u>oonents:</u> bendazole:	as a known or anticipa vailable information. : Test Type: Fe Application R Fertility: NOA Result: No eff Test Type: Fe Application R Fertility: NOA	ertility/early embryonic development oute: Oral EL: 50 mg/kg body weight fects on fertility. ertility/early embryonic development oute: Oral				
Repro Not cl <u>Comp</u> Tricla	identified oductive toxicity assified based on a <u>oonents:</u> bendazole:	as a known or anticipa vailable information. : Test Type: Fe Application R Fertility: NOA Result: No eff Test Type: Fe Application R Fertility: NOA Result: No eff	ertility/early embryonic development oute: Oral EL: 50 mg/kg body weight fects on fertility. ertility/early embryonic development oute: Oral EL: 50 mg/kg body weight fects on fertility.				
Repro Not cl <u>Comp</u> Tricla	identified oductive toxicity assified based on a <u>oonents:</u> bendazole:	as a known or anticipa vailable information. : Test Type: Fe Application R Fertility: NOA Result: No eff Test Type: Fe Application R Fertility: NOA Result: No eff Test Type: Tw	ertility/early embryonic development oute: Oral EL: 50 mg/kg body weight fects on fertility. ertility/early embryonic development oute: Oral EL: 50 mg/kg body weight				
Repro Not cl <u>Comp</u> Tricla	identified oductive toxicity assified based on a <u>oonents:</u> bendazole:	as a known or anticipa vailable information. : Test Type: Fe Application R Fertility: NOA Result: No eff Test Type: Fe Application R Fertility: NOA Result: No eff Test Type: Tw Species: Rat	ertility/early embryonic development oute: Oral EL: 50 mg/kg body weight fects on fertility. ertility/early embryonic development oute: Oral EL: 50 mg/kg body weight fects on fertility.				
Repro Not cl <u>Comp</u> Tricla	identified oductive toxicity assified based on a <u>oonents:</u> bendazole:	as a known or anticipa vailable information. : Test Type: Fe Application R Fertility: NOA Result: No eff Test Type: Fe Application R Fertility: NOA Result: No eff Test Type: Tw Species: Rat Application R	ertility/early embryonic development oute: Oral EL: 50 mg/kg body weight fects on fertility. ertility/early embryonic development oute: Oral EL: 50 mg/kg body weight fects on fertility. wo-generation reproduction toxicity study oute: Oral				
Repro Not cl <u>Comp</u> Tricla	identified oductive toxicity assified based on a <u>oonents:</u> bendazole:	as a known or anticipa vailable information. : Test Type: Fe Application R Fertility: NOA Result: No eff Test Type: Fe Application R Fertility: NOA Result: No eff Test Type: Tw Species: Rat Application R	ertility/early embryonic development oute: Oral EL: 50 mg/kg body weight fects on fertility. ertility/early embryonic development oute: Oral EL: 50 mg/kg body weight fects on fertility.				
Repro Not cl Comp Tricla Effect	identified oductive toxicity assified based on a <u>ponents:</u> bendazole: s on fertility	as a known or anticipal vailable information. : Test Type: Fe Application R Fertility: NOA Result: No eff Test Type: Fe Application R Fertility: NOA Result: No eff Test Type: Tw Species: Rat Application R Fertility: NOA	ertility/early embryonic development oute: Oral EL: 50 mg/kg body weight fects on fertility. ertility/early embryonic development oute: Oral EL: 50 mg/kg body weight fects on fertility. vo-generation reproduction toxicity study oute: Oral EL: 5.5 mg/kg body weight				
Repro Not cl Comp Tricla Effect	identified oductive toxicity assified based on a <u>oonents:</u> bendazole:	as a known or anticipa vailable information. : Test Type: Fe Application R Fertility: NOA Result: No eff Test Type: Fe Application R Fertility: NOA Result: No eff Test Type: Tw Species: Rat Application R Fertility: NOA ent : Test Type: Er	ertility/early embryonic development oute: Oral EL: 50 mg/kg body weight fects on fertility. ertility/early embryonic development oute: Oral EL: 50 mg/kg body weight fects on fertility. wo-generation reproduction toxicity study oute: Oral				
Repro Not cl Comp Tricla Effect	identified oductive toxicity assified based on a <u>ponents:</u> bendazole: s on fertility	as a known or anticipal vailable information. : Test Type: Fe Application R Fertility: NOA Result: No eff Test Type: Fe Application R Fertility: NOA Result: No eff Test Type: Tw Species: Rat Application R Fertility: NOA Result: No eff Test Type: Tw Species: Rat	ertility/early embryonic development oute: Oral EL: 50 mg/kg body weight fects on fertility. ertility/early embryonic development oute: Oral EL: 50 mg/kg body weight fects on fertility. vo-generation reproduction toxicity study oute: Oral EL: 5.5 mg/kg body weight hbryo-fetal development				
Repro Not cl Comp Tricla Effect	identified oductive toxicity assified based on a <u>ponents:</u> bendazole: s on fertility	as a known or anticipal vailable information. : Test Type: Fe Application R Fertility: NOA Result: No eff Test Type: Fe Application R Fertility: NOA Result: No eff Test Type: Tw Species: Rat Application R Fertility: NOA	ertility/early embryonic development oute: Oral EL: 50 mg/kg body weight fects on fertility. ertility/early embryonic development oute: Oral EL: 50 mg/kg body weight fects on fertility. vo-generation reproduction toxicity study oute: Oral EL: 5.5 mg/kg body weight mbryo-fetal development oute: Oral				
Repro Not cl Comp Tricla Effect	identified oductive toxicity assified based on a <u>ponents:</u> bendazole: s on fertility	as a known or anticipal vailable information. : Test Type: Fe Application R Fertility: NOA Result: No eff Test Type: Fe Application R Fertility: NOA Result: No eff Test Type: Tw Species: Rat Application R Fertility: NOA	ertility/early embryonic development oute: Oral EL: 50 mg/kg body weight fects on fertility. ertility/early embryonic development oute: Oral EL: 50 mg/kg body weight fects on fertility. vo-generation reproduction toxicity study oute: Oral EL: 5.5 mg/kg body weight hbryo-fetal development				

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Versio 3.0	'n	Revision Date: 07/06/2024	-	9S Number: 41825-00013	Date of last issue: 04/06/2024 Date of first issue: 12/05/2019
				Species: Rat Application Route Developmental To Test Type: Embry Species: Rabbit Application Route Developmental To Result: Effects on Remarks: Materna Test Type: Embry Species: Rabbit Application Route Developmental To	oxicity: NOAEL: 50 mg/kg body weight vo-fetal development e: Oral oxicity: LOAEL: 10 mg/kg body weight ofetal development. al toxicity observed. vo-fetal development
		dioxide: on fetal development	:	Test Type: Embry Species: Rat Application Route Result: negative	vo-fetal development
	-	alcohol: on fertility	:	Species: Rat Application Route Result: negative	y/early embryonic development : Ingestion on data from similar materials
E	ffects	on fetal development	:	Test Type: Embry Species: Mouse Application Route Result: negative	vo-fetal development
		ctin (combination of a on fertility	ave :	Test Type: Fertilit Species: Rat, mal Application Route Result: Effects on Test Type: Two-g Species: Rat Application Route	e errility. eneration reproduction toxicity study of Oral Development: NOAEL: 0.12 mg/kg body
E	ffects	on fetal development	:		vo-fetal development

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Triclabendazole / Abamectin Formulation

ersion 0	Revision Date: 07/06/2024		DS Number: 41825-00013	Date of last issue: 04/06/2024 Date of first issue: 12/05/2019
			Developmental T Result: Cleft pala	Maternal: NOAEL: 0.05 mg/kg body weight oxicity: NOAEL: 0.2 mg/kg body weight
			Species: Rabbit Application Rout Developmental T Result: Cleft pala survival	yo-fetal development e: Oral oxicity: LOAEL: 2 mg/kg body weight ate, Teratogenic effects., Reduced embryoni se developmental effects were observed
			Test Type: Deve Species: Rat Application Rout Developmental T Result: Teratoge	e: Oral oxicity: LOAEL: 1.6 mg/kg body weight
Repro sessn	oductive toxicity - As- nent	:	fertility, based or	of adverse effects on sexual function and a animal experiments., Some evidence of on development, based on animal
	-single exposure lassified based on avai	lable	information.	
		ns (Li	ver, Blood) throug	h prolonged or repeated exposure if swal-
Com	oonents:			
Targe	abendazole: et Organs ssment	:	Liver, Blood May cause dama exposure.	age to organs through prolonged or repeate
abam	ectin (combination of	f ave	rmectin B1a and	avermectin B1b) (ISO):
	es of exposure	:	Ingestion	

Routes of exposure	:	Ingestion
Target Organs	:	Central nervous system
Assessment	:	Causes damage to organs through prolonged or repeated exposure.

Repeated dose toxicity

Components:

Triclabendazole:

Species	: Rat
NOAEL	: 6.6 mg/kg
LOAEL	: 69 mg/kg

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Expos	ation Route ure time t Organs	: Oral : 13 Weeks : Blood	
Expos	L	: Dog : 3.4 mg/kg : 37 mg/kg : Oral : 13 Weeks : Liver, Blood	
Expos		: Mouse : 29 mg/kg : Oral : 24 Months : Liver	
	L ation Route ure time	: Rat : 4 mg/kg : Oral : 24 Months : No significant	adverse effects were reported
Specie NOAE Applic		: Rat : 1.3 mg/m ³ : inhalation (dus : 13 Weeks	t/mist/fume)
Specie NOAE Applic	L ation Route ure time	: Rat : 1.072 mg/l : inhalation (dus : 28 Days : OECD Test Gi	
Specie NOAE Applic Expos	es L ation Route ure time t Organs	avermectin B1a ar : Rat : 1.5 mg/kg : Oral : 24 Months : Central nervou : Tremors, ataxi	
Specie NOAE Applic Expos	es L ation Route ure time t Organs	Mouse 4.0 mg/kg Oral 24 Months Central nervou Tremors, ataxi	ıs system
Specie	es	: Dog	

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rsion)	Revision Date: 07/06/2024		S Number: 41825-00013	Date of last issue: 04/06/2024 Date of first issue: 12/05/2019
NOAE LOAE		:	0.25 mg/kg 0.5 mg/kg	
Applic	cation Route	:	Oral	
	sure time	:	53 Weeks	
Targe Symp	t Organs	÷	Central nervous s Tremors, weight l	
Rema		:	mortality observe	
Speci		:	Monkey	
NOAE		÷	1.0 mg/kg Oral	
	cation Route sure time	:	14 Weeks	
	t Organs	:	Central nervous s	ystem
Aspir	ation toxicity			
	assified based on availa			
-	rience with human exp	osu	re	
	oonents:			
	bendazole:		.	
Inges	tion	:		ninal pain, Sweating, Headache, Nausea, a, Dizziness, Fatigue, Cough, Fever, pruriti
abam	ectin (combination of a	ave	rmectin B1a and a	vermectin B1b) (ISO):
Inges	tion	:	Symptoms: May of system effects, Sa	ause, Tremors, Diarrhea, central nervous alivation, tearing
CTION	12. ECOLOGICAL INFO	ORN	IATION	
Ecoto	oxicity			
<u>Com</u>	oonents:			
Silico	n dioxide:			
Toxici	ty to fish	:	LC50 (Danio reric Exposure time: 96 Method: OECD T	
	ty to daphnia and other ic invertebrates	:	EC50 (Daphnia m Exposure time: 24 Method: OECD T	
Toxici plants	ty to algae/aquatic	:	mg/l Exposure time: 72 Method: OECD T	
			NOEC (Desmode mg/l Exposure time: 72 Method: OECD T	



Versio 3.0	on	Revision Date: 07/06/2024		S Number: 41825-00013	Date of last issue: 04/06/2024 Date of first issue: 12/05/2019
				Remarks: Based of	on data from similar materials
E	Benzyl	alcohol:			
	Foxicity		:	LC50 (Pimephales Exposure time: 96	s promelas (fathead minnow)): 460 mg/l S h
		to daphnia and other invertebrates	:	EC50 (Daphnia m Exposure time: 48 Method: OECD Te	
	Foxicity plants	to algae/aquatic	:	EC50 (Pseudokiro mg/l Exposure time: 72 Method: OECD Te	
				NOEC (Pseudokir mg/l Exposure time: 72 Method: OECD Te	
а		to daphnia and other invertebrates (Chron- ty)	:	NOEC (Daphnia r Exposure time: 21 Method: OECD Te	
а	abameo	ctin (combination of a	avei	rmectin B1a and a	avermectin B1b) (ISO):
	Foxicity	•	:		hus mykiss (rainbow trout)): 3.2 µg/l
				LC50 (Lepomis m Exposure time: 96	acrochirus (Bluegill sunfish)): 9.6 µg/l ò h
				LC50 (Ictalurus pu Exposure time: 96	unctatus (channel catfish)): 24 μg/l δ h
				LC50 (Cyprinus ca Exposure time: 96	arpio (Carp)): 42 μg/l δ h
				LC50 (Cyprinodor Exposure time: 96	n variegatus (sheepshead minnow)): 15 μg/l δ h
		to daphnia and other invertebrates	:	EC50 (Americamy Exposure time: 96	
				EC50 (Daphnia m Exposure time: 48	agna (Water flea)): 0.34 μg/l } h
	Foxicity plants	to algae/aquatic	:	EC50 (Pseudokiro mg/l Exposure time: 72	chneriella subcapitata (green algae)): 100 2 h
	Foxicity city)	to fish (Chronic tox-	:	NOEC (Pimephale Exposure time: 32	es promelas (fathead minnow)): 0.52 μg/l 2 d





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	ty to daphnia and other	:		a magna (Water flea)): 0.03 µg/l
ic toxi	ic invertebrates (Chron-		Exposure time:	21 d
	- ,,		NOEC (Mysidor Exposure time:	osis bahia (opossum shrimp)): 0.0035 μg/ 28 d
Toxici	ty to microorganisms	:	EC50: > 1,000 r Exposure time: Test Type: Res	
Persi	stence and degradabili	ity		
Comp	oonents:			
Benzy	yl alcohol:			
Biode	gradability	:	Result: Readily Biodegradation: Exposure time:	92 - 96 %
abam	ectin (combination of	ave	rmectin B1a and	l avermectin B1b) (ISO):
Stabili	ity in water	:	Hydrolysis: 50 %	%(< 12 h)
Bioac	cumulative potential			
•				
Comp	oonents:			
Benzy Partiti	onents: yl alcohol: on coefficient: n- ol/water	:	log Pow: 1.05	
Benzy Partition	yl alcohol: on coefficient: n- ol/water		U U	l avermectin B1b) (ISO):
Benzy Partitie octane abam	yl alcohol: on coefficient: n- ol/water		rmectin B1a and	I avermectin B1b) (ISO): n factor (BCF): 52
Benzy Partitio octano abam Bioaco Partitio	yl alcohol: on coefficient: n- ol/water ectin (combination of a	ave :	rmectin B1a and	
Benzy Partitio octand Bioaco Partitio octand	yl alcohol: on coefficient: n- ol/water ectin (combination of a cumulation on coefficient: n-	ave :	rmectin B1a and Bioconcentratio	
Benzy Partitio octand Bioacd Partitio octand Mobil	yl alcohol: on coefficient: n- ol/water ectin (combination of a cumulation on coefficient: n- ol/water	ave :	rmectin B1a and Bioconcentratio	
Benzy Partitio octand Bioacd Partitio octand Mobil	yl alcohol: on coefficient: n- ol/water ectin (combination of a cumulation on coefficient: n- ol/water ity in soil ponents:	ave :	rmectin B1a and Bioconcentratio log Pow: 4	
Benzy Partitio octand Bioaco Partitio octand Mobil <u>Comp</u> abam	yl alcohol: on coefficient: n- ol/water ectin (combination of a cumulation on coefficient: n- ol/water ity in soil ponents:	ave : :	rmectin B1a and Bioconcentratio log Pow: 4 rmectin B1a and	n factor (BCF): 52
Benzy Partitio octand Bioacc Partitio octand Mobil <u>Comp</u> abam Distrik menta	yl alcohol: on coefficient: n- ol/water ectin (combination of a cumulation on coefficient: n- ol/water ity in soil <u>conents:</u> ectin (combination of a pution among environ-	ave : :	rmectin B1a and Bioconcentratio log Pow: 4 rmectin B1a and	n factor (BCF): 52

according to the OSHA Hazard Communication Standard



Version 3.0	Revision Date: 07/06/2024		Number: 325-00013	Date of last issue: 04/06/2024 Date of first issue: 12/05/2019
Conta	minated packaging	ha	andling site for	rs should be taken to an approved waste recycling or disposal. specified: Dispose of as unused product.
SECTION	14. TRANSPORT INFO	RMAT	ION	
Interr	national Regulations			
UNRT	ſDG			
UN nu	umber	: U	N 3082	
Prope	er shipping name	Ν	0.S.	TALLY HAZARDOUS SUBSTANCE, LIQUID
			1b) (ISO))	
Class		: 9	/ \ - //	
	ng group	: 111		
Label		: 9		
Enviro	onmentally hazardous	: уе	S	
IATA	DGR			
UN/IC			N 3082	
Prope	r shipping name	(;		hazardous substance, liquid, n.o.s. mbination of avermectin B1a and avermectin
Class		: 9	/ //	
Packi	ng group	: 111		
Label			iscellaneous	
aircra		: 96		
ger ai	ng instruction (passen- rcraft) onmentally hazardous	: 96		
	•	: ye	5	
	-Code			
	umber		N 3082	FALLY HAZARDOUS SUBSTANCE, LIQUIE
Fiope	er shipping name		0.S.	I ALLI HAZANDOUS SUBSTANUE, LIQUIL
		(a		nbination of avermectin B1a and avermectin
Class		: 9		
	ng group	: 111		
Label		: 9		
EmS Marin	e pollutant	: F· : ye	A, S-F es	
	port in bulk according			POL 73/78 and the IBC Code
-	estic regulation	- appile	.	
	-			
49 CF Un/IC	F R D/NA number	: U	N 3082	
-		_		

UN/ID/NA number	:	UN 3082
Proper shipping name	:	Environmentally hazardous substance, liquid, n.o.s.
		(abamectin (combination of avermectin B1a and avermectin
		B1b) (ISO))
Class	:	9



Triclabendazole / Abamectin Formulation

Version 3.0	Revision Date: 07/06/2024	SDS Number: 5341825-00013	Date of last issue: 04/06/2024 Date of first issue: 12/05/2019
Packir Labels	ng group	: III : CLASS 9	
ERG	-	: 171	
Marine	e pollutant	: yes(abamecti tin B1b) (ISO)	n (combination of avermectin B1a and avermec-
Rema	rks	: Above applies	s only to containers over 119 gallons or 450
		may be shipp	ground under DOT is non-regulated; however it ed per the applicable hazard classification to i-modal transport involving ICAO (IATA) or IMO.

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

SECTION 15. REGULATORY INFORMATION

CERCLA Reportable Quantity

This material does not contain any components with a CERCLA RQ.

SARA 304 Extremely Hazardous Substances Reportable Quantity

This material does not contain any components with a section 304 EHS RQ.

SARA 302 Extremely Hazardous Substances Threshold Planning Quantity

This material does not contain any components with a section 302 EHS TPQ.

SARA 311/312 Hazards	:	Specific target organ toxicity (single or repeated exposure)
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SARA 313

: This material does not contain any chemical components with known CAS numbers that exceed the threshold (De Minimis) reporting levels established by SARA Title III, Section 313.

US State Regulations

Pennsylvania Right To Know7732-18-5Water7732-18-5Triclabendazole68786-66-3Sodium citrate68-04-2Polyvinyl pyrrolidone9003-39-8Silicon dioxide7631-86-9

California Prop. 65

WARNING: This product can expose you to chemicals including abamectin (combination of avermectin B1a and avermectin B1b) (ISO), which is/are known to the State of California to cause birth defects or other reproductive harm. For more information go to www.P65Warnings.ca.gov.

California List of Hazardous Substances	
Polyvinyl pyrrolidone	9003-39-8
Silicon dioxide	7631-86-9
California Permissible Exposure Limits for Chemical Contaminants	
Silicon dioxide	7631-86-9

according to the OSHA Hazard Communication Standard



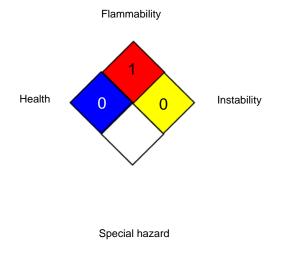
Triclabendazole / Abamectin Formulation

Version 3.0	Revision Date: 07/06/2024		OS Number: 41825-00013	Date of last issue: 04/06/2024 Date of first issue: 12/05/2019
The in AICS	•	oduct :	are reported in the not determined	ne following inventories:
DSL		:	not determined	
IECS	С	:	not determined	

SECTION 16. OTHER INFORMATION



NFPA 704:



HMIS® IV:



HMIS® ratings are based on a 0-4 rating scale, with 0 representing minimal hazards or risks, and 4 representing significant hazards or risks. The "*" represents a chronic hazard, while the "/" represents the absence of a chronic hazard.

Full text of other abbreviations

NIOSH REL OSHA Z-3	USA. NIOSH Recommended Exposure Limits USA. Occupational Exposure Limits (OSHA) - Table Z-3 Min- eral Dusts
US WEEL NIOSH REL / TWA	USA. Workplace Environmental Exposure Levels (WEEL) Time-weighted average concentration for up to a 10-hour workday during a 40-hour workweek
OSHA Z-3 / TWA US WEEL / TWA	8-hour time weighted average 8-hr TWA

AIIC - Australian Inventory of Industrial Chemicals; ASTM - American Society for the Testing of Materials; bw - Body weight; CERCLA - Comprehensive Environmental Response, Compensation, and Liability Act; CMR - Carcinogen, Mutagen or Reproductive Toxicant; DIN - Standard of the German Institute for Standardisation; DOT - Department of Transportation; DSL - Domestic Substances List (Canada); ECx - Concentration associated with x% response; EHS - Extremely Hazardous Substance; ELx - Loading rate associated with x% response; EmS - Emergency Schedule; ENCS - Existing and New Chemical Substances (Japan); ErCx - Concentration associated with x% response; EHS - Extremely Hazardous GLP - Good Laboratory Practice; HMIS - Hazardous Materials Identification System; IARC - International Agency for Research on Cancer; IATA - International Air Transport Association; IBC



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- 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; MSHA - Mine Safety and Health Administration; n.o.s. - Not Otherwise Specified; NFPA - National Fire Protection Association; NO(A)EC - No Observed (Adverse) Effect Concentration; NO(A)EL - No Observed (Adverse) Effect Level; NOELR - No Observable Effect Loading Rate; NTP - National Toxicology Program; 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; RCRA - Resource Conservation and Recovery Act; REACH - Regulation (EC) No 1907/2006 of the European Parliament and of the Council concerning the Registration, Evaluation, Authorisation and Restriction of Chemicals; RQ - Reportable Quantity; SADT - Self-Accelerating Decomposition Temperature; SARA - Superfund Amendments and Reauthorization Act; SDS - Safety Data Sheet; TCSI - Taiwan Chemical Substance Inventory; 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

Sources of key data used to	:	Internal technical data, data from raw material SDSs, OECD
compile the Material Safety Data Sheet		eChem Portal search results and European Chemicals Agency, http://echa.europa.eu/

Revision Date : 07/06/2024

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

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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