according to the OSHA Hazard Communication Standard



Ezetimibe Formulation

Version	Revision Date:	SDS Number:	Date of last issue: 03/20/2023
10.1	09/26/2023	23848-00022	Date of first issue: 10/21/2014

SECTION 1. IDENTIFICATION

Product name	:	Ezetimibe Formulation				
Manufacturer or supplier's details						
Company name of supplier	:	Organon & Co.				
Address	:	30 Hudson Street, 33nd floor				
		Jersey City, New Jersey, U.S.A 07302				
Telephone	:	1-551-430-6000				
Emergency telephone	:	1-215-631-6999				
E-mail address	:	EHSSTEWARD@organon.com				
Recommended use of the chemical and restrictions on use						

Recommended use	:	Pharmaceutical
Restrictions on use	:	Not applicable

SECTION 2. HAZARDS IDENTIFICATION

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

Combustible dust

GHS label elements

Signal Word	:	Warning
Hazard Statements	:	If small particles are generated during further processing, han- dling or by other means, may form combustible dust concentra- tions in air.

Other hazards

Dust contact with the eyes can lead to mechanical irritation.

SECTION 3. COMPOSITION/INFORMATION ON INGREDIENTS

Components		
Chemical name	CAS-No.	Concentration (% w/w)
Cellulose	9004-34-6	>= 20 - < 30
Ezetimibe	163222-33-1	>= 10 - < 20
Sodium n-dodecyl sulfate	151-21-3	>= 1 - < 5
Magnesium stearate	557-04-0	>= 1 - < 5
2-Pyrrolidone	616-45-5	>= 0.1 - < 1

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

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		advice.					
lf inha	led	,	: If inhaled, remove to fresh air.				
In case of skin contact		: In case of con Remove conta Get medical a Wash clothing	Get medical attention if symptoms occur. In case of contact, immediately flush skin with 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			If in eyes, rinse well with water. Get medical attention if irritation develops and persists.				
If swallowed		: If swallowed, I Get medical a	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			Dust contact with the eyes can lead to mechanical irritation.				
•	Protection of first-aiders		onders should pay attention to self-protection, ecommended personal protective equipment ntial for exposure exists (see section 8).				
Notes	to physician		natically and supportively.				

SECTION 5. FIRE-FIGHTING MEASURES

Suitable extinguishing media	:	Water spray Alcohol-resistant foam Carbon dioxide (CO2) Dry chemical
Unsuitable extinguishing media	:	None known.
Specific hazards during fire fighting	:	Avoid generating dust; fine dust dispersed in air in sufficient concentrations, and in the presence of an ignition source is a potential dust explosion hazard. Exposure to combustion products may be a hazard to health.
Hazardous combustion prod- ucts	:	Carbon oxides Nitrogen oxides (NOx) Fluorine compounds Sulfur oxides 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	:	In the event of fire, wear self-contained breathing apparatus. Use personal protective equipment.

SECTION 6. ACCIDENTAL RELEASE MEASURES

Personal precautions, protec- :	Use personal protective equipment.	
tive equipment and emer-	Follow safe handling advice (see section 7) and perso	onal

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gency	gency procedures		protective equipm	ent recommendations (see section 8).	
Environmental 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.		
Methods and materials for containment and cleaning up		:	container for disp Avoid dispersal of with compressed Dust deposits sho surfaces, as these released into the Local or national disposal of this m employed in the of determine which n Sections 13 and 1	dust in the air (i.e., clearing dust surfaces	

SECTION 7. HANDLING AND STORAGE

:	Static electricity may accumulate and ignite suspended dust causing an explosion. Provide adequate precautions, such as electrical grounding	
	and bonding, or inert atmospheres.	
:	Use only with adequate ventilation.	
:	Do not get on skin or clothing.	
	Do not breathe dust.	
	Do not swallow.	
	Avoid contact with eyes.	
	Handle in accordance with good industrial hygiene and safety	
	practice, based on the results of the workplace exposure	
	assessment	
	Minimize dust generation and accumulation.	
	Keep container closed when not in use.	
	Keep away from heat and sources of ignition.	
	Take precautionary measures against static discharges.	
	Take care to prevent spills, waste and minimize release to the environment.	
:	Keep in properly labeled containers.	
	Store in accordance with the particular national regulations.	
:	Do not store with the following product types: Strong oxidizing agents	
	: : : : : : : : : : : : : : : : : : : :	

SECTION 8. EXPOSURE CONTROLS/PERSONAL PROTECTION

Ingredients with workplace control parameters

inert or nuisance dust	50 Million particles per cubic foot
------------------------	-------------------------------------

according to the OSHA Hazard Communication Standard



ACGIH

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			Value type (Form of exposure): TWA (total dust) Basis: OSHA Z-3							
				15 mg/m³ Value type (Form of exposure): TWA (total dust) Basis: OSHA Z-3						
			5 mg/m³ Value type (Fo Basis: OSHA 2	5 mg/m³ Value type (Form of exposure): TWA (respirable fraction) Basis: OSHA Z-3						
					oot : TWA (respirable fra	ction)				
	Dust, nuisance dust and par- ticulates		10 mg/m³ Value type (Form of exposure): PEL (Total dust) Basis: CAL PEL							
			5 mg/m³ Value type (Fo Basis: CAL PE		: PEL (respirable dus	t fraction)				
C	Compo	nents	CAS-No.	Value type (Form of exposure)	Control parame- ters / Permissible concentration	Basis				
C	Cellulo	se	9004-34-6	TWA	10 mg/m ³	ACGIH				
				TWA (Res- pirable)	5 mg/m ³	NIOSH REL				
				TWA (total)	10 mg/m ³	NIOSH REL				
				TWA (total dust)	15 mg/m ³	OSHA Z-1				
				TWA (respir- able fraction)	5 mg/m ³	OSHA Z-1				
E	Ezetimi	ibe	163222-33-1	TWA	25 µg/m3 (OEB 3)	Internal				
				Wipe limit	250 µg/100 cm ²	Internal				
N	Magnes	sium stearate	557-04-0	TWA (Inhal- able particu- late matter)	10 mg/m ³	ACGIH				
			i	/ / /						

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 containment devices). Minimize open handling.

3 mg/m³

TWA (Res-

pirable particulate mat-

ter)

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Perso	onal protective equip	ment	
	iratory protection	maintain vapo concentration unknown, app Follow OSHA use NIOSH/M by air purifying hazardous ch supplied resp release, expo	ocal exhaust ventilation is recommended to or exposures below recommended limits. Where is are above recommended limits or are propriate respiratory protection should be worn. respirator regulations (29 CFR 1910.134) and ISHA approved respirators. Protection provided g respirators against exposure to any emical is limited. Use a positive pressure air irator if there is any potential for uncontrolled sure levels are unknown, or any other where air purifying respirators may not provide tection.
Hand	protection		
Ma	aterial	: Chemical-resi	istant gloves
	emarks protection	If the work en mists or aeros Wear a faces	ble gloving. glasses with side shields or goggles. vironment or activity involves dusty conditions, sols, wear the appropriate goggles. hield or other full face protection if there is a lirect contact to the face with dusts, mists, or
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 rformed (e.g., sleevelets, apron, gauntlets, its) to avoid exposed skin surfaces. ate degowning techniques to remove potentially clothing.
Hygie	ene 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 systems and safety showers close to the

SECTION 9. PHYSICAL AND CHEMICAL PROPERTIES

Appearance	:	powder
Color	:	off-white
Odor	:	No data available
Odor Threshold	:	No data available
рН	:	No data available
Melting point/freezing point	:	No data available

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	nitial boiling point and ange	boiling :	No data available	e
F	lash point	:	Not applicable	
E	vaporation rate	:	No data available	e
F	lammability (solid, gas	s) :	May form explos handling or othe	ive dust-air mixture during processing, r means.
F	lammability (liquids)	:	No data available	e
	Ipper explosion limit / ammability limit	Upper :	No data available	e
	ower explosion limit / ammability limit	Lower :	No data available	e
V	apor pressure	:	No data available	e
R	Relative vapor density	:	No data available	e
F	Relative density	:	No data available	e
C	Density	:	No data available	e
S	olubility(ies) Water solubility	:	No data available	e
	Partition coefficient: n-	:	No data available	e
-	utoignition temperatu	re :	No data available	e
D	Decomposition tempera	ature :	No data available	e
V	/iscosity Viscosity, kinematic	:	No data available	e
E	xplosive properties	:	Not explosive	
C	Oxidizing properties	:	The substance o	r mixture is not classified as oxidizing.
N	Iolecular weight	:	No data available	e
F	Particle size	:	No data available	e

SECTION 10. STABILITY AND REACTIVITY

Reactivity	:	Not classified as a reactivity hazard.
Chemical stability	:	Stable under normal conditions.

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Pos tions	sibility of hazardous reac- s	:	handling or other	ive dust-air mixture during processing, ⁻ means. rong oxidizing agents.
Con	Conditions to avoid		Heat, flames and Avoid dust forma	•
Haz	Incompatible materials Hazardous decomposition products		Oxidizing agents No hazardous decomposition products are known.	

SECTION 11. TOXICOLOGICAL INFORMATION

Information on likely routes Inhalation Skin contact Ingestion Eye contact	of	exposure
Acute toxicity		
Not classified based on availa	ble	information.
Product: Acute oral toxicity	:	Acute toxicity estimate: > 5,000 mg/kg Method: Calculation method
Components:		
Cellulose:		
Acute oral toxicity	:	LD50 (Rat): > 5,000 mg/kg
Acute inhalation toxicity	:	LC50 (Rat): > 5.8 mg/l Exposure time: 4 h Test atmosphere: dust/mist
Acute dermal toxicity	:	LD50 (Rabbit): > 2,000 mg/kg
Ezetimibe:		
Acute oral toxicity	:	LD50 (Rat): > 5,000 mg/kg
		LD50 (Mouse): > 5,000 mg/kg
		LD50 (Dog): > 3,000 mg/kg
Acute inhalation toxicity	:	Remarks: No data available
Acute dermal toxicity	:	Remarks: No data available
Acute toxicity (other routes of administration)	:	LD50 (Rat): > 2,000 mg/kg Application Route: Intraperitoneal
		LD50 (Mouse): > 1,000 - < 2,000 mg/kg Application Route: Intraperitoneal

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0		_		
	Im n-dodecyl sulfate			
Acute	oral toxicity	:	LD50 (Rat): 1, Method: OECI	200 mg/kg D Test Guideline 401
Acute dermal toxicity		:		2,000 mg/kg D Test Guideline 402 ed on data from similar materials
Magn	esium stearate:			
Acute	oral toxicity	:	LD50 (Rat): > 3	2,000 mg/kg
	-			D Test Guideline 423
				he substance or mixture has no acute oral to
			icity Remarks: Base	ed on data from similar materials
			Nemarks. DdS	
Acute	dermal toxicity	:	LD50 (Rabbit):	
			Remarks: Base	ed on data from similar materials
2-Pyr	rolidone:			
Acute	oral toxicity	:	LD50 (Rat): > 2	2,000 mg/kg
	-			D Test Guideline 401
				he substance or mixture has no acute oral to
			icity	
Acute	dermal toxicity	:	LD50 (Rabbit):	: > 2,000 mg/kg
			Method: OEC	D Test Guideline 402
			Assessment: T toxicity	he substance or mixture has no acute derma
Skin	corrosion/irritation			
	assified based on ava	ailable	information.	
Comp	oonents:			
Ezetir	nibe:			
Speci		:	Rabbit	
Resul	t	:	No skin irritatio	on
Sodiu	Im n-dodecyl sulfate	e :		
Speci		:	Rabbit	
Resul	t	:	Skin irritation	
Magn	esium stearate:			
-		:	Rabbit	
Speci			A I I I I I I I	
Speci Resul Rema		:	No skin irritatio	n from similar materials

2-Pyrrolidone:

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Specie Metho Resul	bd	:	Rabbit OECD Test Guide No skin irritation	eline 404		
	us eye damage/eye i assified based on ava					
<u>Comp</u>	oonents:					
Ezetir	nibe:					
Speci Resul			Rabbit No eye irritation			
Sodiu	ım n-dodecyl sulfate	:				
Specie Resul Metho	t	:	Rabbit Irreversible effects on the eye OECD Test Guideline 405			
Magn	esium stearate:					
Specie Resul Rema	t	:	Rabbit No eye irritation Based on data from similar materials			
2-Pvr	rolidone:					
Specie Resul	es		Rabbit Irritation to eyes, r	reversing within 7 days		
Respi	iratory or skin sensit	tizatior	1			
	sensitization assified based on ava	ailable in	nformation.			
_	iratory sensitization assified based on ava	ailable in	nformation.			
Comp	oonents:					
Ezetir Test T Specie Resul	Гуре es	:	Maximization Tes Guinea pig negative	t		
Sodiu	ım n-dodecyl sulfate	:				

Test Type : Maximization Test

root rypo	•	
Routes of exposure	:	Skin contact
Species	:	Guinea pig
Result	:	negative
Remarks	:	Based on data from similar materials

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ersion 0.1	Revision Date: 09/26/2023	SDS Nur 23848-00	
Magn	esium stearate:		
Test T Route Specie Metho Result Rema	s of exposure es id t	: Skin : Guin : OEC : nega	mization Test contact ea pig D Test Guideline 406 ative ed on data from similar materials
2-Pyrı	rolidone:		
Test T Route Specie Metho Result Rema	s of exposure es od	: Skin : Mous : OEC : nega	D Test Guideline 429
	cell mutagenicity assified based on av	ailable inform	nation
	oonents:		
Cellul	ose:		
Genot	oxicity in vitro		Type: Bacterial reverse mutation assay (AMES) ult: negative
			Type: In vitro mammalian cell gene mutation test ult: negative
Genot	oxicity in vivo	cytog Spec Appli	Type: Mammalian erythrocyte micronucleus test (in viv genetic assay) cies: Mouse ication Route: Ingestion ult: negative
Ezetir	nibe:		
Genot	oxicity in vitro	Meta	Type: Bacterial reverse mutation assay (AMES) abolic activation: with and without metabolic activation ult: negative
		Test	Type: Chromosomal aberration system: Human lymphocytes ult: negative
Genot	oxicity in vivo	Spec Cell t Appli	Type: Micronucleus test cies: Mouse type: Bone marrow ication Route: Oral ult: negative
Sodiu	m n-dodecyl sulfat	e :	
	oxicity in vitro		Type: Bacterial reverse mutation assay (AMES)

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rsion 1	Revision Date: 09/26/2023	SDS Number: 23848-00022	Date of last issue: 03/20/2023 Date of first issue: 10/21/2014
		Method: OE Result: nega	CD Test Guideline 471 ttive
		Test Type: Ir Result: nega	n vitro mammalian cell gene mutation test ative
Genot	toxicity in vivo	Species: Mo	Route: Ingestion
Magn	esium stearate:		
Genot	toxicity in vitro	Result: nega	
		Remarks: Ba	ased on data from similar materials
		Method: OE Result: nega	
			ased on data from similar materials
		Test Type: E Result: nega	Bacterial reverse mutation assay (AMES)
			ased on data from similar materials
2-Pyr	rolidone:		
Genot	toxicity in vitro	: Test Type: E Result: nega	Bacterial reverse mutation assay (AMES) ttive
			n vitro mammalian cell gene mutation test CD Test Guideline 476
			ased on data from similar materials
			Chromosome aberration test in vitro CD Test Guideline 473 Itive
Genot	toxicity in vivo	: Test Type: N cytogenetic a Species: Mo	
			Route: Intraperitoneal injection CD Test Guideline 474 ttive
	nogenicity assified based on ava	ailable information.	
Comp	oonents:		
Cellu	lose:		
Speci	es	: Rat	

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ersion .1	Revision Date: 09/26/2023	SDS Number: 23848-00022	Date of last issue: 03/20/2023 Date of first issue: 10/21/2014				
	ation Route ure time	: Ingestion : 72 weeks : negative					
Ezetin	nibe:						
	ation Route ure time	: Rat, female : oral (feed) : 104 weeks : negative					
	ation Route ure time	: Rat, male : oral (feed) : 104 weeks : negative					
	ation Route ure time	: Mouse : oral (feed) : 104 weeks : negative					
Sodiu	m n-dodecyl sulfate:						
	ation Route ure time d	: Rat : Ingestion : 2 Years : OECD Test G : negative : Based on data	uideline 453 a from similar materials				
2-Pyrr	olidone:						
Specie Applica	es ation Route ure time	: Mouse : Ingestion : 18 month(s) : negative : Based on data	a from similar materials				
IARC			sent at levels greater than or equal to 0.1% is or confirmed human carcinogen by IARC.				
OSHA		No component of this product present at levels greater than or equal to 0.1% is on OSHA's list of regulated carcinogens.					
NTP		No ingredient of this product present at levels greater than or equal to 0.1% is identified as a known or anticipated carcinogen by NTP.					
-	ductive toxicity assified based on avai	lable information.					
<u>Comp</u>	<u>onents:</u>						
Cellul	ose: s on fertility	: Test Type: On	e-generation reproduction toxicity study				

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				Species: Rat Application Route Result: negative	: Ingestion
E	ffects	on fetal development	:	Test Type: Fertilit Species: Rat Application Route Result: negative	y/early embryonic development : Ingestion
Е	zetim	ibe:			
E	ffects	on fertility	:	Species: Rat, mal Fertility: NOAEL:	y/early embryonic development e and female > 1,000 mg/kg body weight on fertility., No fetotoxicity.
E	ffects	on fetal development	:	Test Type: Develo Species: Rat Application Route Developmental To Result: No advers	: Oral oxicity: NOAEL: > 1,000 mg/kg body weight
				Test Type: Develo Species: Rabbit Application Route Developmental To Result: No advers	: Oral oxicity: NOAEL: > 1,000 mg/kg body weight
S	odiun	n n-dodecyl sulfate:			
E	ffects	on fertility	:	Species: Rat Application Route Method: OECD To Result: negative	
E	ffects	on fetal development	:	Species: Rat Application Route Result: negative	o-fetal development : Ingestion on data from similar materials
М	lagne	sium stearate:			
E	ffects	on fertility	:	reproduction/deve Species: Rat Application Route Method: OECD To Result: negative	
E	ffects	on fetal development	:	Test Type: Embry Species: Rat	o-fetal development

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ersion 0.1	Revision Date: 09/26/2023	SDS Number: 23848-00022	Date of last issue: 03/20/2023 Date of first issue: 10/21/2014
		Result: neg	Route: Ingestion ative Based on data from similar materials
2-Pvr	rolidone:		
	s on fertility	Species: Ra Application Result: pos	Route: Ingestion
Effect	s on fetal development	Species: Ra	Route: Ingestion
Repro sessn	oductive toxicity - As- nent	fertility, bas	ance of adverse effects on sexual function and ted on animal experiments., Clear evidence of ects on development, based on animal s.
	-single exposure assified based on availa	ble information	
	-repeated exposure		
STOT	-repeated exposure assified based on availa	ble information.	
STOT Not cl	• •	ble information.	
STOT Not cl Repe	assified based on availa	ble information.	
STOT Not cl Repe	assified based on availa ated dose toxicity conents:	ble information.	
STOT Not cl Repe <u>Comp</u> Cellu Speci	lassified based on availa ated dose toxicity <u>conents:</u> lose: es	: Rat	
STOT Not cl Repe <u>Comp</u> Cellu Speci NOAE	lassified based on availa ated dose toxicity <u>conents:</u> lose: es EL	: Rat : >= 9,000 m	ıg/kg
STOT Not cl Repe <u>Comp</u> Cellu Speci NOAE Applic	lassified based on availa ated dose toxicity <u>conents:</u> lose: es	: Rat	ıg/kg
STOT Not cl Repe <u>Comp</u> Cellu Speci NOAE Applic Expos	assified based on availa ated dose toxicity <u>conents:</u> lose: es EL cation Route	: Rat : >= 9,000 m : Ingestion	ıg/kg
STOT Not cl Repe Comp Cellu Speci NOAE Applic Expos	assified based on availa ated dose toxicity <u>ponents:</u> lose: es EL cation Route sure time mibe: es	: Rat : >= 9,000 m : Ingestion	ıg/kg
STOT Not cl Repe Comp Cellu Speci NOAE Applic Expos Speci NOAE	assified based on availa ated dose toxicity <u>ponents:</u> lose: es EL cation Route sure time mibe: es EL	: Rat : >= 9,000 m : Ingestion : 90 Days : Dog : 1,000 mg/k	
STOT Not cl Repe Comp Cellu Speci NOAE Applic Expos Speci NOAE Applic	assified based on availa ated dose toxicity <u>conents:</u> lose: es EL cation Route sure time mibe: es EL cation Route	: Rat : >= 9,000 m : Ingestion : 90 Days : Dog : 1,000 mg/k : Oral	
STOT Not cl Repe Comp Cellu Speci NOAE Applic Expos Speci NOAE Applic	assified based on availa ated dose toxicity <u>conents:</u> lose: es EL cation Route sure time mibe: es EL cation Route sure time	: Rat : >= 9,000 m : Ingestion : 90 Days : Dog : 1,000 mg/k : Oral : 90 d	
STOT Not cl Repe <u>Comp</u> Cellu Speci NOAE Applic Expos Rema	assified based on availa ated dose toxicity <u>bonents:</u> lose: es EL cation Route sure time mibe: es EL cation Route sure time arks	: Rat : >= 9,000 m : Ingestion : 90 Days : Dog : 1,000 mg/k : Oral : 90 d	g
STOT Not cl Repe Comp Cellu Speci NOAE Applic Expos Rema Speci NOAE	assified based on availa ated dose toxicity <u>ponents:</u> lose: es EL cation Route sure time mibe: es EL cation Route sure time arks es	 Rat >= 9,000 m Ingestion 90 Days Dog 1,000 mg/k Oral 90 d No significa Rat 1,500 mg/k 	g ant adverse effects were reported
STOT Not cl Repe Comp Cellu Speci NOAE Applic Expos Rema Speci NOAE Applic Expos Rema	assified based on availa ated dose toxicity <u>ponents:</u> lose: es EL cation Route sure time mibe: es EL cation Route sure time arks es EL cation Route sure time	 Rat >= 9,000 m Ingestion 90 Days Dog 1,000 mg/k Oral 90 d No significat Rat 1,500 mg/k Oral 	g ant adverse effects were reported
STOT Not cl Repe Comp Cellu Speci NOAE Applic Expos Rema Speci NOAE Applic Expos Rema	assified based on availa ated dose toxicity <u>ponents:</u> lose: es EL cation Route sure time mibe: es EL cation Route sure time arks es EL cation Route sure time arks	 Rat >= 9,000 m Ingestion 90 Days Dog 1,000 mg/k Oral 90 d No significat Rat 1,500 mg/k Oral 90 d 	g ant adverse effects were reported
STOT Not cl Repe Comp Cellu Speci NOAE Applic Expos Rema Speci NOAE Applic Expos Rema	assified based on availa ated dose toxicity <u>conents:</u> lose: es EL cation Route sure time mibe: es EL cation Route sure time arks es EL cation Route sure time arks	 Rat >= 9,000 m Ingestion 90 Days Dog 1,000 mg/k Oral 90 d No significat Rat 1,500 mg/k Oral 90 d 	g ant adverse effects were reported g
STOT Not cl Repe Comp Cellu Speci NOAE Applic Expos Rema Speci NOAE Applic Expos Rema Speci NOAE	assified based on availa ated dose toxicity <u>bonents:</u> lose: es EL cation Route sure time mibe: es EL cation Route sure time arks es EL cation Route sure time arks es EL cation Route sure time arks es EL cation Route sure time arks es	 Rat >= 9,000 m Ingestion 90 Days Dog 1,000 mg/k Oral 90 d No significat Rat 1,500 mg/k Oral 90 d No significat 	g ant adverse effects were reported g

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	iposi emar	ure time ks	:	90 d No significant adv	rerse effects were reported
NC Ap Ex		- ition Route ure time		Dog 300 mg/kg Oral 1 y No significant adv	verse effects were reported
So	odiur	n n-dodecyl sulfate:			
NC Ap Ex		- ition Route ure time		Rat 488 mg/kg Ingestion 90 Days Based on data fro	om similar materials
Ма	agne	sium stearate:			
NC Ap Ex		- ition Route ure time		Rat > 100 mg/kg Ingestion 90 Days Based on data fro	om similar materials
2-1	Pyrro	olidone:			
NC Ap Ex		- ition Route ure time		Rat 207 mg/kg Ingestion 3 Months OECD Test Guide	eline 408
As	spira	tion toxicity			
No	ot cla	ssified based on avail	able	information.	
<u>Cc</u>	ompo	onents:			
Ez	etim	ibe:			

Not applicable

Experience with human exposure

Components:

Ezetimibe:

Ingestion

: Symptoms: Headache, Nausea, Vomiting, Diarrhea, flatulence, muscle pain, upper respiratory tract infection, Back pain, joint pain

according to the OSHA Hazard Communication Standard



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SECTION	N 12. ECOLOGICAL INFO	ORI	IATION	
Ecot	toxicity			
Com	ponents:			
Cell	ulose:			
Toxi	city to fish	:	Exposure time: 48	ipes (Japanese medaka)): > 100 mg/l 3 h on data from similar materials
Ezet	imibe:			
Τοχί	city to fish	:	Exposure time: 96 Method: OECD T	
	city to daphnia and other atic invertebrates	:	Exposure time: 48 Method: OECD T	
Toxic plant	city to algae/aquatic ts	:	0.317 mg/l Exposure time: 96 Method: OECD T	
			mg/l Exposure time: 96 Method: OECD T	
Toxic icity)	city to fish (Chronic tox-	:	NOEC (Pimephal Exposure time: 33 Method: OECD T	
			Exposure time: 7	on variegatus (sheepshead minnow)): 4 mg/l d city at the limit of solubility.
aqua	city to daphnia and other atic invertebrates (Chron- xicity)	:	Exposure time: 2	nagna (Water flea)): 0.282 mg/l l d city at the limit of solubility.
Τοχί	city to microorganisms	:	EC50: > 4.4 mg/l Exposure time: 3 Test Type: Respin Method: OECD T Remarks: No toxi	ation inhibition
			NOEC: 4.4 mg/l	

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			Exposure time: 3 Test Type: Respir Method: OECD Te Remarks: No toxic	ation inhibition
Sodiu	m n-dodecyl sulfate:			
Toxicit	ty to fish	:	LC50 (Pimephales Exposure time: 96	s promelas (fathead minnow)): 29 mg/l ì h
	ty to daphnia and other cinvertebrates	:	EC50 (Ceriodaphi Exposure time: 48	nia dubia (water flea)): 5.55 mg/l 3 h
Toxicit plants	ty to algae/aquatic	:	ErC50 (Desmodes Exposure time: 72	smus subspicatus (green algae)): > 120 mg/l ? h
			NOEC (Desmode Exposure time: 72	smus subspicatus (green algae)): 30 mg/l ? h
Toxicit icity)	ty to fish (Chronic tox-	:	NOEC (Pimephale mg/l Exposure time: 42	es promelas (fathead minnow)): >= 1.357 2 d
	ty to daphnia and other c invertebrates (Chron-	:	NOEC (Ceriodaph Exposure time: 7	nnia dubia (water flea)): 0.88 mg/l d
	ty to microorganisms	:	EC50: 135 mg/l Exposure time: 3	h
Magne	esium stearate:			
Toxicit	ty to fish	:	Exposure time: 48 Method: DIN 3841	
	ty to daphnia and other c invertebrates	:	Exposure time: 47 Test substance: V Method: Directive	Vater Accommodated Fraction 67/548/EEC, Annex V, C.2. on data from similar materials
Toxicit plants	ty to algae/aquatic	:	mg/l Exposure time: 72 Test substance: V Method: OECD Te Remarks: Based o No toxicity at the l	Vater Accommodated Fraction est Guideline 201 on data from similar materials imit of solubility. irchneriella subcapitata (green algae)): > 1

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-	Revision Date: 09/26/2023	-	0S Number: 848-00022	Date of last issue: 03/20/2023 Date of first issue: 10/21/2014
			Method: OECD T	Vater Accommodated Fraction est Guideline 201 on data from similar materials
Toxicity	/ to microorganisms	:	Exposure time: 16 Test substance: V	nas putida): > 100 mg/l 5 h Vater Accommodated Fraction on data from similar materials
2-Pyrro	olidone:			
Toxicity	/ to fish	:	LC50 (Danio reric Exposure time: 96 Method: OECD Te	
	/ to daphnia and other invertebrates	:	EC50 (Daphnia m Exposure time: 48	agna (Water flea)): > 500 mg/l s h
Toxicity plants	/ to algae/aquatic	:	ErC50 (Desmode Exposure time: 72	smus subspicatus (green algae)): > 500 mg/l ? h
			EC10 (Desmodes Exposure time: 72	mus subspicatus (green algae)): 22.2 mg/l ? h
Toxicity	/ to microorganisms	:	EC50: > 1,000 mg Exposure time: 30 Method: OECD To) min
Dereier				
Persis	tence and degradabili	ity		
	tence and degradabilition	ity		
	onents:	ity		
<u>Compo</u> Cellulo	onents:	ity :	Result: Readily bi	odegradable.
<u>Compo</u> Cellulo	onents: ose: radability	ity :	Result: Readily bi	odegradable.
Compo Cellulo Biodeg Ezetim	onents: ose: radability	ity :	Result: Readily bi Result: Not readily Biodegradation: 6 Exposure time: 28	/ biodegradable. 5.8 %
Compo Cellulo Biodeg Ezetim Biodeg	onents: ose: radability ibe:	ity : :	Result: Not readily Biodegradation:	/ biodegradable. 5.8 % 5 d 4.5 d)
Compo Cellulo Biodeg Ezetim Biodeg Stability	onents: pse: radability ibe: radability y in water	ity :	Result: Not readily Biodegradation: 6 Exposure time: 28 Hydrolysis: 50 %(/ biodegradable. 5.8 % 5 d 4.5 d)
Compo Cellulo Biodeg Ezetim Biodeg Stability	onents: ose: radability ibe: radability	ity : :	Result: Not readily Biodegradation: 6 Exposure time: 28 Hydrolysis: 50 %(Method: OECD To Result: Readily bi Biodegradation: 9 Exposure time: 28	y biodegradable. 5.8 % 5 d 4.5 d) est Guideline 111 odegradable. 95 %
Compo Cellulo Biodeg Ezetim Biodeg Stability Sodiur Biodeg	onents: pse: radability ibe: radability y in water n n-dodecyl sulfate:	ity : :	Result: Not readily Biodegradation: 6 Exposure time: 28 Hydrolysis: 50 %(Method: OECD To Result: Readily bi Biodegradation: 9 Exposure time: 28	/ biodegradable. 5.8 % 5 d 4.5 d) est Guideline 111 odegradable. 95 %

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-	rrolidone: egradability	:	Result: Readily b Remarks: Based	iodegradable. on data from similar materials
Bioa	ccumulative potential			
<u>Com</u>	ponents:			
	i mibe: ccumulation	:	Bioconcentration Exposure time: 9	s macrochirus (Bluegill sunfish) factor (BCF): 173 7 d rest Guideline 305
	tion coefficient: n- nol/water	:	log Pow: 4.36	
Partit	um n-dodecyl sulfate: tion coefficient: n- nol/water	:	log Pow: 0.83	
Partit	nesium stearate: tion coefficient: n- nol/water	:	log Pow: > 4	
Partit	rrolidone: tion coefficient: n- nol/water	:		est Guideline 107
Mobi	ility in soil			
<u>Com</u>	ponents:			
Distri	i mibe: bution among environ- al compartments	:		est Guideline 106
	r adverse effects ata available			
SECTION	13. DISPOSAL CONSI	DEF	RATIONS	

Disposal methods		
Waste from residues	:	Dispose of in accordance with local regulations. Do not dispose of waste into sewer.
Contaminated packaging	:	Empty containers should be taken to an approved waste handling site for recycling or disposal. If not otherwise specified: Dispose of as unused product.

according to the OSHA Hazard Communication Standard



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SECTION 14. TRANSPORT INFORMATION

International Regulations

UNRTDG		
UN number	:	UN 3077
Proper shipping name	:	ENVIRONMENTALLY HAZARDOUS SUBSTANCE, SOLID, N.O.S. (Ezetimibe)
Class	:	9
Packing group	:	11
Labels	:	9
Environmentally hazardous	:	yes
IATA-DGR		
UN/ID No.	:	UN 3077
Proper shipping name	:	Environmentally hazardous substance, solid, n.o.s. (Ezetimibe)
Class	:	9
Packing group	:	
Labels	:	Miscellaneous
Packing instruction (cargo aircraft)	:	956
Packing instruction (passen- ger aircraft)	:	956
Environmentally hazardous	:	yes
IMDG-Code		
UN number	:	UN 3077
Proper shipping name	:	ENVIRONMENTALLY HAZARDOUS SUBSTANCE, SOLID, N.O.S.
-		(Ezetimibe)
Class	:	9
Packing group	:	
Labels	:	9
EmS Code	:	F-A, S-F
Marine pollutant	:	yes

Transport in bulk according to Annex II of MARPOL 73/78 and the IBC Code

Not applicable for product as supplied.

Domestic regulation

49 CFR		
UN/ID/NA number	:	UN 3077
Proper shipping name	:	Environmentally hazardous substance, solid, n.o.s. (Ezetimibe)
Class	:	9
Packing group	:	III
Labels	:	CLASS 9
ERG Code	:	171
Marine pollutant	:	yes(Ezetimibe)
Remarks	:	Above applies only to containers over 119 gallons or 450 liters.
		Shipment by ground under DOT is non-regulated; however it

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may be shipped per the applicable hazard classification to facilitate multi-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

: Combustible dust

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

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

SARA 311/312 Hazards

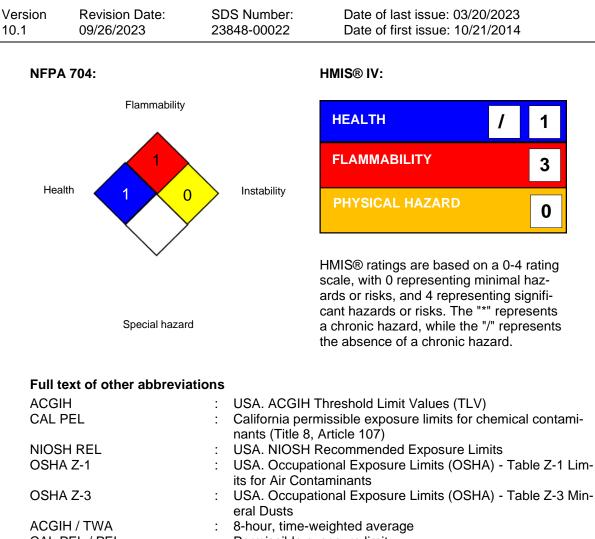
Pennsylvania Right To Know	N	
D-Glucose, 4-O-β-E Cellulose Ezetimibe Croscarmellose soc Polyvinyl pyrrolidon	64044-51-5 9004-34-6 163222-33-1 74811-65-7 9003-39-8	
California List of Hazardous	Substances	
Polyvinyl pyrrolidon	e	9003-39-8
California Permissible Exposure Limits for Chemical Contaminants		
Cellulose Magnesium stearat	e	9004-34-6 557-04-0
The ingredients of this prod	uct are reported in the following invento	ries:
AICS	: not determined	
DSL	: not determined	
IECSC	: not determined	

SECTION 16. OTHER INFORMATION

Further information

according to the OSHA Hazard Communication Standard

Ezetimibe Formulation



	•	e neur, une neighted average
CAL PEL / PEL	:	Permissible exposure limit
NIOSH REL / TWA	:	Time-weighted average concentration for up to a 10-hour
		workday during a 40-hour workweek
OSHA Z-1 / TWA	:	8-hour time weighted average
OSHA Z-3 / TWA	:	8-hour time weighted average

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% growth rate response; ERG - Emergency Response Guide; GHS - Globally Harmonized System; GLP - Good Laboratory Practice; HMIS - Hazardous Materials Identification System; 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 Pre-

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vention 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 compile the Material 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/

09/26/2023

Revision Date :

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.

US / Z8