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



Simvastatin Formulation

Version	Revision Date:	SDS Number:	Date of last issue: 26.09.2023
6.0	06.04.2024	24367-00023	Date of first issue: 21.10.2014

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

1.1	Product identifier Trade name	:	Simvastatin Formulation
1.2	Relevant identified uses of th Use of the Sub- stance/Mixture		ubstance or mixture and uses advised against Pharmaceutical
	Recommended restrictions on use	:	Not applicable
1.3	Details of the supplier of the	saf	ety data sheet
	Company	:	Organon & Co. 30 Hudson Street, 33nd floor 07302 Jersey City, New Jersey, U.S.A
	Telephone	:	+1-551-430-6000
	E-mail address of person	:	EHSSTEWARD@organon.com

1.4 Emergency telephone number

responsible for the SDS

+1-215-631-6999

SECTION 2: Hazards identification

2.1 Classification of the substance or mixture

Classification (REGULATION (EC) No 1272/2008)

Skin sensitisation, Category 1 Specific target organ toxicity - repeated exposure, Category 2 Long-term (chronic) aquatic hazard, Category 3 H317: May cause an allergic skin reaction.H373: May cause damage to organs through prolonged or repeated exposure.H412: Harmful to aquatic life with long lasting effects.

2.2 Label elements

Labelling (REGULATION (EC) No 1272/2008)

1

Hazard pictograms



Signal word

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



Simvastatin Formulation

Version 6.0	Revision Date: 06.04.2024	SDS Numbe 24367-00023	
Haza	rd statements	: H317 H373 H412	May cause an allergic skin reaction. May cause damage to organs through prolonged or repeated exposure. Harmful to aquatic life with long lasting effects.
Preca	autionary statements	: Preventic P260 P273 P280	Do not breathe dust. Avoid release to the environment. Wear protective gloves.
			e: Get medical advice/ attention if you feel unwell. 313 If skin irritation or rash occurs: Get medical advice/ attention. 364 Take off contaminated clothing and wash it before reuse.

Hazardous components which must be listed on the label:

Simvastatin

2.3 Other hazards

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

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

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

Dust contact with the eyes can lead to mechanical irritation. May form explosive dust-air mixture during processing, handling or other means.

SECTION 3: Composition/information on ingredients

3.2 Mixtures

Components

Components			
Chemical name	CAS-No.	Classification	Concentration
	EC-No.		(% w/w)
	Index-No.		
	Registration number		
Simvastatin	79902-63-9	Skin Irrit. 2; H315	>= 2.5 - < 10
		Skin Sens. 1; H317	
		STOT RE 1; H372	
		(Liver, muscle, optic	
		nerve, Eye)	
		Aquatic Chronic 2;	
		H411	

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



Simvastatin Formulation

Version 6.0	Revision Date: 06.04.2024	SDS Number: 24367-00023	Date of last issue: 26.09.2023 Date of first issue: 21.10.2014	
Citric	acid monohydrate	5949-29-1	Eye Irrit. 2; H319 STOT SE 3; H335	>= 1 - < 10
Subst	ances with a workplace	ce exposure limit :		
Ascor	bic acid	50-81-7 200-066-2		>= 1 - < 10

For explanation of abbreviations see section 16.

SECTION 4: First aid measures

4.1 Description of first aid measures General advice In the case of accident or if you feel unwell, seek medical ad-: vice immediately. When symptoms persist or in all cases of doubt seek medical advice. Protection of first-aiders First Aid responders should pay attention to self-protection, : and use the recommended personal protective equipment when the potential for exposure exists (see section 8). If inhaled If inhaled, remove to fresh air. : Get medical attention. In case of skin contact : 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, DO NOT induce vomiting, : Get medical attention if symptoms occur. Rinse mouth thoroughly with water. 4.2 Most important symptoms and effects, both acute and delayed Risks May cause an allergic skin reaction. : May cause damage to organs through prolonged or repeated exposure. Dust contact with the eyes can lead to mechanical irritation. 4.3 Indication of any immediate medical attention and special treatment needed Treatment : Treat symptomatically and supportively.

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



Simvastatin Formulation

Version	Revision Date:	SDS Number:	Date of last issue: 26.09.2023
6.0	06.04.2024	24367-00023	Date of first issue: 21.10.2014

SECTION 5: Firefighting measures

5.1 Extinguishing media

media

Suitable extinguishing media	:	Water spray Alcohol-resistant foam Carbon dioxide (CO2) Dry chemical
Unsuitable extinguishing	:	None known.

5.2 Special hazards arising from the substance or mixture

5.2 3	Special nazards arising from	the	e substance or mixture
	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
5.3 A	Advice for firefighters		
	Special protective equipment for firefighters	:	In the event of fire, wear self-contained breathing apparatus. Use personal protective equipment.
	Specific extinguishing meth- ods	:	Use extinguishing measures that are appropriate to local cir- cumstances and the surrounding environment. Use water spray to cool unopened containers. Remove undamaged containers from fire area if it is safe to do so.

Evacuate area.

SECTION 6: Accidental release measures

6.1 Personal precautions, protective equipment and emergency procedures

Personal precautions	:	Use personal protective equipment. Follow safe handling advice (see section 7) and personal pro- tective equipment recommendations (see section 8).
6.2 Environmental precautions Environmental precautions	:	Avoid release to the environment. Prevent further leakage or spillage if safe to do so. Retain and dispose of contaminated wash water.

Local authorities should be advised if significant spillages

6.3 Methods and material for containment and cleaning up

Methods for cleaning up	cleaning up :	Sweep up or vacuum up spillage and collect in suitable con-
		tainer for disposal.

cannot be contained.

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



Simvastatin Formulation

Version	Revision Date:	SDS Number:	Date of last issue: 26.09.2023
6.0	06.04.2024	24367-00023	Date of first issue: 21.10.2014
		with compress Dust deposits es, as these m leased into the Local or nation posal of this m employed in th mine which re Sections 13 an	al of dust in the air (i.e., clearing dust surfaces sed air). should not be allowed to accumulate on surfac- nay form an explosive mixture if they are re- e atmosphere in sufficient concentration. nal regulations may apply to releases and dis- naterial, as well as those materials and items ne cleanup of releases. You will need to deter- gulations are applicable. nd 15 of this SDS provide information regarding r national requirements.

6.4 Reference to other sections

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

SECTION 7: Handling and storage

7.1 Precautions for safe handling

Technical measures	 Static electricity may accumulate and ignite suspended dust causing an explosion. Provide adequate precautions, such as electrical grounding and handling, or inert atmospheres.
Local/Total ventilation	and bonding, or inert atmospheres.
Advice on safe handling	 Use only with adequate ventilation. Do not get on skin or clothing.
Advice on sale handling	Do not breathe dust.
	Do not swallow.
	Avoid contact with eyes.
	Wash skin thoroughly after handling.
	Handle in accordance with good industrial hygiene and safety
	practice, based on the results of the workplace exposure as-
	sessment
	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.
	Do not eat, drink or smoke when using this product.
	Take care to prevent spills, waste and minimize release to the environment.
Hygiene measures	: If exposure to chemical is likely during typical use, provide eye
	flushing systems and safety showers close to the working
	place. When using do not eat, drink or smoke. Contaminated
	work clothing should not be allowed out of the workplace.
	Wash contaminated clothing before re-use.
	The effective operation of a facility should include review of
	engineering controls, proper personal protective equipment,
	appropriate degowning and decontamination procedures,
	industrial hygiene monitoring, medical surveillance and the use of administrative controls.
7.2 Conditions for safe storage, i	ncluding any incompatibilities
Poquiromente for storage	Koop in properly labelled containers. Store in accordance with

Requirements for storage	:	Keep in properly labelled containers. Store in accordance with
areas and containers		the particular national regulations.

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



Simvastatin Formulation

Version 6.0	Revision Date: 06.04.2024	SDS Number: 24367-00023	Date of last issue: 26.09.2023 Date of first issue: 21.10.2014	
Advic	e on common storage	Strong oxidizi	substances and mixtures	
•	ic end use(s) fic use(s)	: No data availa	able	

SECTION 8: Exposure controls/personal protection

8.1 Control parameters

Occupational Exposure Limits

dusts non-specific	4 mg/m3 Value type (Form of exposure): OELV - 8 hrs (TWA) (Respirable dust) Basis: IE OEL
	10 mg/m3 Value type (Form of exposure): OELV - 8 hrs (TWA) (inhalable

Value type (Form of exposure): OELV - 8 hrs (TWA) (inhalable dust) Basis: IE OEL

Components	CAS-No.	Value type (Form of exposure)	Control parameters	Basis
Simvastatin	79902-63-9	TWA	25 µg/m3 (OEB 3)	Internal
	Further inform	nation: DSEN		
		Wipe limit	250 µg/100 cm ²	Internal
Starch	9005-25-8	OELV - 8 hrs (TWA) (Respira- ble dust)	4 mg/m3	IE OEL
		OELV - 8 hrs (TWA) (inhalable dust)	10 mg/m3	IE OEL
Cellulose	9004-34-6	OELV - 8 hrs (TWA)	10 mg/m3	IE OEL
Ascorbic acid	50-81-7	TWA	5000 µg/m3 (OEB 1)	Internal

Predicted No Effect Concentration (PNEC) according to Regulation (EC) No. 1907/2006:

Substance name	Environmental Compartment	Value
Citric acid monohydrate	Fresh water	0.44 mg/l
	Marine water	0.044 mg/l
	Sewage treatment plant	1000 mg/l
	Fresh water sediment	34.6 mg/kg dry weight (d.w.)
	Marine sediment	3.46 mg/kg dry

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



Simvastatin Formulation

Version 6.0	Revision Date: 06.04.2024	SDS Number: 24367-00023	Date of last issue: 26.09.2023 Date of first issue: 21.10.2014
11			weight (d.w.)
		Soil	33.1 mg/kg dry weight (d.w.)

8.2 Exposure controls

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.

Personal protective equipm	ent	
Eye/face protection	:	Wear safety glasses with side shields or goggles. If the work environment or activity involves dusty conditions, mists or aerosols, wear the appropriate goggles. Wear a faceshield or other full face protection if there is a potential for direct contact to the face with dusts, mists, or aerosols.
Hand protection		
Material	:	Chemical-resistant gloves
Remarks Skin and body protection	:	Consider double gloving. Work uniform or laboratory coat. Additional body garments should be used based upon the task being performed (e.g., sleevelets, apron, gauntlets, dis- posable suits) to avoid exposed skin surfaces. Use appropriate degowning techniques to remove potentially contaminated clothing.
Respiratory protection	:	If adequate local exhaust ventilation is not available or expo- sure assessment demonstrates exposures outside the rec- ommended guidelines, use respiratory protection. Equipment should conform to I.S. EN 143
Filter type	:	Particulates type (P)

SECTION 9: Physical and chemical properties

9.1 Information on basic physical and chemical properties

Physical state	:	powder
Colour	:	No data available
Odour	:	odourless
Odour Threshold	:	No data available
Melting point/freezing point	:	No data available
Initial boiling point and boiling range	:	No data available

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



Vers 6.0	sion	Revision Date: 06.04.2024		S Number: 367-00023	Date of last issue: 26.09.2023 Date of first issue: 21.10.2014
	Flamm	ability (solid, gas)	:	May form explos dling or other me	ive dust-air mixture during processing, han- ans.
	Flamm	ability (liquids)	:	No data availabl	9
		explosion limit / Upper ability limit	:	No data available	
		explosion limit / Lower ability limit	:	No data availabl	Ð
	Flash p	point	:	Not applicable	
	Auto-ig	nition temperature	:	No data availabl	9
	Decom	position temperature	:	No data availabl	9
	рН		:	No data available	9
	Viscos Viso	ity cosity, kinematic	:	Not applicable	
		ity(ies) ter solubility	:	No data availabl	2
	Partitic octano	n coefficient: n- I/water	:	Not applicable	
	Vapou	r pressure	:	Not applicable	
	Relativ	e density	:	No data available	9
	Densit	y	:	No data available	9
	Relativ	e vapour density	:	Not applicable	
		e characteristics ticle size	:	No data available	9
9.2	Other i	nformation			
	Explos	ives	:	Not explosive	
	Oxidizi	ng properties	:	The substance of	r mixture is not classified as oxidizing.
	Evapo	ration rate	:	Not applicable	

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



Simvastatin Formulation

Version	Revision Date:	SDS Number:	Date of last issue: 26.09.2023
6.0	06.04.2024	24367-00023	Date of first issue: 21.10.2014

SECTION 10: Stability and reactivity

10.1 Reactivity

Not classified as a reactivity hazard.

10.2 Chemical stability

Stable under normal conditions.

10.3 Possibility of hazardous reactions

Hazardous reactions	 May form explosive dust-air mixture during processing, han- dling or other means. Can react with strong oxidizing agents.
10.4 Conditions to avoid	
Conditions to avoid	: Heat, flames and sparks.

Avoid dust formation.

10.5 Incompatible materials

Materials to avoid : Oxidizing agents

10.6 Hazardous decomposition products

No hazardous decomposition products are known.

SECTION 11: Toxicological information

11.1 Information on hazard classes as defined in Regulation (EC) No 1272/2008

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

Acute toxicity

Not classified based on available information.

Components:

Simvastatin:

Acute oral toxicity	:	LD50 (Rat): 5,000 mg/kg LD50 (Mouse): 3,800 mg/kg
Citric acid monohydrate:		
Acute oral toxicity	:	LD50 (Mouse): 5,400 mg/kg
Acute dermal toxicity	:	LD50 (Rat): > 2,000 mg/kg Method: OECD Test Guideline 402 Assessment: The substance or mixture has no acute dermal toxicity

Ascorbic acid:

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



Simvastatin Formulation

Acute o	oral toxicity				
		:	LD50 (Rat): 11,90	00 mg/kg	
Skin co	orrosion/irritation	ilabla	information		
	ssified based on ava onents:	liable	inionnation.		
Simva			Dalla		
Specie: Remar		:	Rabbit Moderate skin irri	tation	
Citric a	acid monohydrate:				
Specie	S	:	Rabbit		
Result		•	No skin irritation		
Ascorb	bic acid:				
Specie	S	:	Rabbit		
Method	1	:	OECD Test Guide	eline 404	
Result		:	No skin irritation		
Compo Simvas Species Remar	S	:	Rabbit slight irritation		
Citric a	acid monohydrate:				
Specie	-	:	Rabbit		
Result		:	Irritation to eyes,	reversing within 21 days	
Ascort	bic acid:				
Specie		:	Rabbit		
Methoo Result	1	:	OECD Test Guide No eye irritation	eline 405	
Incount		•			
Respir	atory or skin sensit	isatio	on		
	ensitisation				
May cause an allergic skin reaction.					
Respir	atory sensitisation				
Not classified based on available information.					

Components:

Simvastatin:

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



Version 6.0	Revision Date: 06.04.2024		9S Number: 367-00023	Date of last issue: 26.09.2023 Date of first issue: 21.10.2014
Asses Result		:	Probability or evic positive	lence of skin sensitisation in humans
Test T	ure routes es		Maurer optimisati Skin contact Guinea pig negative	on test
	cell mutagenicity assified based on availa	able	information.	
<u>Comp</u>	onents:			
Simva	statin:			
Genote	oxicity in vitro	:	Test Type: Bacter Result: negative	rial reverse mutation assay (AMES)
			Test Type: Alkalir Result: negative	ne elution assay
			Test Type: Chrom Result: negative	nosomal aberration
			Test Type: In vitro Result: negative	o mammalian cell gene mutation test
Genote	oxicity in vivo	:	Test Type: Micror Species: Mouse Application Route Result: negative	
Germ sessm	cell mutagenicity- As- ent	:	Weight of evidend cell mutagen.	ce does not support classification as a germ
II Citric	acid monohydrate:			
	oxicity in vitro	:	Test Type: Bacter Result: negative	rial reverse mutation assay (AMES)
			Test Type: in vitro Result: positive	o micronucleus test
			Test Type: Bacter Result: negative	rial reverse mutation assay (AMES)
Genote	oxicity in vivo	:		enicity (in vivo mammalian bone-marrow chromosomal analysis) :: Ingestion

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



Version 6.0	Revision Date: 06.04.2024	SDS Number: 24367-00023	Date of last issue: 26.09.2023 Date of first issue: 21.10.2014
	Ascorbic acid: Genotoxicity in vitro		Bacterial reverse mutation assay (AMES) ative
		Test Type: I Result: neg	n vitro mammalian cell gene mutation test ative
		Test Type: (Result: neg	Chromosome aberration test in vitro ative
Geno	toxicity in vivo	cytogenetic Species: Mo	Route: Ingestion
	nogenicity		
	lassified based on ava ponents:	allable information.	
	astatin:		
Speci Applio Expos Targe	ies cation Route sure time et Organs or Type	: Mouse : Oral : < 92 weeks : Harderian g : Liver, Lungs : The significa	
Expo	cation Route sure time or Type	: Rat : Oral : 2 Years : Liver, Thyro : The significa	id ance of these findings for humans is not certain.
Asco	rbic acid:		
	cation Route sure time	: Mouse : Ingestion : 2 Years : negative	
-	oductive toxicity lassified based on ava	vilable information	
	ponents:		
	astatin:		
	ts on fertility	: Test Type: I Species: Ra Application Fertility: LO	it, male

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



Version 6.0	Revision Date: 06.04.2024		DS Number: 367-00023	Date of last issue: 26.09.2023 Date of first issue: 21.10.2014
Effect	Effects on foetal develop- : ment		Species: Rat Application Route Embryo-foetal tox	vo-foetal development e: Oral kicity: NOAEL: 25 mg/kg body weight genic effects, No adverse effects
			Species: Rabbit Application Route Embryo-foetal tox	vo-foetal development e: Oral kicity: NOAEL: 10 mg/kg body weight genic effects, No adverse effects
			Species: Rat Application Route Embryo-foetal tox Result: Teratoger	cicity: LOAEL: 60 mg/kg body weight
Citric	acid monohydrate:			
	ts on foetal develop-	:	Test Type: Embry Species: Rat Application Route Result: negative	vo-foetal development e: Ingestion
Asco	rbic acid:			
	ts on foetal develop-	:	Test Type: Embry Species: Rat Application Route Result: negative	vo-foetal development :: Ingestion
	Γ - single exposure lassified based on avail	able	information.	
Com	ponents:			
	acid monohydrate: ssment	:	May cause respira	atory irritation.
	F - repeated exposure cause damage to organ	s thr	ough prolonged or	repeated exposure.
	ponents:			
	astatin:			
Targe	et Organs ssment	:	Liver, muscle, opt Causes damage t exposure.	tic nerve, Eye to organs through prolonged or repeated

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



Simvastatin Formulation

Version 6.0	Revision Date: 06.04.2024	SDS Number: 24367-00023	Date of last issue: 26.09.2023 Date of first issue: 21.10.2014
Repe	eated dose toxicity		
<u>Com</u>	ponents:		
Simv	vastatin:		
Spec		: Rat	
NOA		: 5 mg/kg	
LOAE		: 30 mg/kg	
	cation Route	: Oral	
	sure time	: 14 - 104 Week	
large	et Organs	: Liver, l'estis, iv	lusculo-skeletal system, Eye
Spec		: Dog	
LOAE		: 10 mg/kg	
Appli	cation Route	: Oral	
	sure time	: 14 - 104 Week	-
large	et Organs	: Liver, Testis, E	уе
Spec	ies	: Rabbit	
NOA	EL	: 30 mg/kg	
LOAE		: 50 mg/kg	
	cation Route	: Oral	
Targe	et Organs	: Liver, Kidney	
Citric	c acid monohydrate:		
Spec	ies	: Rat	
NOA		: 4,000 mg/kg	
LOAE		: 8,000 mg/kg	
	cation Route	: Ingestion	
Expo	sure time	: 10 Days	
Asco	orbic acid:		
Spec	ies	: Rat, male	
NOA	EL	: >= 8,100 mg/kg	9
	cation Route	: Ingestion	
Expo	sure time	: 13 Weeks	
Asni	ration toxicity		
-	lassified based on ava	ailable information.	
	mation on other haz		

Endocrine disrupting properties

Product:

Assessment : The subst ered to ha

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

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



Simvastatin Formulation

Version	Revision Date:	SDS Number:	Date of last issue: 26.09.2023
6.0	06.04.2024	24367-00023	Date of first issue: 21.10.2014

Experience with human exposure

Components:

Simvastatin:

•	
Skin contact	: Remarks: May produce an allergic reaction.
Ingestion	: Target Organs: Liver
-	Symptoms: upper respiratory tract infection, Headache, Ab-
	dominal pain, constipation, Nausea
	Target Organs: Musculo-skeletal system
= =	

SECTION 12: Ecological information

12.1 Toxicity

Components:

Simvastatin:

Toxicity to fish	:	LC50 (Pimephales promelas (fathead minnow)): 2.91 mg/l Exposure time: 96 h Method: OECD Test Guideline 203
Toxicity to daphnia and other aquatic invertebrates	:	EC50 (Daphnia magna (Water flea)): 3.5 mg/l Exposure time: 48 h Method: OECD Test Guideline 202
Toxicity to algae/aquatic plants	:	EC50 (Pseudokirchneriella subcapitata (green algae)): > 25 mg/l Exposure time: 96 h
		NOEC (Pseudokirchneriella subcapitata (green algae)): 25 mg/l Exposure time: 96 h
Toxicity to microorganisms	:	EC50 : > 30 mg/l Exposure time: 3 h Test Type: Respiration inhibition Method: OECD Test Guideline 209
		NOEC : 21 mg/l Exposure time: 3 h Test Type: Respiration inhibition Method: OECD Test Guideline 209
Citric acid monohydrate:		
Toxicity to fish	:	LC50 (Pimephales promelas (fathead minnow)): > 100 mg/l Exposure time: 96 h
Toxicity to daphnia and other aquatic invertebrates	:	EC50 (Daphnia magna (Water flea)): 1,535 mg/l Exposure time: 24 h

Ascorbic acid:

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



Version 6.0	Revision Date: 06.04.2024		DS Number: 4367-00023	Date of last issue: 26.09.2023 Date of first issue: 21.10.2014
Toxic	ity to fish	:	Exposure time: 9	chus mykiss (rainbow trout)): 1,020 mg/l 96 h Fest Guideline 203
Toxic	ity to microorganisms	:	EC50 : 140 mg/l Exposure time: 1 Method: DIN 38	
12.2 Persi	istence and degradab	ility		
Com	ponents:			
Simv	astatin:			
	egradability	:	Result: rapidly de	egradable
Stabi	lity in water	:	Hydrolysis: 50 %	(3.2 d)
Citric	acid monohydrate:			
Biode	egradability	:	Result: Readily to Biodegradation: Exposure time: 2 Method: OECD	97 %
Asco	rbic acid:			
Biode	egradability	:	Result: Readily to Biodegradation: Exposure time: 5 Method: OECD	97 %
12.3 Bioa	ccumulative potential			
Com	ponents:			
Simv	astatin:			
Partit	ion coefficient: n- ol/water	:	log Pow: > 4.07	
Citric	acid monohydrate:			
	ion coefficient: n- ol/water	:	log Pow: -1.72	
	rbic acid:			
	ion coefficient: n- ol/water	:	log Pow: -1.85	
12.4 Mobi	lity in soil			
	ata available			
12.5 Resu	Ilts of PBT and vPvB a	asse	essment	
Prod				
Asses	ssment	:	This substance/r	nixture contains no components considered

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



Simvastatin Formulation

Version 6.0	Revision Date: 06.04.2024	SDS Number: 24367-00023	Date of last issue: 26.09.2023 Date of first issue: 21.10.2014			
		•	rsistent, bioaccumulative and toxic (PBT), or and very bioaccumulative (vPvB) at levels of			
12.6 Endo	12.6 Endocrine disrupting properties					
Produ	uct:					
Asses	ssment	ered to have e REACH Article	/mixture does not contain components consid- ndocrine disrupting properties according to 57(f) or Commission Delegated regulation 0 or Commission Regulation (EU) 2018/605 at or higher.			

12.7 Other adverse effects

No data available

SECTION 13: Disposal considerations

13.1 Waste treatment methods

Product	 Dispose of in accordance with local regulations. According to the European Waste Catalogue, Waste Codes are not product specific, but application specific. Waste codes should be assigned by the user, preferably in discussion with the waste disposal authorities. Do not dispose of waste into sewer.
Contaminated packaging	 Empty containers should be taken to an approved waste han- dling site for recycling or disposal. If not otherwise specified: Dispose of as unused product.

SECTION 14: Transport information

14.1 UN number or ID number

ADN	:	Not regulated as a dangerous good
ADR	:	Not regulated as a dangerous good
RID	:	Not regulated as a dangerous good
IMDG	:	Not regulated as a dangerous good
ΙΑΤΑ	:	Not regulated as a dangerous good
14.2 UN proper shipping name		
ADN	:	Not regulated as a dangerous good
ADN ADR	:	Not regulated as a dangerous good Not regulated as a dangerous good
	:	5 5 5
ADR	::	Not regulated as a dangerous good
ADR RID	::	Not regulated as a dangerous good Not regulated as a dangerous good

14.3 Transport hazard class(es)

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



Simvastatin Formulation

Version 6.0	Revision Date: 06.04.2024	SDS Number:Date of last issue: 26.09.202324367-00023Date of first issue: 21.10.2014	
ADN		: Not regulated as a dangerous good	
ADR		: Not regulated as a dangerous good	
RID		: Not regulated as a dangerous good	
IMDO	3	: Not regulated as a dangerous good	
ΙΑΤΑ	۱.	: Not regulated as a dangerous good	
14.4 Pack	king group		
ADN		: Not regulated as a dangerous good	
ADR		: Not regulated as a dangerous good	
RID		: Not regulated as a dangerous good	
IMDO	3	: Not regulated as a dangerous good	
ΙΑΤΑ	(Cargo)	: Not regulated as a dangerous good	
ΙΑΤΑ	(Passenger)	: Not regulated as a dangerous good	
-	ronmental hazards egulated as a dangero	is good	
-	c ial precautions for ι αpplicable	er	

14.7 Maritime transport in bulk according to IMO instruments

Remarks : Not applicable for product as supplied.

SECTION 15: Regulatory information

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

REACH - Restrictions on the manufacture, placing on the market and use of certain dangerous substances, mixtures and articles (Annex XVII)	:	Conditions of restriction for the fol- lowing entries should be considered: Number on list 75
		Substance(s) or mixture(s) are listed here according to their appearance in the regulation, irrespective of their use/purpose or the conditions of the restriction. Please refer to the condi- tions in corresponding Regulation to determine whether an entry is appli- cable to the placing on the market or not. If you intend to use this product as tattoo ink, please contact your ven- dor.
REACH - Candidate List of Substances of Very High Concern for Authorisation (Article 59).	:	Not applicable
Regulation (EC) No 1005/2009 on substances that de-	:	Not applicable

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



Simvastatin Formulation

Version 6.0	Revision Date: 06.04.2024	SDS Number: 24367-00023	Date of last issue: 26.09.2023 Date of first issue: 21.10.2014			
plete	the ozone layer					
Regul	ation (EU) 2019/1021	on persistent organic	pollu- : Not applicable			
	(recast) ation (EU) No 649/201	2 of the European Pa	arlia- : Not applicable			
•	Regulation (EU) No 649/2012 of the European Parlia- : Not applicable ment and the Council concerning the export and import					
	of dangerous chemicals					
REACH - List of substances subject to authorisation : Not applicable						
(Anne	(Annex XIV)					
Seveso III: Directive 2012/18/EU of the European Parliament and of the Council on the control of major-accident hazards involving dangerous substances.						

Not applicable

Other regulations:

Take note of Directive 94/33/EC on the protection of young people at work or stricter national regulations, where applicable.

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

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

15.2 Chemical safety assessment

A Chemical Safety Assessment has not been carried out.

SECTION 16: Other information

Other information	Items where changes have been made to the previous version are highlighted in the body of this document by two vertical lines.
Full text of H-Statements	
H315	Causes skin irritation.
H317	May cause an allergic skin reaction.
H319	Causes serious eye irritation.
H335	May cause respiratory irritation.
H372	: Causes damage to organs through prolonged or repeated
	exposure.
H411	: Toxic to aquatic life with long lasting effects.
Full text of other abbreviation	S
Aquatic Chronic	: Long-term (chronic) aquatic hazard
Eye Irrit.	Eye irritation
Skin Irrit.	Skin irritation
Skin Sens.	: Skin sensitisation
STOT RE	: Specific target organ toxicity - repeated exposure
STOT SE	: Specific target organ toxicity - single exposure
IE OEL	 Ireland. List of Chemical Agents and Carcinogens with Occu- pational Exposure Limit Values - Code of Practice, Schedule 1 and 2



Simvastatin Formulation

Version	Revision Date:	SDS Number:	Date of last issue: 26.09.2023
6.0	06.04.2024	24367-00023	Date of first issue: 21.10.2014

IE OEL / OELV - 8 hrs (TWA) : Occupational exposure limit value (8-hour reference period)

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

Further information

Aquatic Chronic 3

Sources of key data used to compile the Safety Data Sheet	:	Internal technical data, data from raw material SDSs, OECD eChem Portal search results and European Chemicals Agen- cy, http://echa.europa.eu/
Classification of the mixture	: :	Classification procedure:
Skin Sens. 1	H3 [,]	7 Calculation method
STOT RE 2	H37	3 Calculation method

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

Calculation method

H412

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



Simvastatin Formulation

Commission Regulation (EU) 2020/878

Version	Revision Date:	SDS Number:	Date of last issue: 26.09.2023
6.0	06.04.2024	24367-00023	Date of first issue: 21.10.2014

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.

IE / EN