

according to Regulation (EC) No. 1907/2006

Montelukast Tablet Formulation

Version	Revision Date:	SDS Number:	Date of last issue: 02.10.2020
2.16	09.04.2021	23075-00018	Date of first issue: 17.10.2014

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

1.1 Product identifier

Trade name : Montelukast Tablet Formulation

1.2 Relevant identified uses of the substance or mixture and uses advised against

Use of the Substance/Mixture : Pharmaceutical

1.3 Details of the supplier of the safety data sheet

Company	:	Organon & Co. Shotton Lane NE23 3JU Cramlington NU - Great Britain
Telephone	:	44 1 670 59 30 00
E-mail address of person responsible for the SDS	:	EHSSTEWARD@organon.com

1.4 Emergency telephone number

215-631-6999

SECTION 2: Hazards identification

2.1 Classification of the substance or mixture

Classification (REGULATION (EC) No 1272/2008)

Not a hazardous substance or mixture.

2.2 Label elements

Labelling (REGULATION (EC) No 1272/2008)

Not a hazardous substance or mixture.

Additional Labelling

EUH210 Safety data sheet available on request.

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.

Dust contact with the eyes can lead to mechanical irritation.

Contact with dust can cause mechanical irritation or drying of the skin.

May form explosive dust-air mixture during processing, handling or other means.



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SECTION 3: Composition/information on ingredients

3.2 Mixtures

Components

oomponenta			
Chemical name	CAS-No.	Classification	Concentration
	EC-No.		(% w/w)
	Index-No.		
	Registration number		
Montelukast	151767-02-1	Eye Irrit. 2; H319	>= 1 - < 10

For explanation of abbreviations see section 16.

SECTION 4: First aid measures

4.1 Description of first aid measures

4.1 Description of mist and measu	are	5		
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	:	Wash with water and soap. Get medical attention if symptoms occur.		
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 ar	nd e	effects, both acute and delayed		
Risks	:	Contact with dust can cause mechanical irritation or drying of the skin. 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.		



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SECTION 5: Firefighting measures

5.1 Extinguishing media

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

5.2 Special hazards arising from the substance or mixture

one opposition naeurao unioning nom		
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 Metal oxides
5.3 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

SECTION 6: Accidental release measures

6.1 Personal precautions, protective equipment and emergency procedures

SO.

Evacuate area.

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 cannot be contained.

6.3 Methods and material for containment and cleaning up

Methods for cleaning up	:	Sweep up or vacuum up spillage and collect in suitable con- tainer for disposal.
		Avoid dispersal of dust in the air (i.e., clearing dust surfaces

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		es, as these m leased into the Local or nation posal of this m employed in th mine which re Sections 13 an	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 bonding, or inert atmospheres.
Local/Total ventilation Advice on safe handling	 Use only with adequate ventilation. Do not breathe dust. 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 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. Take care to prevent spills, waste and minimize release to the
Hygiene measures	 environment. If exposure to chemical is likely during typical use, provide eye flushing systems and safety showers close to the working place. When using do not eat, drink or smoke. Wash 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, including any incompatibilities

Requirements for storage areas and containers	:	Keep in properly labelled containers. Store in accordance with the particular national regulations.
Advice on common storage	:	Do not store with the following product types: Strong oxidizing agents

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7.3 Specific end use(s)

Specific use(s)

: No data available

SECTION 8: Exposure controls/personal protection

8.1 Control parameters

Occupational Exposure Limits

Components	CAS-No.	Value type (Form	Control parameters	Basis		
		of exposure)				
Cellulose	9004-34-6	TWA (inhalable dust)	10 mg/m3	GB EH40		
	halable dust a sampling is un MDHS14/4 G ble, thoracic a hazardous to in air equal to mg.m-3 8-hou ject to COSH have been as the appropria of sizes. The entry into the depend on the fractions for li ble dust appro- and mouth du respiratory tra- to the gas exc material are g	are those fractions of indertaken in accorda eneral methods for s and inhalable aeroso health includes dust or greater than 10 n ur TWA of respirable H if people are exposision signed specific WEL te limits., Most indus behaviour, deposition human respiratory s e nature and size of mit-setting purposes painates to the fraction ring breathing and is act. Respirable dust a change region of the jiven in MDHS14/4.,	ses of these limits, respirabl airborne dust which will be ance with the methods descr ampling and gravimetric and ls., The COSHH definition of of any kind when present at ng.m-3 8-hour TWA of inhala dust. This means that any d sed to dust above these leve s and exposure to these mu trial dusts contain particles of n and fate of any particular p ystem, and the body respons the particle. HSE distinguish termed 'inhalable' and 'resp on of airborne material that a therefore available for depo approximates to the fraction lung. Fuller definitions and e Where dusts contain compo- elevant limits should be comp	collected when ibed in alysis or respira- f a substance a concentration able dust or 4 ust will be sub- ils. Some dusts st comply with of a wide range particle after se that it elicits, es two size irable'., Inhala- enters the nose osition in the that penetrates explanatory nents that have		
		TWA (Respirable 4 mg/m3 GB EH40 dust)				
	halable dust a sampling is un MDHS14/4 G ble, thoracic a hazardous to in air equal to mg.m-3 8-hou ject to COSHI have been as the appropria of sizes. The entry into the depend on the fractions for li ble dust appro	hation: For the purpo are those fractions of ndertaken in accorda eneral methods for s and inhalable aeroso health includes dust or greater than 10 n ur TWA of respirable H if people are exposisioned specific WEL te limits., Most indus behaviour, depositio human respiratory s e nature and size of mit-setting purposes oximates to the fracti	ses of these limits, respirabl airborne dust which will be ance with the methods descr ampling and gravimetric and ls., The COSHH definition of of any kind when present at ng.m-3 8-hour TWA of inhala dust. This means that any d sed to dust above these leve s and exposure to these mu trial dusts contain particles of n and fate of any particular p ystem, and the body respons the particle. HSE distinguish termed 'inhalable' and 'resp on of airborne material that of s therefore available for depo	collected when ibed in alysis or respira- a substance a concentration able dust or 4 ust will be sub- is. Some dusts st comply with of a wide range particle after se that it elicits, es two size irable'., Inhala- enters the nose		



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		to the gas exc material are g	hange region of the iven in MDHS14/4.	approximates to the fraction t lung. Fuller definitions and e Where dusts contain compor elevant limits should be comp 20 mg/m3	xplanatory nents that have
			dust)	20 mg/m3	GD EH40
Montol	ukot	sampling is ur MDHS14/4 Ge ble, thoracic a hazardous to in air equal to mg.m-3 8-hou ject to COSHI have been as the appropriat of sizes. The B entry into the depend on the fractions for life ble dust appro- and mouth du respiratory tra to the gas exc material are g their own assi	ndertaken in accord eneral methods for and inhalable aerose health includes dus or greater than 10 ar TWA of respirable d if people are expo- signed specific WE be limits., Most indus behaviour, deposition human respiratory se anature and size of mit-setting purpose oximates to the frac- ring breathing and ct. Respirable dust change region of the iven in MDHS14/4. gned WEL, all the r	f airborne dust which will be of ance with the methods descri sampling and gravimetric ana ols., The COSHH definition of t of any kind when present at mg.m-3 8-hour TWA of inhala e dust. This means that any du sed to dust above these leve _s and exposure to these must strial dusts contain particles of on and fate of any particular p system, and the body response the particle. HSE distinguishes the particle. HSE distinguishes therefore available for depo approximates to the fraction the e lung. Fuller definitions and e Where dusts contain compor elevant limits should be comp	bed in lysis or respira- a substance a concentration ble dust or 4 ust will be sub- ls. Some dusts st comply with f a wide range article after se that it elicits, es two size rable'., Inhala- enters the nose sition in the hat penetrates xplanatory nents that have blied with.
Montel	ukast	151767-02- 1	TWA	40 µg/m3 (OEB 3)	Internal
			Wipe limit	400 µg/100 cm ²	Internal

8.2 Exposure controls

Engineering measures

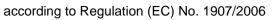
All engineering controls should be implemented by facility design and operated in accordance with GMP principles to protect products, workers, and the environment.

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 equipment

Eye 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	:	Consider double gloving.





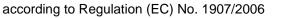
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d body protection	task being perform posable suits) to a	arments should be used based upon the med (e.g., sleevelets, apron, gauntlets, dis- avoid exposed skin surfaces. degowning techniques to remove potentially
	sure assessment ommended guide Equipment should	exhaust ventilation is not available or expo- demonstrates exposures outside the rec- lines, use respiratory protection. d conform to BS EN 143
l	d body protection tory protection	d body protection : Work uniform or I Additional body g task being perform posable suits) to Use appropriate of contaminated clo tory protection : If adequate local sure assessment ommended guide Equipment should

SECTION 9: Physical and chemical properties

9.1 Information on basic physical and chemical properties

Appearance Colour Odour Odour Threshold	:	tablet coloured odourless No data available
рН	:	No data available
Melting point/freezing point	:	No data available
Initial boiling point and boiling range	:	No data available
Flash point	:	Not applicable
Evaporation rate	:	No data available
Flammability (solid, gas)	:	May form explosive dust-air mixture during processing, han- dling or other means.
Flammability (liquids)	:	No data available
Upper explosion limit / Upper flammability limit	:	No data available
Lower explosion limit / Lower flammability limit	:	No data available
Vapour pressure	:	No data available
Relative vapour density	:	No data available
Relative density	:	No data available
Density	:	No data available
Solubility(ies) Water solubility Partition coefficient: n- octanol/water	:	No data available No data available





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Auto	-ignition temperature	:	No data available	e
Deco	omposition temperature	:	No data available	e
Visco V	osity iscosity, kinematic	:	No data available	e
Expl	osive properties	:	Not explosive	
Oxidizing properties		:	The substance of	r mixture is not classified as oxidizing.
	r information cular weight	:	No data availabl	e
Parti	cle size	:	No data availabl	9

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 re	actio	ns	
Hazardous reactions		May form explosive dust-air mixture during processing	han

Hazardous reactions	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.

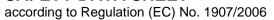
SECTION 11: Toxicological information

11.1 Information on toxicological effects

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

Acute toxicity

Not classified based on available information.





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<u>Comp</u>	oonents:				
Monte	elukast:				
Acute	oral toxicity	: LD50 (Rat):	> 5,000 mg/kg		
		LD50 (Mous	e): > 5,000 mg/kg		
Acute	inhalation toxicity	: Remarks: No	: Remarks: No data available		
Acute	dermal toxicity	: Remarks: No	o data available		
-	corrosion/irritation assified based on av	ailable information.			
	oonents:				
	elukast:				
Speci		: Rabbit			
Resul	t	: Mild skin irrit	ation		
	us eye damage/eye assified based on av				
<u>Comp</u>	oonents:				
Monte	elukast:				
Speci Resul		: Rabbit : Severe irrita	tion		
Respi	iratory or skin sens	tisation			
-	sensitisation				
_	assified based on av	ailable information.			
Respi	iratory sensitisatior				
Not cl	assified based on av	ailable information.			
<u>Comp</u>	oonents:				
Monte	elukast:				
Rema	ırks	: No data ava	ilable		
	cell mutagenicity assified based on av	ailable information.			
<u>Comp</u>	oonents:				
Monte	elukast:				
Geno	toxicity in vitro	: Test Type: E Result: nega	Bacterial reverse mutation assay (AMES) Itive		
			n vitro mammalian cell gene mutation test : Chinese hamster fibroblasts		

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		Result: negativ	e
			omosomal aberration hinese hamster ovary cells e
		Test Type: Alka Test system: ra Result: negativ	
Genc	otoxicity in vivo	: Test Type: Chr Species: Mous Cell type: Bone Application Rou Result: negativ	e marrow ute: Oral
	inogenicity lassified based on ava	ilable information.	
Com	ponents:		
Spec Appli	cation Route sure time	: Rat : Oral : 2 Years : negative	
Spec Appli	ies cation Route sure time	: Mouse : Oral : 92 weeks : negative	
-	oductive toxicity lassified based on ava	ilable information.	
Com	ponents:		
Mont	elukast:		
Effec	ts on fertility		nale
		Test Type: Fer Species: Rat, f Application Rou Fertility: LOAE Symptoms: Re	emale ute: Oral L: 200 mg/kg body weight
		Test Type: Fer Species: Rat, f Application Ro	emale
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Fertility: NOAEL: 100 mg/kg body weight Symptoms: Reduced fertility

STOT - single exposure

Not classified based on available information.

STOT - repeated exposure

Not classified based on available information.

Repeated dose toxicity

Components:

Montelukast:

Species NOAEL Application Route Exposure time Remarks	:	Monkey, male and female 150 - 300 mg/kg Oral 53 Weeks No significant adverse effects were reported
Species NOAEL Application Route Exposure time Remarks	:	Rat 50 mg/kg Oral 53 Weeks No significant adverse effects were reported
Species NOAEL Application Route Exposure time Remarks	:	Mouse 50 mg/kg Oral 14 Weeks No significant adverse effects were reported

Aspiration toxicity

Not classified based on available information.

Experience with human exposure

Components:

Montelukast:

Skin contact	: Remarks: May irritate skin.
Eye contact	: Symptoms: Severe irritation
Ingestion	: Symptoms: upper respiratory tract infection, pharyngitis, Headache, Cough, Abdominal pain, Diarrhoea, Fever

SECTION 12: Ecological information

12.1 Toxicity

Components:

Montelukast:

Toxicity to fish

: LC50 (Pimephales promelas (fathead minnow)): > 0.0778 mg/l Exposure time: 96 h

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				Method: OECD To Remarks: No toxio	est Guideline 203 city at the limit of solubility
		to daphnia and other invertebrates	:	Exposure time: 48 Method: OECD Te	
	Toxicity plants	to algae/aquatic	:	mg/l Exposure time: 72 Method: OECD To	
				mg/l Exposure time: 72 Method: OECD To	
-	Toxicity	to microorganisms	:	EC50 : > 100 mg/ Exposure time: 3 Test Type: Respir Method: OECD To Remarks: No toxio	h ation inhibition
	Toxicity icity)	to fish (Chronic tox-	:	Method: OECD To	2 d ales promelas (fathead minnow)
ä		to daphnia and other invertebrates (Chron- ty)	:		d magna (Water flea) city at the limit of solubility
12.2 Persistence and degradability					
<u>(</u>	Compo	nents:			
	Montel Biodegr	ukast: adability	:	Result: not rapidly Biodegradation: (Exposure time: 28)%
;	Stability	in water	:	Hydrolysis: 50 %(21.7 h)



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12.3 Bioa	12.3 Bioaccumulative potential						
<u>Con</u>	<u>iponents:</u>						
Montelukast: Partition coefficient: n- octanol/water		: log Pow: > 4	4.3				
	b ility in soil lata available						
12.5 Res	ults of PBT and vPvB a	ssessment					
Proc	duct:						
Asse	essment	to be either	nce/mixture contains no components considered persistent, bioaccumulative and toxic (PBT), or ent and very bioaccumulative (vPvB) at levels of ner.				
12.6 Oth	er adverse effects						
Proc	duct:						
End tial	ocrine disrupting poten-	ered to have REACH Art (EU) 2017/2	nce/mixture does not contain components consid- e endocrine disrupting properties according to cle 57(f) or Commission Delegated regulation 2100 or Commission Regulation (EU) 2018/605 at % or higher.				
SECTIO	N 13: Disposal consi	derations					

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.
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

Not regulated as a dangerous good

14.2 UN proper shipping name

Not regulated as a dangerous good

14.3 Transport hazard class(es)

Not regulated as a dangerous good

14.4 Packing group

Not regulated as a dangerous good

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14.5 Environmental hazards

Not regulated as a dangerous good

14.6 Special precautions for user

Not applicable

14.7 Transport in bulk according to Annex II of Marpol and the IBC Code

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, preparations and articles (Annex XVII)	:	Not applicable
REACH - Candidate List of Substances of Very High Concern for Authorisation (Article 59).	:	Not applicable
REACH - List of substances subject to authorisation (Annex XIV)	:	Not applicable
Regulation (EC) No 1005/2009 on substances that deplete the ozone layer	:	Not applicable
Regulation (EU) 2019/1021 on persistent organic pollu- tants (recast)	:	Not applicable
Regulation (EC) No 649/2012 of the European Parlia- ment and the Council concerning the export and import of dangerous chemicals	:	Not applicable
Seveso III: Directive 2012/18/EU of the European Parlian major-accident hazards involving dangerous substances.		t and of the Council on the control of

Not 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					
H319	:	Causes serious eye irritation.			
Full text of other abbreviations					
Eye Irrit.	:	Eye irritation			



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	40 40 / TWA 40 / STEL	: L	ong-term exp	L - Workplace Exposure Limits osure limit (8-hour TWA reference period) oosure limit (15-minute reference period)

ADN - European Agreement concerning the International Carriage of Dangerous Goods by Inland Waterways; ADR - European 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; TRGS -Technical Rule for Hazardous Substances; TSCA - Toxic Substances Control Act (United States); UN - United Nations; vPvB - Very Persistent and Very Bioaccumulative

Further information

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

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Version	Revision Date:	SDS Number:	Date of last issue: 02.10.2020
2.16	09.04.2021	23075-00018	Date of first issue: 17.10.2014