

Rizatriptan Formulation

Version 2.10	Revision Date: 10.10.2020		S Number: 2448-00013	Date of last issue: 23.03.2020 Date of first issue: 10.12.2015	
SECTION	1. PRODUCT AND COI	MPA		TION	
Produ	uct name	:	Rizatriptan Forr	nulation	
Manu	afacturer or supplier's o	deta	ils		
Com	bany	:	Organon & Co.		
Addre	Address		30 Hudson Street, 33nd floor Jersey City, New Jersey, U.S.A 07302		
Telep	phone	:	551-430-6000		
Emer	gency telephone numbe	r:	215-631-6999		
E-ma	il address	:	EHSSTEWARD	@organon.com	
Reco	mmended use of the c	hem	ical and restrict	ions on use	
Reco	mmended use	:	Pharmaceutical		
GHS Repre	2. HAZARDS IDENTIFI	:	Category 2	rdio-vascular system)	
	ated exposure (Oral)				
GHS	label elements				
Haza	rd pictograms	:			
Signa	al word	:	Warning		
Haza	rd statements	:	H373 May caus	ed of damaging the unborn child. e damage to organs (Cardio-vascular system) jed or repeated exposure if swallowed.	
Preca	autionary statements	:	P202 Do not ha and understood P260 Do not bre		

P281 Use personal protective equipment as required.

Response:

P308 + P313 IF exposed or concerned: Get medical advice/ attention.

Storage:





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P405 Store locked up.

Disposal:

P501 Dispose of contents/ container to an approved waste disposal plant.

Other hazards which do not result in classification

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.

SECTION 3. COMPOSITION/INFORMATION ON INGREDIENTS

Substance / Mixture : Mixture

Components

Chemical name	CAS-No.	Concentration (% w/w)
Cellulose	9004-34-6	>= 30 -< 60
Starch	9005-25-8	>= 10 -< 30
Rizatriptan	145202-66-0	>= 3 -< 10

SECTION 4. 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.
If inhaled	:	If inhaled, remove to fresh air. Get medical attention.
In case of skin contact	:	In case of contact, immediately flush skin with soap and plenty of water. Remove contaminated clothing and shoes. Get medical attention. Wash clothing before reuse. Thoroughly clean shoes before reuse.
In case of eye contact	:	
If swallowed	:	If swallowed, DO NOT induce vomiting. Get medical attention. Rinse mouth thoroughly with water.
Most important symptoms and effects, both acute and delayed	:	
Protection of first-aiders	:	Dust contact with the eyes can lead to mechanical irritation. First Aid responders should pay attention to self-protection, and use the recommended personal protective equipment when the potential for exposure exists (see section 8).
Notes to physician	:	Treat symptomatically and supportively.

SECTION 5. FIREFIGHTING MEASURES



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Suitable extinguishing media		:	Water spray Alcohol-resistant Carbon dioxide (Dry chemical	
Unsu media	itable extinguishing	:	None known.	
	Specific hazards during fire- fighting		concentrations, a potential dust exp	dust; fine dust dispersed in air in sufficient and in the presence of an ignition source is a plosion hazard. bustion products may be a hazard to health.
Haza ucts	rdous combustion prod-	:	Carbon oxides	
Speci ods	Specific extinguishing meth- ods		cumstances and Use water spray	g measures that are appropriate to local cir- the surrounding environment. to cool unopened containers. Iged containers from fire area if it is safe to d
	Special protective equipment for firefighters		In the event of fir	e, wear self-contained breathing apparatus. stective equipment.
ECTION	6. ACCIDENTAL RELE	AS	E MEASURES	
tive e	onal precautions, protec- quipment and emer- / procedures	:	Follow safe hand	tective equipment. lling advice (see section 7) and personal pro t recommendations (see section 8).
Envir	Environmental precautions		Retain and dispo	eakage or spillage if safe to do so. se of contaminated wash water. should be advised if significant spillages
	ods and materials for inment and cleaning up	:	tainer for disposa Avoid dispersal of with compressed Dust deposits she es, as these may leased into the at Local or national posal of this mate employed in the of mine which regul	of dust in the air (i.e., clearing dust surfaces

SECTION 7. HANDLING AND STORAGE

Technical measures	: Static electricity may accumulate and ignite suspended dust
	causing an explosion.
	Provide adequate precautions, such as electrical grounding

certain local or national requirements.

Sections 13 and 15 of this SDS provide information regarding



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Local/Total ventilation Advice on safe handling		 Use only with a Do not breather Do not swallow Avoid contact Avoid prolonge Wash skin thou Handle in accor practice, based sessment Minimize dust Keep containe Keep away fro Take precaution Do not eat, drive 	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			
Hygiene measures :		: If exposure to a flushing system place. When using do Wash contamin The effective o engineering co appropriate de industrial hygie	chemical is likely during typical use, provide eye ns and safety showers close to the working o not eat, drink or smoke. nated clothing before re-use. peration of a facility should include review of ntrols, proper personal protective equipment, gowning and decontamination procedures, ene monitoring, medical surveillance and the trative controls.			
Cond	itions for safe storage	: Keep in proper Store locked u	ly labelled containers.			
Mate	rials to avoid		ith the following product types:			

SECTION 8. EXPOSURE CONTROLS/PERSONAL PROTECTION

Components	CAS-No.	Value type (Form of exposure)	Control parame- ters / Permissible concentration	Basis
Cellulose	9004-34-6	TWA	10 mg/m3	AU OEL
		ation: This value < 1% crystalline	is for inhalable dust silica	containing no
		TWA	10 mg/m3	ACGIH
Starch	9005-25-8	TWA	10 mg/m3	AU OEL
		ation: This value < 1% crystalline	is for inhalable dust silica	containing no
		TWA	10 mg/m3	ACGIH
Rizatriptan	145202-66-0	TWA	10 µg/m3 (OEB 3)	Internal
		Wipe limit	100 µg/100 cm ²	Internal

Components with workplace control parameters

Engineering measures

: All engineering controls should be implemented by facility design and operated in accordance with GMP principles to



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		Containment to are required to	/		
Perso	onal protective equip	ment			
Resp	iratory protection	sure assessm	 If adequate local exhaust ventilation is not available or expo- sure assessment demonstrates exposures outside the rec- ommended guidelines, use respiratory protection. 		
	lter type protection	: Particulates ty	/pe		
Ma	aterial	: Chemical-resi	stant 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 per posable suits)	or laboratory coat. dy garments should be used based upon the rformed (e.g., sleevelets, apron, gauntlets, dis-) to avoid exposed skin surfaces. ate degowning techniques to remove potentially clothing.		

SECTION 9. PHYSICAL AND CHEMICAL PROPERTIES

Appearance	:	powder
Colour	:	pink
Odour	:	odourless
Odour Threshold	:	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



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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
Density	: No data available
Solubility(ies) Water solubility	: No data available
Partition coefficient: n- octanol/water	: No data available
Auto-ignition temperature	: No data available
Decomposition temperature	: No data available
Viscosity Viscosity, kinematic	: No data available
Explosive properties	: Not explosive
Oxidizing properties	: The substance or mixture is not classified as oxidizing.
Molecular weight	: No data available
Particle size	: No data available

SECTION 10. STABILITY AND REACTIVITY

Reactivity Chemical stability Possibility of hazardous reac- tions		Not classified as a reactivity hazard. Stable under normal conditions. May form explosive dust-air mixture during processing, han- dling or other means. Can react with strong oxidizing agents.
Conditions to avoid	:	Heat, flames and sparks. Avoid dust formation.
Incompatible materials	:	Oxidizing agents
Hazardous decomposition products	:	No hazardous decomposition products are known.

SECTION 11. TOXICOLOGICAL INFORMATION

Exposure routes	: Inhalation Skin contact Ingestion Eye contact





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Acu	ite toxicity				
Not	classified based on ava				
	duct:				
Acu	te oral toxicity	:	Acute toxicity of Method: Calcu	estimate: > 2,000 mg/kg lation method	
Con	nponents:				
Cell	lulose:				
Acu	te oral toxicity	:	LD50 (Rat): >	5,000 mg/kg	
Acu	te inhalation toxicity	:	LC50 (Rat): > Exposure time Test atmosphe	:4 h	
Acu	te dermal toxicity	:	LD50 (Rabbit)	> 2,000 mg/kg	
Star	rch:				
Acu	te oral toxicity	:	LD50 (Rat): >	5,000 mg/kg	
Acu	te dermal toxicity	:	LD50 (Rabbit)	> 2,000 mg/kg	
Riza	atriptan:				
Acu	te oral toxicity	:	LD50 (Rat): 2,	227 mg/kg	
			LD50 (Mouse)	: 700 - 1,631 mg/kg	
-	n corrosion/irritation classified based on ava	ilabla	information		
		liable	information.		
	nponents:				
	atriptan:		Dabbit		
Res	ecies sult	:	Rabbit No skin irritatio	on	
	ious eye damage/eye i				
	classified based on ava	ilable	information.		
	nponents:				
	rch:		Dabbit		
Spe Res	ecies sult	:	Rabbit No eye irritatic	n	
	atriptan:				
	ecies narks	:	Bovine cornea Moderate eye		



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Respi	iratory or skin sensi	itisatio	'n			
Skin s	sensitisation					
Not classified based on available information. Respiratory sensitisation						
	oonents:					
Starc	h:					
Test T			Maximisation Tes	st		
	sure routes	:	Skin contact			
Speci		:	Guinea pig			
Resul	t	:	negative			
Rizatı	riptan:					
Test 1	Гуре	:	Maximisation Tes	st		
	sure routes	:	Dermal			
Speci		:	Guinea pig			
Asses Resul	ssment	:	Does not cause s	kin sensitisation.		
Resul	L		negative			
Chro	nic toxicity					
Germ	cell mutagenicity					
Not cl	assified based on ava	ailable	information.			
Comr						
	oonents:					
Cellu						
Cellu		:	Test Type: Bacte Result: negative	rial reverse mutation assay (AMES)		
Cellu	lose:	:	Result: negative	rial reverse mutation assay (AMES) o mammalian cell gene mutation test		
Cellu Genot	lose:	:	Result: negative Test Type: In vitre Result: negative	o mammalian cell gene mutation test		
Cellu Genot	lose: toxicity in vitro	:	Result: negative Test Type: In vitre Result: negative	o mammalian cell gene mutation test nalian erythrocyte micronucleus test (in vivo		
Cellu Genot	lose: toxicity in vitro	:	Result: negative Test Type: In vitre Result: negative Test Type: Mammer cytogenetic assar Species: Mouse	o mammalian cell gene mutation test nalian erythrocyte micronucleus test (in vive y)		
Cellu Genot	lose: toxicity in vitro	:	Result: negative Test Type: In vitre Result: negative Test Type: Mamr cytogenetic assa	o mammalian cell gene mutation test nalian erythrocyte micronucleus test (in vive y)		
Cellu Genot	lose: toxicity in vitro toxicity in vivo	:	Result: negative Test Type: In vitre Result: negative Test Type: Mamer cytogenetic assa Species: Mouse Application Route	o mammalian cell gene mutation test nalian erythrocyte micronucleus test (in vive y)		
Cellul Genot	lose: toxicity in vitro toxicity in vivo	:	Result: negative Test Type: In vitre Result: negative Test Type: Mamr cytogenetic assa Species: Mouse Application Route Result: negative	o mammalian cell gene mutation test nalian erythrocyte micronucleus test (in vivo y) e: Ingestion		
Cellul Genot	lose: toxicity in vitro toxicity in vivo	:	Result: negative Test Type: In vitre Result: negative Test Type: Mamr cytogenetic assa Species: Mouse Application Route Result: negative	o mammalian cell gene mutation test nalian erythrocyte micronucleus test (in vivo y)		
Cellul Genot Genot Starc Genot	lose: toxicity in vitro toxicity in vivo h: toxicity in vitro	:	Result: negative Test Type: In vitre Result: negative Test Type: Mamr cytogenetic assa Species: Mouse Application Route Result: negative Test Type: Bacte	o mammalian cell gene mutation test nalian erythrocyte micronucleus test (in vive y) e: Ingestion		
Cellul Genot Genot Starc Genot	lose: toxicity in vitro toxicity in vivo h: toxicity in vitro riptan:	:	Result: negative Test Type: In vitre Result: negative Test Type: Mamr cytogenetic assa Species: Mouse Application Route Result: negative Test Type: Bacte Result: negative	o mammalian cell gene mutation test nalian erythrocyte micronucleus test (in vive y) e: Ingestion rial reverse mutation assay (AMES)		
Cellul Genot Genot Starc Genot	lose: toxicity in vitro toxicity in vivo h: toxicity in vitro	:	Result: negative Test Type: In vitre Result: negative Test Type: Mamr cytogenetic assa Species: Mouse Application Route Result: negative Test Type: Bacte Result: negative	o mammalian cell gene mutation test nalian erythrocyte micronucleus test (in vivo y) e: Ingestion		
Cellul Genot Genot Starc Genot	lose: toxicity in vitro toxicity in vivo h: toxicity in vitro riptan:	:	Result: negative Test Type: In vitre Result: negative Test Type: Mamr cytogenetic assay Species: Mouse Application Route Result: negative Test Type: Bacte Result: negative	o mammalian cell gene mutation test nalian erythrocyte micronucleus test (in vive y) e: Ingestion rial reverse mutation assay (AMES)		



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		Test Type: In v Result: negativ	vitro mammalian cell gene mutation test ve
		Test Type: Ch Result: negativ	romosome aberration test in vitro
Genot	toxicity in vivo	: Test Type: Ma cytogenetic as Species: Mous Application Ro Result: negativ	se oute: Oral
	nogenicity assified based on avai	lable information.	
<u>Comp</u>	oonents:		
	es cation Route sure time	: Rat : Ingestion : 72 weeks : negative	
Rizatı	riptan:		
	cation Route sure time EL	: Mouse : Oral : 100 weeks : 125 mg/kg boo : negative	dy weight
	cation Route sure time EL	: Rat : Oral : 106 weeks : 106 mg/kg boo : negative	dy weight
•	oductive toxicity acted of damaging the	unborn child.	
-	oonents:		
Cellul	lose:		
Effect	s on fertility	: Test Type: On Species: Rat Application Ro Result: negativ	
Effect ment	s on foetal develop-	: Test Type: Fer Species: Rat Application Ro Result: negativ	rtility/early embryonic development



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Rizat	riptan:		
Effects on fertility		Species: Rat, f Application Ro Fertility: LOAE Symptoms: alt	ute: Oral L: 100 mg/kg body weight ered estrus cycles ects on fertility and early embryonic develop-
		Species: Rat, r Application Ro Fertility: NOAE	ute: Oral L: 250 mg/kg body weight ects on fertility and early embryonic develop-
Effect ment	s on foetal develop-	Species: Rat Application Ro Developmenta	bryo-foetal development ute: Oral I Toxicity: LOAEL: 10 mg/kg body weight atogenic effects, Embryo-foetal toxicity
		Species: Rabb Application Ro Developmenta Result: No tera	
Repro sessn	oductive toxicity - As- nent	: Some evidenc animal experin	e of adverse effects on development, based on nents.
стот	- single exposure		
	assified based on avail	able information.	
<u>Comp</u>	oonents:		
Rizat	riptan:		
Asses	ssment	: May cause dro	wsiness or dizziness.

STOT - repeated exposure

May cause damage to organs (Cardio-vascular system) through prolonged or repeated exposure if swallowed.

Components:

Rizatriptan:

Target Organs Assessment	Cardio-vascular system Causes damage to organs through prolonged or repeated exposure.
	exposure.



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F	Repeat	ted dose toxicity			
<u>c</u>	Compo	onents:			
c	Cellulo	ose:			
N A			:	Rat >= 9,000 mg/kg Ingestion 90 Days	
S	Starch	:			
N A E		- ition Route ire time		Rat >= 2,000 mg/kg Skin contact 28 Days OECD Test Guide	eline 410
F	Rizatri	ptan:			
L A E		tion Route Ire time	:	Rat 1 mg/kg Oral 14 Weeks Dilatation of the p	upil, Increased pulse rate, Redness
L A E		tion Route Ire time		Dog 0.05 mg/kg Intravenous 2 Weeks Dilatation of the p	upil, Increased pulse rate, Redness
L A E		tion Route Ire time		Dog 0.2 mg/kg Oral 1 yr Dilatation of the p	upil
Ν	Not cla	tion toxicity ssified based on availa ence with human exp			
<u>c</u>	Compo	onents:			
	Rizatri				

Ingestion

: Target Organs: Cardio-vascular system Symptoms: asthenia, Fatigue, Pain, Dizziness, Weakness, Drowsiness



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CTION	12. ECOLOGICAL INFO	ORI	MATION	
Ecoto	xicity			
	oonents:			
Cellul				
	ty to fish	:	Exposure time: 48	pes (Japanese medaka)): > 100 mg/l 3 h on data from similar materials
Rizatr	iptan:			
	ty to fish	:	LC50 (Pimephales Exposure time: 96	s promelas (fathead minnow)): > 1,000 mg 3 h
	ty to daphnia and other ic invertebrates	:	EC50 (Daphnia m Exposure time: 48	agna (Water flea)): 1,000 mg/l 8 h
Toxici plants	ty to algae/aquatic	:	EC50 (Pseudokiro mg/l Exposure time: 72 Method: OECD Te	
			NOEC (Pseudokir mg/l Exposure time: 72 Method: OECD Te	
Toxici icity)	ty to fish (Chronic tox-	:	NOEC (Pimephale Exposure time: 32 Method: OECD Te	
	ty to daphnia and other ic invertebrates (Chron- city)	:	NOEC (Daphnia r Exposure time: 21 Method: OECD Te	
Toxici	ty to microorganisms	:	EC50: > 1,000 mg Exposure time: 3 Test Type: Respir Method: OECD Te	h ation inhibition
			NOEC: 1,000 mg/ Exposure time: 3 Test Type: Respir Method: OECD Te	h ation inhibition
Persis	stence and degradabili	ity		
<u>Comp</u>	oonents:			
Cellul Biode	ose: gradability	:	Result: Readily bi	odegradable.



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	riptan: gradability	:	Biodegradation: Exposure time:	
Bioac	cumulative potential			
Comp	oonents:			
Partiti	riptan: on coefficient: n- ol/water	:	log Pow: -0.649	
Mobil	lity in soil			
Comp	oonents:			
Distrik	riptan: oution among environ- al compartments	:	log Koc: 3.83 Method: OECD	Test Guideline 106
Other	adverse effects			
	ta available			
	ita available 13. DISPOSAL CONSI	DEF	RATIONS	
ECTION Dispo Waste		DEF	Dispose of in ac Empty containe dling site for rec	ccordance with local regulations. rs should be taken to an approved waste han cycling or disposal. specified: Dispose of as unused product.
ECTION Dispo Waste Conta	13. DISPOSAL CONSI osal methods e from residues	:	Dispose of in ac Empty containe dling site for rec If not otherwise	rs should be taken to an approved waste han cycling or disposal.
ECTION Dispo Waste Conta	 13. DISPOSAL CONSI Desal methods Definition for the formation of the formation of	:	Dispose of in ac Empty containe dling site for rec If not otherwise	rs should be taken to an approved waste har cycling or disposal.
ECTION Dispo Waste Conta	13. DISPOSAL CONSI osal methods e from residues iminated packaging	:	Dispose of in ac Empty containe dling site for rec If not otherwise	rs should be taken to an approved waste har cycling or disposal.
ECTION Dispo Waste Conta ECTION Interr UNRT	 13. DISPOSAL CONSI Desal methods e from residues aminated packaging 14. TRANSPORT INFO national Regulations TDG 	: DRM	Dispose of in ac Empty containe dling site for red If not otherwise	rs should be taken to an approved waste har cycling or disposal.
ECTION Dispo Waste Conta ECTION Intern UNRT Not re IATA-	 13. DISPOSAL CONSI Desal methods Desal me	: : DRM	Dispose of in ac Empty containe dling site for red If not otherwise	rs should be taken to an approved waste har cycling or disposal.
ECTION Dispo Waste Conta ECTION Interr UNRT Not re IATA- Not re IMDG	 13. DISPOSAL CONSI Desal methods a from residues b from residues b from residues a from residues b from resi	: : DRM s go	Dispose of in ad Empty containe dling site for red If not otherwise	rs should be taken to an approved waste han cycling or disposal.
ECTION Dispo Waste Conta ECTION Intern UNRT Not re IATA- Not re IMDG Not re Trans	 13. DISPOSAL CONSI 5. DISPOSAL CONSI 5. Disposal methods a from residues b from residues a from residues b f	: : DRM s go s go s go g to	Dispose of in ad Empty containe dling site for red If not otherwise ATION od od od Annex II of MAR	rs should be taken to an approved waste har cycling or disposal.
ECTION Waste Conta ECTION Intern UNRT Not re IATA- Not re IMDG Not re Trans Not ap	 13. DISPOSAL CONSI 23. DISPOSAL CONSI 24. TRANSPORT INFO 24. TRANSPORT INFO 25. DGR 29. Uated as a dangerous 25. Code 29. Uated as a dangerous 	: : DRM s go s go s go g to	Dispose of in ad Empty containe dling site for red If not otherwise ATION od od od Annex II of MAR	rs should be taken to an approved waste han cycling or disposal. specified: Dispose of as unused product.





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SECTION 15. REGULATORY INFORMATION

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

Prohibition/Licensing Requirements

: There is no applicable prohibition, authorisation and restricted use requirements, including for carcinogens referred to in Schedule 10 of the model WHS Act and Regulations.

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

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

SECTION 16. OTHER INFORMATION

Further information		
Revision Date Sources of key data used to compile the Safety Data Sheet	:	10.10.2020 Internal technical data, data from raw material SDSs, OECD eChem Portal search results and European Chemicals Agen- cy, http://echa.europa.eu/
Date format	:	dd.mm.yyyy
Full text of other abbreviation	ns	
ACGIH AU OEL	:	USA. ACGIH Threshold Limit Values (TLV) Australia. Workplace Exposure Standards for Airborne Con-
		taminants.
ACGIH / TWA	:	8-hour, time-weighted average
AU OEL / TWA	:	Exposure standard - time weighted average

AIIC - Australian Inventory of Industrial Chemicals; ANTT - National Agency for Transport by Land of Brazil; ASTM - American Society for the Testing of Materials; bw - Body weight; CMR - Carcinogen, Mutagen or Reproductive Toxicant; DIN - Standard of the German Institute for Standardisation; DSL - Domestic Substances List (Canada); ECx - Concentration associated with x% response; ELx - Loading rate associated with x% response; EmS - Emergency Schedule; ENCS - Existing and New Chemical Substances (Japan); ErCx - Concentration associated with x% growth rate response; ERG - Emergency Response Guide; GHS - Globally Harmonized System; GLP - Good Laboratory Practice; IARC - International Agency for Research on Cancer; IATA - International Air Transport Association; IBC - International Code for the Construction and Equipment of Ships carrying Dangerous Chemicals in Bulk; IC50 - Half maximal inhibitory concentration; ICAO - International Civil Aviation Organization; IECSC - Inventory of Existing Chemical Substances in China; IMDG - International Maritime Dangerous Goods; IMO - International Maritime Organization; ISHL - Industrial Safety and Health Law (Japan); ISO - International Organisation for Standardization; KECI - Korea Existing Chemicals Inventory; LC50 - Lethal Con-



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centration to 50 % of a test population; LD50 - Lethal Dose to 50% of a test population (Median Lethal Dose); MARPOL - International Convention for the Prevention of Pollution from Ships; n.o.s. - Not Otherwise Specified; Nch - Chilean Norm; NO(A)EC - No Observed (Adverse) Effect Concentration; NO(A)EL - No Observed (Adverse) Effect Level; NOELR - No Observable Effect Loading Rate; NOM - Official Mexican Norm; NTP - National Toxicology Program; NZIoC - New Zealand Inventory of Chemicals; OECD - Organization for Economic Co-operation and 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; SADT - Self-Accelerating Decomposition Temperature; SDS - Safety Data Sheet; TCSI - Taiwan Chemical Substance Inventory; TDG - Transportation of Dangerous Goods; TSCA - Toxic Substances Control Act (United States); UN - United Nations; UNRTDG - United Nations Recommendations on the Transport of Dangerous Goods; vPvB - Very Persistent and Very Bioaccumulative; WHMIS - Workplace Hazardous Materials Information System

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

AU / EN