according to Regulation (EC) No. 1907/2006

ARADUR® HY 1300 CH

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

1.1 Product identifier	
Trade name	: ARADUR® HY 1300 CH
Unique Formula Identifier (UFI)	: WPT0-F0WT-5009-MK3M

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

Use of the	: Component used for the manufacture of electrical insulation
Substance/Mixture	parts

1.3 Details of the supplier of the safety data sheet

Company Address	 Huntsman Advanced Materials (Europe)BVBA Everslaan 45 3078 Everberg Belgium
Telephone Telefax	: +41 61 299 20 41 : +41 61 299 20 40
E-mail address of person responsible for the SDS	: Global_Product_EHS_AdMat@huntsman.com

1.4 Emergency telephone number

a Emergency telephone number		
Emergency telephone number	Centres Antipoison et de Toxicovigilance: ANGERS: 02 41 48 21 21 BORDEAUX: 05 56 96 40 80 LILLE: 0 825 812 822 LYON: 04 72 11 69 11 MARSEILLE 04 91 75 25 25 NANCY: 03 83 32 36 36 PARIS: 01 40 05 48 48 RENNES: 02 99 59 22 22 STRASBOURG: 03 88 37 37 37 TOULOUSE: 05 61 77 74 47 EUROPE: $+32$ 35 75 1234 France ORFILA: $+33(0)145425959$ ASIA: $+65$ 6336-6011 China: $+86$ 20 39377888 +86 532 83889090 India: $+$ 91 22 42 87 5333 Australia: 1800 786 152 New Zealand: 0800 767 437 USA: $+1$ 800-424-9300	

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SECTION 2: Hazards identification

2.1 Classification of the substance or mixture

Classification (REGULATION (EC) No 1272/2008)

Acute toxicity, Category 4H302: Harmful if swallowed.Acute toxicity, Category 4H312: Harmful in contact with skin.Skin corrosion, Sub-category 1BH314: Causes severe skin burns and eye damage.Serious eye damage, Category 1H318: Causes serious eye damage.Skin sensitisation, Category 1H317: May cause an allergic skin reaction.Reproductive toxicity, Category 2H361d: Suspected of damaging the unborn child.Long-term (chronic) aquatic hazard, Category 2H411: Toxic to aquatic life with long lasting effects.

2.2 Label elements

Labelling (REGULATION (EC) No 1272/2008)

Hazard pictograms :	
Signal word :	Danger
Hazard statements :	 H302 + H312 Harmful if swallowed or in contact with skin. H314 Causes severe skin burns and eye damage. H317 May cause an allergic skin reaction. H361d Suspected of damaging the unborn child. H411 Toxic to aquatic life with long lasting effects.
Precautionary statements :	Prevention:P273Avoid release to the environment.P280Wear protective gloves/ protective clothing/ eyeprotection/ face protection/ hearing protection.
	Response: P303 + P361 + P353 IF ON SKIN (or hair): Take off immediately all contaminated clothing. Rinse skin with water. P304 + P340 + P310 IF INHALED: Remove person to fresh air and keep comfortable for breathing. Immediately call a POISON CENTER/ doctor. P305 + P351 + P338 + P310 IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing. Immediately call a POISON CENTER/ doctor. P391 Collect spillage.

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Hazardous components which must be listed on the label:

Propylidynetrimethanol, propoxylated, reaction products with ammonia Amines, polyethylenepoly-, triethylenetetramine fraction salicylic acid

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

SECTION 3: Composition/information on ingredients

3.2 Mixtures

Chemical nature

: Polyamines

Hazardous components

Chemical name	CAS-No. EC-No. Index-No. Registration number	Classification	Concent ration (% w/w)
Propylidynetrimethanol, propoxylated, reaction products with ammonia	39423-51-3 500-105-6 01-2119556886-20	Acute Tox. 4; H302 Acute Tox. 4; H312 Eye Dam. 1; H318 Aquatic Chronic 2; H411 Acute toxicity estimate Acute oral toxicity: 550 mg/kg Acute dermal toxicity: 1 000,1 mg/kg	>= 70 - < 90
Amines, polyethylenepoly-, triethylenetetramine fraction	90640-67-8 292-588-2 01-2119487919-13	Acute Tox. 4; H302 Acute Tox. 4; H312 Skin Corr. 1B; H314 Eye Dam. 1; H318 Skin Sens. 1; H317 Aquatic Chronic 3; H412	>= 10 - < 20
salicylic acid	69-72-7 200-712-3 607-732-00-5 01-2119486984-17	Acute Tox. 4; H302 Eye Dam. 1; H318 Repr. 2; H361d	>= 3 - < 10

For explanation of abbreviations see section 16.



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SECTION 4: First aid measures

4.1 Description of first aid measures				
General advice	:	Move out of dangerous area. Consult a physician. Show this safety data sheet to the doctor in attendance. Treat symptomatically. Get medical attention if symptoms occur.		
Protection of first-aiders	:	First Aid responders should pay attention to self-protection and use the recommended protective clothing If potential for exposure exists refer to Section 8 for specific personal protective equipment. Avoid inhalation, ingestion and contact with skin and eyes. No action shall be taken involving any personal risk or without suitable training. It may be dangerous to the person providing aid to give mouth-to-mouth resuscitation.		
If inhaled	:	If inhaled, remove to fresh air. Get medical attention if symptoms occur.		
In case of skin contact	:	Immediate medical treatment is necessary as untreated wounds from corrosion of the skin heal slowly and with difficulty. If on skin, rinse well with water. If on clothes, remove clothes.		
In case of eye contact	:	Small amounts splashed into eyes can cause irreversible tissue damage and blindness. In the case of contact with eyes, rinse immediately with plenty of water and seek medical advice. Continue rinsing eyes during transport to hospital. Remove contact lenses. Keep eye wide open while rinsing. If eye irritation persists, consult a specialist.		
If swallowed	:	Keep respiratory tract clear. Do NOT induce vomiting. Never give anything by mouth to an unconscious person. If symptoms persist, call a physician. Take victim immediately to hospital.		

4.2 Most important symptoms and effects, both acute and delayed

None known.

4.3 Indication of any immediate medical attention and special treatment needed

Treatment

: Treat symptomatically.

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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	:	Exercise caution when using a high volume water jet as it may scatter and spread fire
5.2	Special hazards arising from	the	substance or mixture
	Specific hazards during firefighting	:	Do not allow run-off from fire fighting to enter drains or water courses.
	Hazardous combustion products	:	Ammonia Carbon oxides Nitrogen oxides (NOx)
5.3	Advice for firefighters		
	Special protective equipment for firefighters	:	Wear self-contained breathing apparatus for firefighting if necessary.
	Specific extinguishing methods	:	Use extinguishing measures that are appropriate to local circumstances and the surrounding environment.
	Further information	:	Collect contaminated fire extinguishing water separately. This must not be discharged into drains. Fire residues and contaminated fire extinguishing water must be disposed of in accordance with local regulations.

SECTION 6: Accidental release measures

• •	ve equipment and emergency procedures Use personal protective equipment. Refer to protective measures listed in sections 7 and 8.	
6.2 Environmental precautions		
Environmental precautions :	Prevent product from entering drains. Prevent further leakage or spillage if safe to do so. If the product contaminates rivers and lakes or drains inform respective authorities.	
6.3 Methods and material for containment and cleaning up		

Methods for cleaning up	:	Neutralise with acid.
		Soak up with inert absorbent material (e.g. sand, silica gel,
		acid binder, universal binder, sawdust).

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Keep in suitable, closed containers for disposal.

6.4 Reference to other sections

For disposal considerations see section 13., See Section 1 for emergency contact information., For personal protection see section 8.

SECTION 7: Handling and storage

7.1 Precautions for safe handling	g	
Advice on safe handling	:	Repeated or prolonged skin contact may cause skin irritation and/or dermatitis and sensitisation of susceptible persons. Persons suffering from asthma, eczema or skin problems should avoid contact, including dermal contact, with this product. Avoid exposure - obtain special instructions before use. Avoid contact with skin and eyes. For personal protection see section 8. Smoking, eating and drinking should be prohibited in the application area. To avoid spills during handling keep bottle on a metal tray. Dispose of rinse water in accordance with local and national regulations. Do not breathe vapours or spray mist.
Advice on protection against fire and explosion	:	Normal measures for preventive fire protection.
Hygiene measures	:	When using do not eat or drink. When using do not smoke. Wash hands before breaks and at the end of workday.

7.2 Conditions for safe storage, including any incompatibilities

	Requirements for storage areas and containers	:	Keep container tightly closed in a dry and well-ventilated place. Containers which are opened must be carefully resealed and kept upright to prevent leakage. Observe label precautions. Keep in properly labelled containers.
	Advice on common storage	:	Do not store near acids.
	Further information on storage stability	:	Stable under normal conditions.
	Recommended storage temperature	:	2 - 40 °C
7.3	Specific end use(s) Specific use(s)	:	No data available



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SECTION 8: Exposure controls/personal protection

8.1 Control parameters

Contains no substances with occupational exposure limit values.

Derived No Effect Level (DNEL) according to Regulation (EC) No. 1907/2006:

Substance name	End Use	Exposure routes	Potential health effects	Value
Propylidynetrimethan ol, propoxylated, reaction products with ammonia	Workers	Inhalation	Long-term systemic effects	14,1 mg/m3
	Workers	Dermal	Long-term systemic effects	1,6 mg/kg bw/day
salicylic acid	Workers	Inhalation	Long-term systemic effects	5 mg/m3
	Workers	Inhalation	Long-term local effects	5 mg/m3
	Workers	Dermal	Long-term systemic effects	2,3 mg/kg bw/day
	Consumers	Inhalation	Long-term systemic effects	4 mg/m3
	Consumers	Dermal	Long-term systemic effects	1 mg/kg bw/day
	Consumers	Oral	Long-term systemic effects	1 mg/kg bw/day
	Consumers	Oral	Acute effects, Short- term exposure	4 mg/kg bw/day
Amines, polyethylenepoly-, triethylenetetramine fraction	Workers	Inhalation	Long-term systemic effects	0,54 mg/m3
	Consumers	Inhalation	Long-term systemic effects	0,096 mg/m3
	Consumers	Oral	Long-term systemic effects	14 mg/kg bw/day

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

Substance name	Environmental Compartment	Value
Propylidynetrimethanol, propoxylated, reaction products	Fresh water	0,004 mg/l
with ammonia		
	Intermittent use/release	0,044 mg/l
	Marine water	0 mg/l
	Fresh water sediment	0,022 mg/kg dry
		weight (d.w.)
	Marine sediment	0,002 mg/kg dry
		weight (d.w.)
	Sewage treatment plant	10 mg/l
	Soil	0,002 mg/kg dry
		weight (d.w.)
salicylic acid	Marine water	0,02 mg/l
	Sewage treatment plant	162 mg/l



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	Fresh water sediment	1,42 mg/kg dry weight (d.w.)
	Marine sediment	0,142 mg/kg dry weight (d.w.)
	Soil	0,166 mg/kg dry weight (d.w.)
	Secondary Poisoning	
Amines, polyethylenepoly-, triethylenetetramine fraction	Fresh water	0,027 mg/l
	Marine water	0,003 mg/l
	Sewage treatment plant	0,13 mg/l
	Fresh water sediment	8,572 mg/kg dry weight (d.w.)
	Marine sediment	0,857 mg/kg dry weight (d.w.)
	Soil	1,25 mg/kg dry weight (d.w.)

8.2 Exposure controls

Personal protective equipment

Eye/face protection	:	Eye wash bottle with pure water Tightly fitting safety goggles Wear face-shield and protective suit for abnormal processing problems.	
Hand protection Material Break through time		butyl-rubber > 8 h	
Material Break through time	-	Nitrile rubber 10 - 480 min	
Material Break through time		Ethyl Vinyl Alcohol Laminate (EVAL) > 8 h	
Remarks	:	The selected protective gloves have to satisfy the specifications of Regulation (EU) 2016/425 and the standard EN 374 derived from it. Gloves should be discarded and replaced if there is any indication of degradation or chemical breakthrough. Take note of the information given by the producer concerning permeability and break through times, and of special workplace conditions (mechanical strain, duration of contact). Chemical-resistant, impervious gloves complying with an approved standard should be worn at all times when handling chemical products if a risk assessment indicates this is necessary. The suitability for a specific workplace should be discussed with the producers of the protective gloves.	
Skin and body protection	:	Impervious clothing Choose body protection according to the amount and concentration of the dangerous substance at the work place.	

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Respiratory protection		ventilation is pro that exposures a	protection unless adequate local exhaust ovided or exposure assessment demonstrates are within recommended exposure guidelines Ild conform to EN 14387
Filter type : Co		: Combined partic	culates and ammonia/amines type (K-P)

SECTION 9: Physical and chemical properties

9.1 Information on basic physical and chemical properties

Physical state	: liquid
Colour	: light brown
Odour	: amine-like
Odour Threshold	: No data is available on the product itself.
рН	: 11 (20 °C) Concentration: 500 g/l
Melting point/freezing point	: No data is available on the product itself.
Boiling point	: > 200 °C
Flash point	: > 150 °C Method: Pensky-Martens closed cup
Flammability (solid, gas)	: No data is available on the product itself.
Upper explosion limit / Upper flammability limit	: No data is available on the product itself.
Lower explosion limit / Lower flammability limit	: No data is available on the product itself.
Vapour pressure	: <1 hPa (20 °C)
Relative vapour density	: No data is available on the product itself.
Relative density	: 1 (25 °C)
Density	: 1 g/cm3 (25 °C)
Solubility(ies) Water solubility	: partly miscible (20 °C)
Solubility in other solvents	: No data is available on the product itself.
Partition coefficient: n- octanol/water	: No data is available on the product itself.



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Auto	-ignition temperature	: No data is ava	ilable on the product itself.	
Decomposition temperature		: >200 °C		
Viscosity Viscosity, dynamic		: 160 - 200 mPa	a.s (25 °C)	
9.2 Other	r information			
No data available				
SECTIO	N 10: Stability and re	activity		
10.1 Rea	ctivity			
No d	angerous reaction know	n under conditions of	normal use.	
10.2 Che	mical stability			
Stable under normal conditions.				
10.3 Pos	sibility of hazardous re	actions		
Haza	ardous reactions	: No hazards to	be specially mentioned.	
10.4 Con	ditions to avoid			

10.4 Conditions to avoid

Conditions to avoid : None known.

10.5 Incompatible materials

Materials to avoid	: Strong acids
	Strong bases
	Strong oxidizing agents

10.6 Hazardous decomposition products

Hazardous decomposition products	: ammonia, anhydrous Aldehydes Nitrogen oxides (NOx) carbon monoxide carbon dioxide Ketones
----------------------------------	--

SECTION 11: Toxicological information

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

Acute toxicity

Product:		
Acute oral toxicity	:	Acute toxicity estimate: 624,28 mg/kg Method: Calculation method
Acute dermal toxicity	:	Acute toxicity estimate: 1 213 mg/kg Method: Calculation method



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Components:	
Propylidynetrimethanol, p Acute oral toxicity	 copoxylated, reaction products with ammonia: LD50 (Rat, female): 550 mg/kg Method: OECD Test Guideline 425
	Acute toxicity estimate: 550 mg/kg Method: Calculation method
Acute dermal toxicity	: LD50 (Rat, male and female): > 1 000 mg/kg Method: OECD Test Guideline 402
	Acute toxicity estimate: 1 000,1 mg/kg Method: Calculation method
Amines, polyethylenepoly	, triethylenetetramine fraction:
Acute oral toxicity	: LD50 (Rat, male and female): 1 716,2 mg/kg
	Method: OECD Test Guideline 401 Assessment: The component/mixture is moderately toxic afte single ingestion.
Acute dermal toxicity	: LD50 (Rabbit, male and female): 1 465,4 mg/kg
	Method: OECD Test Guideline 402 Assessment: The component/mixture is moderately toxic afte single contact with skin.
salicylic acid:	
Acute oral toxicity	: LD50 (Rat, male): 891 mg/kg Method: OECD Test Guideline 401 GLP: no
	Assessment: The component/mixture is moderately toxic afte single ingestion.
Acute inhalation toxicity	: LC50 (Rat, male): > 0,9 mg/l
	Exposure time: 1 h Test atmosphere: dust/mist
	Assessment: The substance or mixture has no acute inhalation toxicity
Acute dermal toxicity	: LD50 (Rat, male and female): > 2 000 mg/kg Method: OECD Test Guideline 402
	GLP: yes Assessment: The substance or mixture has no acute dermal toxicity
Skin corrosion/irritation	
Components:	
	opoxylated, reaction products with ammonia:
Species Method Result	 Rabbit OECD Test Guideline 404 Mild skin irritation

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Speci Metho Resul	bd	: reconstructe : OECD Test (: No skin irrita	
Amin	es, polyethylenepoly	/-, triethylenetetrar	nine fraction:
Speci			d human epidermis (RhE)
Asses Metho	ssment	: Causes burn	s. Guideline 435
Resul			er 3 minutes to 1 hour of exposure
Speci		: Rabbit	
	ssment	: Causes burn	
Metho	Da	: OECD lest	Guideline 404

: Corrosive after 3 minutes to 1 hour of exposure

eal	licy	lic	acid:
5 a	псу	ΠC	aciu.

Result

Species	:	Rabbit
Assessment	:	No skin irritation
Method	:	OECD Test Guideline 404
Result	:	No skin irritation
GLP	:	yes

Serious eye damage/eye irritation

Components:

Propylidynetrimethanol, propoxylated, reaction products with ammonia:

Method	:	OECD Test Guideline 405
Result	:	Irreversible effects on the eye

Amines, polyethylenepoly-, triethylenetetramine fraction:

Species	:	Rabbit
Assessment	:	Risk of serious damage to eyes.
Method	:	OECD Test Guideline 405
Result	:	Irreversible effects on the eye

salicylic acid:

Species	:	Rabbit
Assessment	:	Risk of serious damage to eyes.
Result	:	Irreversible effects on the eye

Respiratory or skin sensitisation

Components:

Propylidynetrimethanol, propoxylated, reaction products with ammonia:

Species Assessment Method	:	Skin Guinea pig Did not cause sensitisation on laboratory animals. OECD Test Guideline 406 Did not cause sensitisation on laboratory animals.
Result	:	Did not cause sensitisation on laboratory animals.



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Exposure routes	: Skin
Species	: Guinea pig
Assessment	: Probability or evidence of skin sensitisation in humans
Method	: OECD Test Guideline 406
Result	: Probability or evidence of skin sensitisation in humans
salicylic acid:	
Test Type	: Local lymph node assay (LLNA)
Exposure routes	: Skin
Species	: Mouse
Method	: OECD Test Guideline 429
Result	: Does not cause skin sensitisation.
Germ cell mutagenicity	
Components:	
	propoxylated, reaction products with ammonia:
Genotoxicity in vitro	: Test Type: reverse mutation assay
	Test system: Salmonella typhimurium Metabolic activation: with and without metabolic activatio
	Method: OECD Test Guideline 471
	Result: negative
	hoodit. hogalivo
	Test Type: unscheduled DNA synthesis assay
	Test system: rat hepatocytes
	Metabolic activation: Metabolic activation
	Method: OECD Test Guideline 482
	Result: negative
	Test Type: In vitro mammalian cell gene mutation test
	Test system: Chinese hamster ovary cells
	Metabolic activation: with and without metabolic activatio
	Method: OECD Test Guideline 476
	Result: negative
Genotoxicity in vivo	: Test Type: In vivo micronucleus test
	Species: Mouse (male and female)
	Cell type: Bone marrow
	Application Route: Intraperitoneal injection
	Dose: 2.5 mg/kg Method: OECD Test Guideline 474
	Result: negative
Amines, polvethylenepo	ly-, triethylenetetramine fraction:
Genotoxicity in vitro	: Test Type: reverse mutation assay
•	Test system: Salmonella tryphimurium and E. coli
	Metabolic activation: with and without metabolic activation
	Method: OECD Test Guideline 471
	Result: positive

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		Metabolic activ	uman lymphocytes ation: with and without metabolic activation Test Guideline 487
Geno	toxicity in vivo	Species: Mouse Cell type: Bone Application Rou Dose: 0 - 600 n	ute: Intraperitoneal injection ng/kg Test Guideline 474
salicy	/lic acid:		
Geno	toxicity in vitro	Test system: S Metabolic activ	erse mutation assay almonella tryphimurium and E. coli ation: with and without metabolic activation Test Guideline 471 e
		Test system: C Metabolic activ	omosome aberration test in vitro hinese hamster ovary cells ation: with and without metabolic activation Test Guideline 473 e
		Test system: m Metabolic activ	itro mammalian cell gene mutation test ouse lymphoma cells ation: with and without metabolic activation Test Guideline 476 e
Geno	toxicity in vivo	: Test Type: siste Species: Mouse Cell type: Bone Application Rou Dose: 350 mg/l Method: OPPT Result: negativ	marrow ute: Oral <g S 870.5915</g
		Species: Mouse Cell type: Bone	marrow ute: Intraperitoneal injection 0 mg/kg S 870.5915
		Species: Mouse Cell type: Bone Application Rou	

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Dose: 50/100/200 mg/kg Method: OECD Test Guideline 475 Result: negative

Species: Mouse (male) Cell type: Bone marrow Application Route: Oral Dose: 350 mg/kg Method: OECD Test Guideline 475 Result: negative

Carcinogenicity

Components:

Amines, polyethylenepoly-, triethylenetetramine fraction:

Species Application Route NOAEL Method Result		Mouse, male Dermal >= 50 mg/kg bw/day OECD Test Guideline 451 negative
Species Application Route Exposure time NOAEL	:	Mouse, male Dermal 104 weeks >= 20 mg/kg bw/day

•	2 20 mg/kg bw/uay
:	OECD Test Guideline 451

: negative

salicylic acid:

Method

Result

Species	:	Rat, male and female
Application Route	:	Oral
Exposure time	:	24 month(s)
Dose	:	0,50,250,500,1000 mg/kg
Frequency of Treatment	:	7 daily
NOAEL	:	500 mg/kg bw/day
Result	:	negative
Remarks	:	Information given is based on data obtained from similar substances.

Reproductive toxicity

Components:

Propylidynetrimethanol, propoxylated, reaction products with ammonia:

Effects on fertility	: Test Type: Reproduction / Developmental Toxicity Screening Test
	Species: Rat, male and female
	Application Route: Dermal
	Dose: 0, 10, 50, 100 mg/kg
	General Toxicity - Parent: NOAEL: > 100 mg/kg body weight
	General Toxicity F1: NOAEL: > 100 mg/kg body weight
	Method: OECD Test Guideline 421
	Result: No effects on fertility and early embryonic
	development were detected.

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Effects on foetal development		:	 Test Type: Pre-natal Species: Rat, female Application Route: Oral Dose: 0/10/100/125/200 milligram per kilogram Duration of Single Treatment: 16 d General Toxicity Maternal: NOEL: 125 mg/kg body weight Developmental Toxicity: NOEL: 125 mg/kg body weight Method: OECD Test Guideline 414 	
Amines, polyeth	ylenepoly-, tı	riet	hylenetetramine	fraction:
Effects on foetal development		:		: Oral mg/kg bw/day Treatment: 10 d Maternal: NOAEL: >= 750 mg/kg body wei oxicity: NOAEL: >= 750 mg/kg body weigh est Guideline 414
				: Dermal g/kg bw/day Treatment: 13 d Maternal: NOAEL: 50 mg/kg body weight oxicity: NOAEL: >= 125 mg/kg body weigh est Guideline 414
Reproductive toxi Assessment	city -	:	under further eval	ects of Triethylenetetramine (TETA) are uation as part of the EU REACH program aminoethyl ethanolamine (AEEA) content.
salicylic acid:				
Effects on foetal development		:	General Toxicity N Developmental To Method: OECD Te	: Oral Treatment: 3 - 13 d Maternal: NOAEL: 125 mg/kg body weight oxicity: NOAEL: 250 mg/kg body weight est Guideline 414 tion given is based on data obtained from
Reproductive toxi Assessment	city -	:	Some evidence of animal experiment	f adverse effects on development, based o ts.
STOT - single ex No data available	-			
STOT - repeated No data available				



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Repeated dose toxicity

Components:

Propylidynetrimethanol, propoxylated, reaction products with ammonia:

:	Rat, male and female
:	>= 100 mg/kg
:	Oral
:	90 d
:	0, 10, 75, 100, 150, 200 mg/kg
:	OECD Test Guideline 408
:	Rat, male and female
:	> 160 mg/kg
:	Dermal
:	90 d 6 h
:	5 days/week
:	0/16/50/160 mg/kg bw7day
:	OECD Test Guideline 411

Amines, polyethylenepoly-, triethylenetetramine fraction:

Annies, poryetitytenepory, t		
Species NOAEL Application Route Exposure time Number of exposures Dose Method Target Organs Remarks		Rat, male and female 350 mg/kg Oral 28 d 7 d 100/350/1000 mg/kg bw/day OECD Test Guideline 407 Lungs Information given is based on data obtained from similar substances.
Species NOAEL Application Route Target Organs Remarks	:	Dog, male and female 125 mg/kg Oral Lungs Information given is based on data obtained from similar substances.
Species NOAEL Application Route Method Remarks	:	Dog, male and female 50 mg/kg Oral Subchronic toxicity Information given is based on data obtained from similar substances.
Species NOAEL Application Route Exposure time Dose Method Target Organs Remarks		Rat, male and female 50 mg/kg Oral 26 weeks 50/175/600 mg/kg bw/day OECD Test Guideline 408 Lungs Information given is based on data obtained from similar substances.

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	EL cation Route sure time od	 Mouse, male a 92 mg/kg, 600 Oral 120/600/3000 p OECD Test Gu Information give substances. 	opm
Spec NOAI Applic Expo	EL cation Route sure time per of exposures od	: Chronic toxicity	, 1000 mg/kg bw
Expo	C cation Route sure time per of exposures od	 Rat, female 700 mg/m3 inhalation (vape) 7 h 4 Weeks 5 days/week 635 mg/m3 OECD Test Guits no Information given substances. 	

Aspiration toxicity

No data available

11.2 Information on other hazards

Endocrine disrupting properties

Product:

Assessment

: 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

Experience with human exposure

No data available

Toxicology, Metabolism, Distribution

No data available

Neurological effects

No data available



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Further information

No data available

SECTION 12: Ecological information

12.1 Toxicity

Components:				
Propylidynetrimethanol, propo	DescriptionDescriptionLC50 (Oncorhynchus mykiss (rainbow trout)): > 100 mg/lExposure time: 96 hTest Type: static testTest substance: Fresh waterMethod: OECD Test Guideline 203GLP: yes			
Toxicity to daphnia and other : aquatic invertebrates	EC50 (Daphnia magna (Water flea)): 13 mg/l Exposure time: 48 h Test Type: static test Test substance: Fresh water Method: OECD Test Guideline 202 GLP: yes			
Toxicity to algae/aquatic : plants	ErC50 (Selenastrum capricornutum (green algae)): 4,4 mg/l Exposure time: 72 h Test Type: static test Test substance: Fresh water Method: OECD Test Guideline 201 GLP: yes			
	NOEC (Selenastrum capricornutum (green algae)): 1 mg/l Exposure time: 72 h Test Type: static test Test substance: Fresh water Method: OECD Test Guideline 201 GLP: yes			
Toxicity to microorganisms :	EC50 (activated sludge): ca. 1 000 mg/l Exposure time: 0,5 h Test Type: static test Test substance: Fresh water Method: OECD Test Guideline 209 GLP: yes			
Amines, polyethylenepoly-, triethylenetetramine fraction:				
Toxicity to fish :	LC50 (Poecilia reticulata (guppy)): 570 mg/l Exposure time: 96 h Test Type: semi-static test Test substance: Fresh water Method: Directive 67/548/EEC, Annex V, C.1.			
	LC50 (Leuciscus idus (Golden orfe)): 200 - 500 mg/l Exposure time: 96 h			

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Version Revision Date: SDS Number: 400001008624 3.0 29.11.2022 Date of first issue: 20.11.2018 Print Date 11.12.2023 LC50 (Pimephales promelas (fathead minnow)): 330 mg/l End point: mortality Exposure time: 96 h Test Type: static test Test substance: Fresh water Method: Fish Acute Toxicity Test Toxicity to daphnia and other : EC50 (Daphnia magna (Water flea)): 31,1 mg/l aquatic invertebrates End point: Immobilization Exposure time: 48 h Test Type: static test Test substance: Fresh water Method: Directive 67/548/EEC, Annex V, C.2. Toxicity to algae/aquatic ErC50 (Selenastrum capricornutum (green algae)): 20 mg/l plants Exposure time: 72 h Test Type: semi-static test Test substance: Fresh water Method: OECD Test Guideline 201 EC10 (Selenastrum capricornutum (green algae)): 1,34 mg/l Exposure time: 72 h Test Type: semi-static test Test substance: Fresh water Method: OECD Test Guideline 201 NOEC (Bacteria): >= 100 mg/l Toxicity to microorganisms 2 Exposure time: 28 d Method: OECD Test Guideline 216 EC50 (Bacteria): > 100 mg/l Exposure time: 28 h Method: OECD Test Guideline 216 EC50 (Bacteria): 15,7 mg/l Exposure time: 2 h Test Type: static test Test substance: Fresh water NOEC (Bacteria): 1,3 mg/l Exposure time: 2 h Test Type: static test Test substance: Fresh water Toxicity to daphnia and other : EC10: 1,9 mg/l aquatic invertebrates Exposure time: 21 d Species: Daphnia magna (Water flea) (Chronic toxicity) Test Type: semi-static test Test substance: Fresh water Method: OECD Test Guideline 202 Toxicity to soil dwelling NOEC: ca. 62,5 mg/kg : organisms Exposure time: 56 d

Species: Eisenia fetida (earthworms)



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			Method: OECD T	est Guideline 222
Ecotoxi	icology Assessment			
Chronic	aquatic toxicity	:	Harmful to aquati	c life with long lasting effects.
salicylic	c acid:			
Toxicity	to fish	:	Exposure time: 96 Test Type: flow-th Analytical monitor Test substance: F Method: OECD T GLP: no	nrough test ring: yes Fresh water est Guideline 203 ation given is based on data obtained from
	to daphnia and other invertebrates	:	Exposure time: 44 Test Type: static Analytical monitor Test substance: F	test ring: yes
Toxicity plants	to algae/aquatic	:	EC50 (Desmodes Exposure time: 72 Method: OECD T	
Toxicity	to microorganisms	:	Exposure time: 10 Test Type: static Test substance: F Method: ISO Method:	test Fresh water hod, other ation given is based on data obtained from
aquatic	to daphnia and other invertebrates c toxicity)	:		1 d a magna (Water flea) est Guideline 202
2 2 Perciet	ence and degradabili	itv		
<u>Compo</u>	-	• • •		

Propylidynetrimethanol, propoxylated, reaction products with ammonia:

: Test Type: aerobic
Inoculum: activated sludge
Result: Not readily biodegradable.
Biodegradation: < 5 %

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		Exposure time: Method: OECD	28 d Test Guideline 301F
Stabil	lity in water	pH: 7,5	alf life (DT50): > 1 yr (25 °C) 9 Test Guideline 111 h water
Amin	es, polyethylenepoly	/-, triethylenetetramiı	ne fraction:
Biodegradability		: Inoculum: activ Result: Not rea Biodegradation Exposure time:	ated sludge dily biodegradable. : 0 % 162 d Test Guideline 301D
		Biodegradation Related to: Dis Exposure time:	ated sludge erently biodegradable. : 20 % solved organic carbon (DOC) 84 d 7 Test Guideline 302A
salicy	ylic acid:		
Biodegradability		Biodegradation Related to: Bio Exposure time: Method: OECD	ure 100 mg/l biodegradable. : 88,1 % chemical oxygen demand 14 d Test Guideline 301C nation available.
		Inoculum: activ Result: Inheren Biodegradation Related to: Dis Exposure time:	ated sludge, non-adapted tly biodegradable. : > 90 % solved organic carbon (DOC)
12.3 Bioa	ccumulative potentia	ıl	
<u>Com</u>	ponents:		
Prop	ylidynetrimethanol, p	propoxylated, reactio	n products with ammonia:
	ion coefficient: n- ol/water	: log Pow: -1,13 pH: 12,7 Method: Partitio	

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Amin	es, polyethylenepoly-,	trie	thylenetetramine	e fraction:
	on coefficient: n- ol/water	:	log Pow: -2,08 - Method: QSAR	2,90 (20 °C)
salicy	/lic acid:			
	on coefficient: n- ol/water	:	log Pow: 2,25 (2 Method: OECD	5 °C) Fest Guideline 117
2.4 Mobi	lity in soil			
Com	oonents:			
Amin	es, polyethylenepoly-,	trie	thylenetetramine	e fraction:
	oution among		Koc: 3162,28, lo	g Koc: 3,5 Fest Guideline 106
enviro	onmental compartments		Method. OECD	
salicy	/lic acid:			
	oution among		Koc: 35	
enviro	onmental compartments		Method: OECD	Fest Guideline 121
2.5 Resu	Its of PBT and vPvB as	sses	sment	
Prod	uct:			
Asses	ssment	:	to be either pers	nixture contains no components considered istent, bioaccumulative and toxic (PBT), or nd very bioaccumulative (vPvB) at levels of
2.6 Endo	crine disrupting prope	erties	S	
Prod				
Asses	ssment	:	considered to ha to REACH Article	nixture does not contain components ve endocrine disrupting properties according e 57(f) or Commission Delegated regulation or Commission Regulation (EU) 2018/605 at higher
2.7 Othe	r adverse effects			
Prod	uct:			
	onal ecological nation	:	unprofessional h	al hazard cannot be excluded in the event of andling or disposal. life with long lasting effects.

13.1 Waste treatment methods

Product

 Dispose of contents and container in accordance with all local, regional, national and international regulations.
 Do not dispose of waste into sewer.
 Do not contaminate ponds, waterways or ditches with



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		chemical or us	ed container.
Contaminated packaging		•	ng contents. unused product. empty containers.

SECTION 14: Transport information

· · ·		
14.1 UN number or ID number		
ADN	:	UN 2735
ADR	:	UN 2735
RID	:	UN 2735
IMDG	:	UN 2735
ΙΑΤΑ	:	UN 2735
14.2 UN proper shipping name		
ADN	:	POLYAMINES, LIQUID, CORROSIVE, N.O.S. (TRIETHYLENETETRAMINE)
ADR	:	POLYAMINES, LIQUID, CORROSIVE, N.O.S. (TRIETHYLENETETRAMINE)
RID	:	POLYAMINES, LIQUID, CORROSIVE, N.O.S. (TRIETHYLENETETRAMINE)
IMDG	:	POLYAMINES, LIQUID, CORROSIVE, N.O.S. (TRIETHYLENETETRAMINE)
ΙΑΤΑ	:	Polyamines, liquid, corrosive, n.o.s. (TRIETHYLENETETRAMINE)
14.3 Transport hazard class(es)		
		Class Subsidiary risks
ADN	:	8
ADR	:	8
RID	:	8
IMDG	:	8
ΙΑΤΑ	:	8
14.4 Packing group		
ADN Packing group Classification Code Hazard Identification Number Labels ADR Packing group Classification Code Hazard Identification Number	:	II C7 80 8 II C7 80



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Labe Tunr	els nel restriction code	:	8 (E)		
Clas	ting group sification Code ard Identification Number els	:	II C7 80 8		
Labe	king group	:	II 8 F-A, S-B		
Pack aircra Pack	ting instruction (LQ)		855 Y840 II Corrosive		
Pack (pas: Pack	A (Passenger) ting instruction senger aircraft) ting instruction (LQ) ting group	:	851 Y840 II Corrosive		
14.5 Envi	ironmental hazards				
ADN Envir ADR	ronmentally hazardous	:	yes		
	ronmentally hazardous	:	yes		
	ronmentally hazardous	:	yes		
IMD Marin	G ne pollutant	:	yes(TRIMETHYL TRIAMINE)	OLPROPANE POLYOXY	PROPYLENE

14.6 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.

14.7 Maritime transport in bulk according to IMO instruments

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 - List of substances subject to authorisation : Not applicable (Annex XIV)

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Con REA the r	CH - Candidate List of S cern for Authorisation (Ar CH - Restrictions on the market and use of certain ures and articles (Annex	ticle 59). manufacture, placing o dangerous substance	substances of very high concern (Regulation (EC) No 1907/2006 (REACH), Article 57). on : Conditions of restriction for the
Euro cont	eso III: Directive 2012/18, ppean Parliament and of t rol of major-accident haz gerous substances.	the Council on the	ENVIRONMENTAL HAZARDS
	upational Illnesses (R- 3, France)	: Not applicable	
prote	allations classified for the ection of the environment rironment Code R511-9)		
Take	er regulations: e note of Directive 92/85/ re applicable.	EEC regarding matern	ity protection or stricter national regulations,
	e note of Directive 94/33/ lations, where applicable		f young people at work or stricter national
The DSL	• •	-	the following inventories: f this product are on the Canadian DSL
AIIC		: On the inventory	or in compliance with the inventory
ENC	S	: On the inventory	or in compliance with the inventory
KEC	1	: On the inventory	or in compliance with the inventory
PIC	CS	: On the inventory	or in compliance with the inventory
IEC	SC	: On the inventory	or in compliance with the inventory
TCS	il	: On the inventory	or in compliance with the inventory
TSC	A	: All substances lis	ted as active on the TSCA inventory



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Inventories

AICS (Australia), AIIC (Australia), DSL (Canada), IECSC (China), ENCS (Japan), KECI (Korea), NZIOC (New Zealand), PICCS (Philippines), TCSI (Taiwan), TSCA (United States of America (USA))

15.2 Chemical safety assessment

Chemical Safety Assessments for all substances in this product are either Complete or Not applicable.

SECTION 16: Other information

i un text of in-Statements		
H302 H312 H314 H317 H318 H361d H411 H412	 Harmful if swallowed. Harmful in contact with s Causes severe skin bur May cause an allergic s Causes serious eye dar Suspected of damaging Toxic to aquatic life with Harmful to aquatic life w 	ns and eye damage. kin reaction. nage. the unborn child. long lasting effects.
Full text of other abbreviation	ons	
Acute Tox. Aquatic Chronic Eye Dam. Repr. Skin Corr. Skin Sens.	 Acute toxicity Long-term (chronic) aqu Serious eye damage Reproductive toxicity Skin corrosion Skin sensitisation 	atic hazard
Further information		
Classification of the mixture	e:	Classification procedure:
Acute Tox. 4	H302	Calculation method
Acute Tox. 4	H312	Calculation method
Skin Corr. 1B	H314	Calculation method
Eye Dam. 1	H318	Calculation method
Skin Sens. 1	H317	Calculation method
Repr. 2	H361d	Calculation method
Aquatic Chronic 2	H411	Calculation method

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THE PRODUCT MAY PRESENT HAZARDS AND SHOULD BE USED WITH CAUTION. WHILE CERTAIN HAZARDS ARE DESCRIBED IN THIS PUBLICATION, NO GUARANTEE IS MADE THAT THESE ARE THE ONLY HAZARDS THAT EXIST.

Hazards, toxicity and behaviour of the products may differ when used with other materials and are dependent upon the manufacturing circumstances or other processes. Such hazards, toxicity and behaviour should be determined by the user and made known to handlers, processors and end users.

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