according to Regulation (EC) No. 1907/2006

ARALDITE® 2011 HARDENER

Version	Revision Date:	SDS Number:
1.2	27.09.2021	400001015904



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Date of last issue: 24.01.2018 Date of first issue: 23.01.2018

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

1.1 Product identifier Trade name	: ARALDITE® 2011 HARDENER	
Unique Formula Identifier (UFI)	: ETE2-200N-1009-XXE9	
1.2 Relevant identified uses of the Use of the Substance/Mixture	e substance or mixture and uses advised against : Hardener	
1.3 Details of the supplier of the s	safety data sheet	
Company Address Telephone Telefax	 Huntsman Advanced Materials (Europe)BVBA Everslaan 45 3078 Everberg Belgium +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	r	
Emergency telephone number	: Berlin: 0049 30 19 24 0 & 0049 30 30 68 6 7 11 Bonn: 0049 228 19 27 0 & 0049 228 28 7 3 32 17 Erfurt: 0049 361 73 07 30 Freiburg: 0049 761 16 24 0 Göttingen: 0049 51 19 24 0 & 0049 551 38 31 80 Homburg: 0049 6841 19 24 0 Mainz: 0049 6131 19 24 0 & 0049 6131 23 24 66 München: 0049 89 19 24 0 Nürnberg: 0049 911 39 8 2 45 1 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: $+1/800/424.9300$)

SECTION 2: Hazards identification

2.1 Classification of the substance or mixture

Classification (REGULATION (EC) No 1272/2008)

Skin corrosion, Sub-category 1C

H314: Causes severe skin burns and eye damage.

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S	Serious	s eye damage, Categor	ry 1	1	H318:	Causes serious eye damage.
S	skin se	nsitisation, Category 1			H317:	May cause an allergic skin reaction.
2.2 La	ibel el	ements				
La	abelli	ng (REGULATION (EG	C)	No 1272/20	08)	
H	lazard	pictograms	:	E		
Si	Signal v	word	:	Danger		
H	lazard	statements	:	H314 H317		Causes severe skin burns and eye damage. May cause an allergic skin reaction.
P	recau	tionary statements	:	Prevention P261 P280 Response P301 + P3 P303 + P3 P304 + P3 P305 + P3	: 30 + P3 61 + P3 40 + P3	NOT induce vomiting. 53 IF ON SKIN (or hair): Take off immediately all contaminated clothing. Rinse skin with water. 10 IF INHALED: Remove person to fresh air and keep comfortable for breathing. Immediately call a POISON CENTER/ doctor.

Hazardous components which must be listed on the label: N'-(3-aminopropyl)-N,N-dimethylpropane-1,3-diamine

Amines, polyethylenepoly-, triethylenetetramine fraction

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



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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)
N'-(3-aminopropyl)-N,N- dimethylpropane-1,3-diamine	10563-29-8 234-148-4 01-2119970376-29	Acute Tox. 4; H302 Skin Corr. 1A; H314 Eye Dam. 1; H318 Skin Sens. 1B; H317	>= 5 - < 9,65
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 EUH071	>= 3 - < 5

For explanation of abbreviations see section 16.

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.



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In cas	e of skin contact	wounds from co difficulty.	lical treatment is necessary as untreated prrosion of the skin heal slowly and with well with water. move clothes.
In cas	e of eye contact	tissue damage In the case of c of water and se Continue rinsin Remove contac Keep eye wide	ontact with eyes, rinse immediately with plenty ek medical advice. g eyes during transport to hospital.
lf swa	llowed	If symptoms pe	•

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 Treat symptomatically.

Treatment	
rieatinent	•

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	e substance or mixture
Specific hazards during firefighting	:	Do not allow run-off from fire fighting to enter drains or water courses.
Hazardous combustion products	:	No hazardous combustion products are known
5.3 Advice for firefighters		
Special protective equipment for firefighters	:	Wear self-contained breathing apparatus for firefighting if necessary.
Specific extinguishing	:	Use extinguishing measures that are appropriate to local



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meth	ods	circumstances a	and the surrounding environment.
Furth	er information	must not be disc Fire residues ar	nated fire extinguishing water separately. This charged into drains. Ind contaminated fire extinguishing water must n accordance with local regulations.

SECTION 6: Accidental release measures

6.1 Personal precautions, protective equipment and emergency procedures

1 /1	· · · · · · · · · · · · · · · · · · ·
Personal precautions	: 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

So	utralise with acid. ak up with inert absorbent material (e.g. sand, silica gel, d binder, universal binder, sawdust). ep 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

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. Do not breathe vapours/dust. 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.
Advice on protection against fire and explosion	:	Normal measures for preventive fire protection.



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H	ygiene measures	:		ot eat or drink. When using do not smoke. ore breaks and at the end of workday.
7.2 Co	nditions for safe storage,	, inc	luding any incom	patibilities
	equirements for storage eas and containers	:	place. Containers resealed and kep	ghtly closed in a dry and well-ventilated s which are opened must be carefully of upright to prevent leakage. Observe label p in properly labelled containers.
A	dvice on common storage	:	Do not store near	r acids.
St	orage class (TRGS 510)	:	8A, Combustible,	corrosive hazardous materials
	urther information on orage stability	:	Stable under norr	mal conditions.
	ecommended storage mperature	:	2 - 40 °C	
7.3 Sp	ecific end use(s)			
-	pecific use(s)	:	No data available	9

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
N'-(3-aminopropyl)- N,N-dimethylpropane- 1,3-diamine	Workers	Inhalation	Long-term systemic effects	3,7 mg/m3
	Workers	Inhalation	Acute systemic effects	7,5 mg/m3
	Workers	Inhalation	Long-term local effects	3,7 mg/m3
	Workers	Inhalation	Acute local effects	7,5 mg/m3
	Workers	Dermal	Long-term systemic effects	0,67 mg/kg
	Consumers	Inhalation	Long-term systemic effects	0,65 mg/m3
	Consumers	Inhalation	Long-term local effects	0,65 mg/m3
	Consumers	Oral	Long-term systemic effects	0,2 mg/kg
Amines, polyethylenepoly-,	Workers	Inhalation	Long-term systemic effects	0,54 mg/m3



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triethylenetetramine fraction				
	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
N'-(3-aminopropyl)-N,N- dimethylpropane-1,3-diamine	Marine water	0,92 µg/l
	Freshwater - intermittent	92 µg/l
	Sewage treatment plant	18,1 mg/l
	Fresh water sediment	0,0336 mg/kg dry weight (d.w.)
	Marine sediment	0,0034 mg/kg dry weight (d.w.)
	Soil	0,0013 mg/kg dry weight (d.w.)
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	nt	
Eye 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	:	Solvent-resistant gloves (butyl-rubber)
Material Break through time		Nitrile rubber 10 - 480 min
Remarks	:	Chemical-resistant, impervious gloves complying with an



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		chemical produ necessary. The	dard should be worn at all times when handling acts if a risk assessment indicates this is a suitability for a specific workplace should be the producers of the protective gloves.
Skin	and body protection		thing protection according to the amount and of the dangerous substance at the work place.
Resp	piratory protection	ventilation is pr that exposures	v protection unless adequate local exhaust rovided or exposure assessment demonstrates are within recommended exposure guidelines ould conform to EN 14387
F	ilter type	: Organic vapou	r type (A)

SECTION 9: Physical and chemical properties

9.1 Information on basic physical and chemical properties

Physical state	:	liquid
Colour	:	light yellow
Odour	:	slight
Odour Threshold	:	No data is available on the product itself.
рН	:	11 Concentration: 50 %
Melting point	:	No data available
Boiling point	:	> 200 °C
Flash point	:	110 °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	:	0,04 hPa (20 °C)
Relative vapour density	:	No data is available on the product itself.
Relative density	:	No data is available on the product itself.
Density	:	0,95 g/cm3 (25 °C)

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S		ity(ies) er solubility	:	practically insolu	ble (20 °C)
	Solu	bility in other solvents	:	No data is availa	ble on the product itself.
		n coefficient: n- /water	:	No data is availa	ble on the product itself.
А	Auto-ig	nition temperature	:	No data is availa	ble on the product itself.
C	Decom	position temperature	:	> 200 °C	
V	/iscosi/ Visc	ty osity, dynamic	:	20 000 - 35 000	mPa.s (25 °C)
9.2 Of	ther ir	formation			
E	Explosi	ve properties	:	No data is availa	ble on the product itself.
C	Dxidizi	ng properties	:	No data is availa	ble on the product itself.
В	Burning	g rate	:	No data is availa	ble on the product itself.
E	Evapor	ation rate	:	No data is availa	ble on the product itself.
Γ.	Nolecu	lar weight	:	No data availabl	e

SECTION 10: Stability and reactivity

10.1 Reactivity

No dangerous reaction known under conditions of normal use.

10.2 Chemical stability

Stable under normal conditions.

10.3 Possibility of hazardous reactions

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Hazardous reactions : No hazards to be specially mentioned.
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10.4 Conditions to avoid

Conditions to avoid : None known.

10.5 Incompatible materials

Materials to avoid : None known.

10.6 Hazardous decomposition products

No decomposition if stored and applied as directed.



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SECTION 11: Toxicological information

11.1 Information on hazard classes as defined in Regulation (EC) No 1272/2008 Acute toxicity Acute oral toxicity - Product : Acute toxicity estimate : > 2 000 mg/kg Method: Calculation method Components: Amines, polyethylenepoly-, triethylenetetramine fraction: : (Rat, male and female): Exposure time: 8 h Acute inhalation toxicity Test atmosphere: vapour Method: OECD Test Guideline 403 Acute dermal toxicity -: Acute toxicity estimate : > 2 000 mg/kg Product Method: Calculation method Acute toxicity (other routes of : No data available administration) Skin corrosion/irritation Product: Result: Corrosive after 1 to 4 hours of exposure Serious eye damage/eye irritation Product: Species: Rabbit Assessment: Corrosive **Result: Corrosive** Respiratory or skin sensitisation Components: N'-(3-aminopropyl)-N,N-dimethylpropane-1,3-diamine: Test Type: Maximisation Test Exposure routes: Skin Species: Guinea pig Method: OECD Test Guideline 406 Result: The product is a skin sensitiser, sub-category 1B. GLP: yes Amines, polyethylenepoly-, triethylenetetramine fraction: Exposure routes: Skin Species: Humans Assessment: Probability or evidence of skin sensitisation in humans Result: Probability or evidence of skin sensitisation in humans No data available Assessment:

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Germ cell mutagenicity

Components:

N'-(3-aminopropyl)-N,N-dimethy Genotoxicity in vitro	 Ipropane-1,3-diamine: Test Type: in vitro assay Test system: Human lymphocytes Metabolic activation: with and without metabolic activation Method: OECD Test Guideline 487 Result: negative GLP: yes
	: Test Type: reverse mutation assay Test system: Salmonella typhimurium Metabolic activation: with and without metabolic activation Method: OECD Test Guideline 471 Result: negative
	: Test Type: In vitro mammalian cell gene mutation test Test system: mouse lymphoma cells Metabolic activation: with and without metabolic activation Method: OECD Test Guideline 476 Result: negative GLP: yes
	: 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: negative
Amines, polyethylenepoly-, trietl Genotoxicity in vitro	 hylenetetramine fraction: Test Type: In vitro mammalian cell gene mutation test Test system: Chinese hamster ovary cells Metabolic activation: with and without metabolic activation Method: OECD Test Guideline 476 Result: positive
<u>Components:</u> Amines, polyethylenepoly-, trieth Genotoxicity in vivo	hylenetetramine fraction: : Test Type: In vivo micronucleus test Test species: Mouse (male and female) Cell type: Bone marrow Application Route: Intraperitoneal injection Dose: 0 - 600 mg/kg Method: OECD Test Guideline 474

Result: negative

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Germ cell mutagenicity-Assessment : No data available

Carcinogenicity

Components:

N'-(3-aminopropyl)-N,N-dimethylpropane-1,3-diamine: Species: Mouse, male Application Route: Dermal Exposure time: 20 month(s) Dose: 1.25/56.3 mg/animal Frequency of Treatment: 3 daily No observed adverse effect level: >= 56,3 mg/kg body weight Result: negative Remarks: Information given is based on data obtained from similar substances.

Amines, polyethylenepoly-, triethylenetetramine fraction: Species: Mouse, male Dose: 42 mg/kg Frequency of Treatment: 3 daily No observed adverse effect level: >= 50 mg/kg bw/day Method: OECD Test Guideline 451 Result: negative

Species: Mouse, male Application Route: Dermal Exposure time: 104 weeks Dose: 16.8 mg/kg Frequency of Treatment: 3 daily No observed adverse effect level: >= 20 mg/kg bw/day Method: OECD Test Guideline 451

Carcinogenicity - : No data available Assessment

Reproductive toxicity

Components:

N'-(3-aminopropyl)-N,N-dimethylpropane-1,3-diamine: Effects on fertility : Test Type: Combined Repeated Dose Toxicity Study with the Reproduction / Developmental Toxicity Screening Test Species: Rat, male and female Application Route: Oral Dose: 5, 15 and 50 mg/kg bw/d General Toxicity - Parent: No observed adverse effect level: 15 mg/kg body weight General Toxicity F1: No observed adverse effect level: 15 mg/kg body weight Method: OECD Test Guideline 422 Result: Animal testing did not show any effects on fertility. GLP: yes

Components:

N'-(3-aminopropyl)-N,N-dimethylpropane-1,3-diamine:

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	ts on foetal opment	General Toxicit 15 mg/kg body	ute: Oral d 50 mg/kg bw/d ty Maternal: No observed adverse effect level: weight) Test Guideline 422
Amin	es, polyethylenepoly-,	Duration of Sin General Toxicit >= 750 mg/kg l Developmental 750 mg/kg bod Method: OECD	-natal ute: Oral 750 mg/kg bw/day gle Treatment: 10 d ty Maternal: No observed adverse effect level: body weight I Toxicity: No observed adverse effect level: >=
		Duration of Sin General Toxicit 50 mg/kg body Developmental 125 mg/kg bod	it ute: Dermal 5 mg/kg bw/day gle Treatment: 13 d ty Maternal: No observed adverse effect level: weight I Toxicity: No observed adverse effect level: >= ly weight D Test Guideline 414
N'-(3- Repro	ponents: aminopropyl)-N,N-dim oductive toxicity - ssment		nine: f adverse effects on sexual function and fertility nent, based on animal experiments.
		triethylenetetramine fr	
Repro	oductive toxicity - ssment	: The reprotoxic under further e	effects of Triethylenetetramine (TETA) are valuation as part of the EU REACH program he aminoethyl ethanolamine (AEEA) content.
	Г - single exposure ata available		
	F - repeated exposur e ata available	e	
Repe	ated dose toxicity		
•			

Components:



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N'-(3-aminopropyl)-N,N-dimethylpropane-1,3-diamine: Species: Rat, male and female NOEC: 550 Application Route: Inhalation Test atmosphere: vapour Exposure time: 3 w 6 hNumber of exposures: 5 d/w Dose: 550 mg/m3 Method: Subchronic toxicity Remarks: Based on data from similar materials

Species: Mouse, male NOAEL: >= 56,3 Application Route: Skin contact Number of exposures: 3 d Method: Chronic toxicity Remarks: Based on data from similar materials

Species: Rat, male and female NOAEL: 1000 Application Route: Oral Exposure time: 90 dMethod: OECD Test Guideline 408 Remarks: Based on data from similar materials

Amines, polyethylenepoly-, triethylenetetramine fraction: Species: Rat, male and female NOAEL: 350 mg/kg Application Route: Oral Exposure time: 28 d Number of exposures: 7 d Dose: 100/350/1000 mg/kg bw/day Method: OECD Test Guideline 407 Target Organs: Lungs Remarks: Information given is based on data obtained from similar substances.

Species: Dog, male and female NOAEL: 125 mg/kg Application Route: Oral Remarks: Information given is based on data obtained from similar substances.

Species: Dog, male and female NOAEL: 50 mg/kg Application Route: Oral Method: Subchronic toxicity Remarks: Information given is based on data obtained from similar substances.

Species: Rat, male and female NOAEL: 50 mg/kg Application Route: Oral Exposure time: 26 weeks Dose: 50/175/600 mg/kg bw/day Method: OECD Test Guideline 408 Target Organs: Lungs Remarks: Information given is based on data obtained from similar substances.

Species: Mouse, male and female NOAEL: 92 mg/kg, 600 ppm Application Route: Oral

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Exposure time: 120/600/3000 ppm Method: OECD Test Guideline 408 Remarks: Information given is based on data obtained from similar substances.

Repeated dose toxicity -Assessment : No data available

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

General Information:	No data available
Inhalation:	No data available
Skin contact:	No data available

Eye contact: No data available

- Ingestion: No data available
- **Toxicology**, **Metabolism**, **Distribution** No data available

Neurological effects No data available

Further information

Ingestion: No data available

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SECTION 12: Ecological information

12.1 Toxicity

Components: N'-(3-aminopropyl)-N,N-dimethy	Intenente 1.2 diaminat
Toxicity to fish	 LC50 (Brachydanio rerio (zebrafish)): > 100 mg/l Exposure time: 96 h Test Type: static test Analytical monitoring: yes Test substance: Fresh water Method: OECD Test Guideline 203 GLP: yes
Toxicity to daphnia and other aquatic invertebrates	 EC50 (Daphnia magna (Water flea)): 9,2 mg/l Exposure time: 48 h Test Type: static test Analytical monitoring: no Test substance: Fresh water Method: OECD Test Guideline 202 GLP: yes
Toxicity to algae/aquatic plants	 ErC50 (Selenastrum capricornutum (green algae)): 21 mg/l Exposure time: 72 h Test Type: static test Analytical monitoring: yes Test substance: Fresh water Method: OECD Test Guideline 201 GLP: yes
	NOEC (Selenastrum capricornutum (green algae)): 5,7 mg/l Exposure time: 72 h Test Type: static test Analytical monitoring: yes Test substance: Fresh water Method: OECD Test Guideline 201 GLP: yes
Toxicity to microorganisms	 EC50 (Pseudomonas putida): 181 mg/l Exposure time: 16 h Test Type: static test Analytical monitoring: no Test substance: Fresh water Method: DIN 38 412 Part 8 GLP: no
Amines, polyethylenepoly-, triet	hylenetetramine fraction:
Toxicity to fish	 LC50 (Pimephales promelas (fathead minnow)): 330 mg/l 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



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	aquatic	invertebrates		
	Toxicity plants	/ to algae/aquatic	Exposure tin Test Type: s Test substar Method: OE EC10 (Seler Exposure tin	emi-static test nce: Fresh water CD Test Guideline 201 nastrum capricornutum (green algae)): 1,34 mg/l
				nce: Fresh water CD Test Guideline 201
	Toxicity	/ to microorganisms	: NOEC (Bact Exposure tin	eria): >= 100 mg/l
			Exposure tin	eria): > 100 mg/l ne: 28 h CD Test Guideline 216
			Exposure tin Test Type: s	
			Exposure tin Test Type: s	
	aquatic	/ to daphnia and other invertebrates ic toxicity)	Exposure tin Species: Da Test Type: s Test substar	
	Toxicity organis	/ to soil dwelling sms		
		icology Assessment aquatic toxicity	: This product	has no known ecotoxicological effects.



according to Regulation (EC) No. 1907/2006

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Chronic aquatic toxicity : Harmful to aquatic life with long lasting effects.

12.2 Persistence and degradability

Components:

N'-(3-aminopropyl)-N,N-dimethylpropane-1,3-diamine:

Biodegradability : Test Type: aerobic Result: Readily biodegradable. Biodegradation: 100 % Related to: Dissolved organic carbon (DOC) Exposure time: 28 d Method: OECD Test Guideline 301A GLP: yes

Amines, polyethylenepoly-, triethylenetetramine fraction:

Result: Biodegr Exposu	m: activated sludge Not readily biodegradable. adation: 0 % re time: 162 d : OECD Test Guideline 301D
------------------------------	---

Test Type: aerobic Inoculum: activated sludge Result: Not inherently biodegradable. Biodegradation: 20 % Related to: Dissolved organic carbon (DOC) Exposure time: 84 d Method: OECD Test Guideline 302A

Chemical Oxygen Demand : 1 940 mg/g

(COD)

12.3 Bioaccumulative potential

Components:

N'-(3-aminopropyl)-N,N-dimethylpropane-1,3-diamine: Partition coefficient: noctanol/water BH: 11,6 Method: OECD Test Guideline 107

Amines, polyethylenepoly-, triethylenetetramine fraction:		
Partition coefficient: n-	:	log Pow: -2,08 - 2,90 (20 °C)
octanol/water		Method: QSAR

12.4 Mobility in soil

Components:

Amines, polyethylenepoly-, triet	hy	lenetetramine fraction:
Distribution among	:	Koc: 1584,9 - 5012
environmental compartments		Method: OECD Test Guideline 106

12.5 Results of PBT and vPvB assessment

Product:

- Assessment
- : This substance/mixture contains no components considered



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			persistent, bioaccumulative and toxic (PBT), or nt and very bioaccumulative (vPvB) at levels of er	
12.6 Endo	crine disrupting pro	perties		
Produ	<u>uct:</u>			
Asses	ssment	considered t to REACH A (EU) 2017/2	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	
12.7 Other	r adverse effects			
Produ	<u>uct:</u>			
Additi inform	onal ecological nation		ental hazard cannot be excluded in the event of al handling or disposal. quatic life.	

SECTION 13: Disposal considerations

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 chemical or used container.
Contaminated packaging	:	Empty remaining contents. Dispose of as unused product. Do not re-use 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. (DIMETHYL DIPROPYL TRIAMINE, TRIETHYLENE TETRAMINE)
ADR	:	POLYAMINES, LIQUID, CORROSIVE, N.O.S. (DIMETHYL DIPROPYL TRIAMINE, TRIETHYLENE

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			TETRAMINE)	
RID		:		LIQUID, CORROSIVE, N.O.S. ROPYL TRIAMINE, TRIETHYLENE
IMDO	3	:		LIQUID, CORROSIVE, N.O.S. ROPYL TRIAMINE, TRIETHYLENE
ΙΑΤΑ		:		id, corrosive, n.o.s. ROPYL TRIAMINE, TRIETHYLENE
14.3 Tran	sport hazard class(es)			
ADN		:	8	
ADR		:	8	
RID		:	8	
IMDO	6	:	8	
ΙΑΤΑ		:	8	
14.4 Pack	ting group			
Class	ing group sification Code rd Identification Number Is	:	III C7 80 8	
ADR Pack Class Haza Labe	ing group sification Code rd Identification Number Is el restriction code	:	III C7 80 8 (E)	
Class	ing group sification Code rd Identification Number Is	:	III C7 80 8	
IMDO Pack Labe	G ing group	:	III 8 F-A, S-B	
Pack aircra Pack	ing instruction (LQ) ing group	:	856 Y841 III Corrosive	
IATA Pack	(Passenger) ing instruction	:	852	

(passenger aircraft)

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Pao Lat	cking instruction (LQ) cking group pels	: Y841 : III : Corrosive	
14.5 En	vironmental hazards		
AD Env	N /ironmentally hazardous	: no	
AD Env	R /ironmentally hazardous	: no	
RIE Env) vironmentally hazardous	: no	

14.6 Special precautions for user

IMDG

Marine pollutant

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

: no

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 (Annex XIV)	: Not applicable
REACH - Candidate List of Substances of Very High Concern for Authorisation (Article 59).	: This product does not contain substances of very high concern (Regulation (EC) No 1907/2006 (REACH), Article 57).

Seveso III: Directive 2012/18/EU of the European Parliament and of the Council on the control of major-accident hazards involving dangerous substances. Not applicable

Water hazard class	:	WGK 1 slightly hazardous to water
(Germany)		Classification according to AwSV, Annex 1 (5.2)

Other regulations:

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

The components of this product are reported in the followin	g inventories:
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DSL : This product contains one or several components listed in the Canadian NDSL.

AIIC

: On the inventory, or in compliance with the inventory



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NZIO	0	: On the invento	ry, or in compliance with the inventory
ENCS	6	: On the invento	ry, or in compliance with the inventory
KECI		: Not in complia	nce with the inventory
PICC	S	: Not in complia	nce with the inventory
IECS	с	: On the invento	ry, or in compliance with the inventory
TCSI		: On the invento	ry, or in compliance with the inventory
TSCA	A	: All substances	listed as active on the TSCA inventory

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

Full	text	of	H-Statements	
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H302	:	Harmful if swallowed.
H312	:	Harmful in contact with skin.
H314	:	Causes severe skin burns and eye damage.
H317	:	May cause an allergic skin reaction.
H318	:	Causes serious eye damage.
H412	:	Harmful to aquatic life with long lasting effects.
EUH071	:	Corrosive to the respiratory tract.
Full text of other abbreviatio	ns	
Full text of other abbreviatio Acute Tox.		Acute toxicity
	:	Acute toxicity Chronic aquatic toxicity
Acute Tox.	:	
Acute Tox. Aquatic Chronic	:	Chronic aquatic toxicity

Further information

Classification of the mixture:

Classification procedure:

according to Regulation (EC) No. 1907/2006

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Skin C	Corr. 1C	H314	Based on product data or assessment
Eye D	am. 1	H318	Based on product data or assessment
Skin S	Sens. 1	H317	Calculation method

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