according to Regulation (EC) No. 1907/2006

ARALDITE® CW 1312 GB

Version	Revision Date:	SDS Number:	Date of
2.0	19.06.2023	400001007669	Date of



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Date of last issue: 27.06.2019 Date of first issue: 02.08.2018

Print Date 01.07.2024

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

1.1 Product identifier	
Trade name	: ARALDITE® CW 1312 GB
Unique Formula Identifier (UFI)	: ES0F-60UD-M00E-M7T1

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

Use of the	Compo	nent 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) BV 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

Emergency telephone number	Centres Antipoison et de T ANGERS: 02 41 48 21 21 BORDEAUX: 05 56 96 40 LILLE: 0 825 812 822 LYON: 04 72 11 69 11 MARSEILLE 04 91 75 25 NANCY: 03 83 32 36 36 PARIS: 01 40 05 48 48 RENNES: 02 99 59 22 22 STRASBOURG: 03 88 37 TOULOUSE: 05 61 77 74 EUROPE: +32 35 75 1234 France ORFILA: +33(0)14 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 4 USA: +1 800-424-9300	80 25 37 37 47 47 5425959

according to Regulation (EC) No. 1907/2006

ARALDITE® CW 1312 GB

Version 2.0

Revision Date: 19.06.2023

SDS Number: 400001007669



Enriching lives through innovation

Date of last issue: 27.06.2019 Date of first issue: 02.08.2018

Print Date 01.07.2024

SECTION 2: Hazards identification

2.1 Classification of the substance or mixture

Classification (REGULATION (EC) No 127	72/2008)
Skin irritation, Category 2	H315: Causes skin irritation.
Eye irritation, Category 2	H319: Causes serious eye irritation.
Skin sensitisation, Category 1	H317: May cause an allergic skin reaction.
Long-term (chronic) aquatic hazard, Category 3	H412: Harmful to aquatic life with long lasting effects.

2.2 Label elements

Labelling (REGULATION (EC) No 1272/2008)

Hazard pictograms	
Signal word	Warning
Hazard statements	 H315 Causes skin irritation. H317 May cause an allergic skin reaction. H319 Causes serious eye irritation. H412 Harmful to aquatic life with long lasting effects.
Precautionary statements	Prevention:P261Avoid breathing mist or vapours.P264Wash skin thoroughly after handling.P273Avoid release to the environment.
	P280 Wear protective gloves/ eye protection/ face protectio

Hazardous components which must be listed on the label:

2,2'-[(1-methylethylidene)bis(4,1-phenyleneoxymethylene)]bisoxirane Polypropyleneglycol diglycidylether

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

according to Regulation (EC) No. 1907/2006

ARALDITE® CW 1312 GB

Version	Re
2.0	19

evision Date: 0.06.2023 SDS Number: 400001007669 Date of last issue: 27.06.2019 Date of first issue: 02.08.2018

Print Date 01.07.2024

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

Hazardous components

Chemical name	CAS-No. EC-No. Index-No. Registration number	Classification	Concent ration (% w/w)	
2,2'-[(1-methylethylidene)bis(4,1- phenyleneoxymethylene)]bisoxir ane		Skin Irrit. 2; H315 Eye Irrit. 2; H319 Skin Sens. 1; H317 Aquatic Chronic 2; H411 specific concentration limit Skin Irrit. 2; H315 >= 5 % Eye Irrit. 2; H319 >= 5 %	>= 10 - < 20	
Polypropyleneglycol	9072-62-2	Skin Sens. 1; H317	>= 10 -	
diglycidylether	Polymer		< 20	
Substances with a workplace exp	Substances with a workplace exposure limit :			
kaolin	1332-58-7		>= 1 - <	
	310-194-1		10	

For explanation of abbreviations see section 16.

Both 25068-38-6 and 1675-54-3 can be used to describe the epoxy resin which is produced through the reaction of bisphenol A and epichlorohydrin

SECTION 4: First aid measures

4.1 Description of first aid measures

General advice	:	Move out of dangerous area. 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



according to Regulation (EC) No. 1907/2006

ARALDITE® CW 1312 GB

Version 2.0	Revision Date: 19.06.2023	SDS Number: 400001007669	Date of last issue: 27.06.2019 Date of first issue: 02.08.2018
			Print Date 01.07.2024
		mouth-to-mo	uth resuscitation.
lf inha	aled		nove to fresh air. attention if symptoms occur.
In cas	e of skin contact	lf on skin, rin	n persists, call a physician. se well with water. remove clothes.
In cas	se of eye contact	Remove con Keep eye wid	flush eye(s) with plenty of water. tact lenses. de open while rinsing. n persists, consult a specialist.
lf swa	llowed	Never give a	tory tract clear. nything by mouth to an unconscious person. persist, call a physician.
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.

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	:	Metal oxides Carbon oxides Halogenated compounds Carbon monoxide Carbon dioxide (CO2)
5.3 Advice for firefighters		
Special protective equipment	:	Wear self-contained breathing apparatus for firefighting if

necessary.



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for firefighters

according to Regulation (EC) No. 1907/2006

ARALDITE® CW 1312 GB

Version 2.0	Revision Date: 19.06.2023	SDS Number: 400001007669	Date of last issue: 27.06.2019 Date of first issue: 02.08.2018
			Print Date 01.07.2024
Specific extinguishing methods			g measures that are appropriate to local nd the surrounding environment.
Further information		must not be disc Fire residues and	ated fire extinguishing water separately. This harged into drains. d contaminated fire extinguishing water must a accordance with local regulations.

SECTION 6: Accidental release measures

6.1 Personal precautions, protect	tive	equipment and emergency procedures
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

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

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. Dispose of rinse water in accordance with local and national regulations.
Advice on protection against fire and explosion	:	Normal measures for preventive fire protection.



according to Regulation (EC) No. 1907/2006

ARALDITE® CW 1312 GB

Ver 2.0	sion	Revision Date: 19.06.2023		0S Number: 0001007669	Date of last issue: 27.06.2019 Date of first issue: 02.08.2018		
					Print Date 01.07.2024		
Hygiene measures		e 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	Conditi	ons for safe storage,	inc	uding any incomp	patibilities		
		ements for storage and containers	:	place. Containers	ghtly closed in a dry and well-ventilated which are opened must be carefully t upright to prevent leakage. Keep in properly s.		
	Advice	on common storage	:	For incompatible SDS.	materials please refer to Section 10 of this		
	Recom temper	mended storage ature	:	: 2 - 40 °C			
		r information on e stability	:	Stable under norn	nal conditions.		
7.3	Specific	c end use(s)					
	Specifi	c use(s)	:	No data available			

SECTION 8: Exposure controls/personal protection

8.1 Control parameters

Occupational Exposure Limits

Components	CAS-No.	Value type (Form of exposure)	Control parameters	Basis
calcium carbonate	471-34-1	VME	10 mg/m3	FR VLE
Further information	Indicative exposure limits			
kaolin	1332-58-7	VME	10 mg/m3	FR VLE
Further information	Indicative exposure limits			
		TWA (Respirable	0,1 mg/m3	2004/37/EC
		dust)		
Further information	on Carcinogens or mutagens			

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

	`	0 0	· · /	
Substance name	End Use	Exposure routes	Potential health effects	Value
2,2'-[(1- methylethylidene)bis(4,1- phenyleneoxymethyle ne)]bisoxirane	Workers	Inhalation	Long-term systemic effects	4,93 mg/m3
	Workers	Dermal	Long-term systemic effects	0,75 mg/kg bw/day
	Consumers	Inhalation	Long-term systemic effects	0,87 mg/m3
	Consumers	Dermal	Long-term systemic effects	0,0893 mg/kg bw/day



according to Regulation (EC) No. 1907/2006

ARALDITE® CW 1312 GB

Version 2.0	Revision Date: 19.06.2023	SDS Number: 400001007669		 last issue: 27.06.2019 first issue: 02.08.2018	
				Print Da	ate 01.07.2024
	(Consumers	Oral	Long-term systemic	0,5 mg/kg

	Consumers	Oral	Long-term systemic effects	0,5 mg/kg bw/day
calcium carbonate	Workers	Inhalation	Long-term local effects	6,36 mg/m3
	Consumers	Inhalation	Long-term local effects	1,06 mg/m3

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

Substance name	Environmental Compartment	Value
2,2'-[(1-methylethylidene)bis(4,1-	Fresh water	0,006 mg/l
phenyleneoxymethylene)]bisoxira		
ne		
	Marine water	0,001 mg/l
	Fresh water sediment	0,341 mg/kg dry
		weight (d.w.)
	Marine sediment	0,034 mg/kg dry
		weight (d.w.)
	Soil	0,065 mg/kg dry
		weight (d.w.)
	Sewage treatment plant	10 mg/l
	Secondary Poisoning	11 mg/kg

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	:	Solvent-resistant gloves (butyl-rubber)
Material Break through time	:	Nitrile rubber 10 - 480 min
Material	:	Neoprene gloves
Remarks	:	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.
Respiratory protection	:	Use respiratory protection unless adequate local exhaust ventilation is provided or exposure assessment demonstrates that exposures are within recommended exposure guidelines Equipment should conform to EN 14387



according to Regulation (EC) No. 1907/2006

ARALDITE® CW 1312 GB

Version	Revision Date: 19.06.2023	SDS Number:	Date of last issue: 27.06.2019
2.0		400001007669	Date of first issue: 02.08.2018
			Print Date 01.07.2024

Filter type : Combined particulates and organic vapour type (A-P)

SECTION 9: Physical and chemical properties

9.1 Information on basic phy Physical state	vsical and chemical properties : paste				
Colour	: off-white				
Odour	: slight				
Odour Threshold	: No data is available on the product itself.				
рН	: substance/mixture is non-soluble (in water)				
Melting point/freezing poi	nt : No data is available on the product itself.				
Boiling point	: > 200 °C				
Flash point	: > 200 °C Method: Pensky-Martens closed cup, closed cup				
Flammability (solid, gas)	: No data is available on the product itself.				
Upper explosion limit / Up flammability limit	oper : No data is available on the product itself.				
Lower explosion limit / Lo flammability limit	wer : No data is available on the product itself.				
Vapour pressure	: < 0,0001 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	: 1,85 g/cm3 (20 °C)				
Solubility(ies) Water solubility	: practically insoluble (20 °C)				
Solubility in other solve	ents : No data is available on the product itself.				
Partition coefficient: n- octanol/water	: No data is available on the product itself.				
Auto-ignition temperature	: No data is available on the product itself.				
Decomposition temperatu	re : > 200 °C				
Viscosity					



according to Regulation (EC) No. 1907/2006

ARALDITE® CW 1312 GB

Version	Revision Date: 19.06.2023	SDS Number:	Date of last issue: 27.06.2019
2.0		400001007669	Date of first issue: 02.08.2018
			Print Date 01.07.2024

Viscosity, dynamic

: 15 000 - 22 000 mPa.s (25 °C) Method: ISO 3219

9.2 Other information

No data available

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

Hazardous reactions : No hazards to be specially mentioned.

10.4 Conditions to avoid

Conditions to avoid	:	None known.
---------------------	---	-------------

10.5 Incompatible materials

Materials to avoid	: No	one known.
ivialenais lo avoid	: INC	ne known.

10.6 Hazardous decomposition products

Hazardous decomposition	: aluminium oxide
products	carbon dioxide carbon monoxide
	Halogenated compounds

SECTION 11: Toxicological information

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

Acute toxicity		
Product:		
Acute oral toxicity	:	LD50 (Rat): > 5 000 mg/kg

Components:

2,2'-[(1-methylethylidene)bis(4,1-phenyleneoxymethylene)]bisoxirane:

Acute oral toxicity	:	LD50 (Rat, female): > 2 000 mg/kg Method: OECD Test Guideline 420 Assessment: The substance or mixture has no acute oral toxicity Remarks: No mortality observed at this dose.
Acute dermal toxicity	:	LD50 (Rat, male and female): > 2 000 mg/kg Method: OECD Test Guideline 402



according to Regulation (EC) No. 1907/2006

ARALDITE® CW 1312 GB

ersion .0	Revision Date: 19.06.2023	SDS Number: 400001007669	Date of last issue: 27.06.2019 Date of first issue: 02.08.2018		
			Print Date 01.07.2024		
		Assessment: toxicity	The substance or mixture has no acute dermal		
Polyp	ropyleneglycol digl	ycidylether:			
Acute	oral toxicity		ale and female): > 2 000 mg/kg The substance or mixture has no acute oral		
		Median lethal	dose (Rat, male and female): >5 ml/kg		
Acute dermal toxicity :			ale and female): > 5 000 mg/kg The substance or mixture has no acute dermal		
Skin o	corrosion/irritation				
<u>Comp</u>	onents:				
2,2'-[(1-methylethylidene)bis(4,1-phenyleneo	kymethylene)]bisoxirane:		
	es sure time sment				

opeoleo	
Exposure time	: 4 h
Assessment	: Irritating to skin.
Method	: OECD Test Guideline 404
Result	: Irritating to skin.

Polypropyleneglycol diglycidylether:

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

Serious eye damage/eye irritation

Components:

2,2'-[(1-methylethylidene)bis(4,1-phenyleneoxymethylene)]bisoxirane:

	-			-	-	-	-
Species		:	:	Rabbit	t		
Assessment		:	:	Irritatir	ng to ey	es.	
Method		:	:	OECD	Test G	iuidelin	e 405
Result		:	:	Irritatir	ng to ey	es.	

Polypropyleneglycol diglycidylether:

	-	
Species	:	Rabbit
Assessment	:	No eye irritation
Method	:	OECD Test Guideline 405
Result	:	No eye irritation

Respiratory or skin sensitisation

Product:

Exposure routes	:	Skin
Species	:	Guinea pig
Result	:	Causes sensitisation.



according to Regulation (EC) No. 1907/2006

ARALDITE® CW 1312 GB

Version	Revision Date:
2.0	19.06.2023

SDS Number: 400001007669



Enriching lives through innovation

Date of last issue: 27.06.2019 Date of first issue: 02.08.2018

Print Date 01.07.2024

Components:

$\label{eq:2.2} 2,2'-[(1-methylethylidene)bis(4,1-phenyleneoxymethylene)] bis oxirane:$

:	Local lymph node assay (LLNA)
:	Skin
:	Mouse
:	OECD Test Guideline 429
:	The product is a skin sensitiser, sub-category 1B.

Polypropyleneglycol diglycidylether:

Exposure routes	:	Skin
Species	:	Guinea pig
Result	:	May cause sensitisation by skin contact.

Germ cell mutagenicity

Components:

2,2'-[(1-methylethylidene)bis(4,1-phenyleneoxymethylene)]bisoxirane:			
Genotoxicity in vitro :	Test Type: In vitro mammalian cell gene mutation test Test system: mouse lymphoma cells Metabolic activation: without metabolic activation Result: positive		
	Test Type: reverse mutation assay Test system: Salmonella typhimurium Metabolic activation: with and without metabolic activation Method: Mutagenicity (Salmonella typhimurium - reverse mutation assay) Result: negative		
Genotoxicity in vivo :	Test Type: in vivo assay Species: Mouse (male) Cell type: Germ Application Route: Oral Dose: 3333, 10000 mg/kg Result: negative		
	Test Type: gene mutation test Species: Rat (male) Cell type: Somatic Application Route: Oral Dose: 50,250,500,1000 mg/kg bw/day Method: OECD Test Guideline 488 Result: negative		
Polypropyleneglycol diglycidylether:			
Genotoxicity in vitro :	Test Type: Ames test Test system: Salmonella typhimurium Metabolic activation: Metabolic activation Method: OECD Test Guideline 471		

Result: positive GLP: no

according to Regulation (EC) No. 1907/2006

ARALDITE® CW 1312 GB

Version 2.0 Revision Date: 19.06.2023

SDS Number: 400001007669



Enriching lives through innovation

Date of last issue: 27.06.2019 Date of first issue: 02.08.2018

Print Date 01.07.2024

Carcinogenicity

Components:

2,2'-[(1-methylethylidene)bis(4,1-phenyleneoxymethylene)]bisoxirane:

Species Application Route Exposure time Dose Frequency of Treatment NOAEL Method Result Target Organs	 Rat, male Oral 24 month(s) 0, 2, 15, or 100 mg/kg bw/day 7 days/week 15 mg/kg bw/day OECD Test Guideline 453 negative Digestive organs
Species Application Route Exposure time Dose Frequency of Treatment NOEL Method Result Target Organs	 Mouse, male Dermal 24 month(s) 0, 0.1, 10, 100 mg/kg bw/day 3 days/week 0,1 mg/kg body weight OECD Test Guideline 453 negative Digestive organs
Species Application Route Exposure time Dose Frequency of Treatment NOEL Method Result	 Rat, female Dermal 24 month(s) 0.1, 100, 1000 mg/kg bw/day 5 days/week 100 mg/kg body weight OECD Test Guideline 453 negative
Species Application Route Exposure time Dose Frequency of Treatment NOAEL Method Result Target Organs	 Rat, female Oral 24 month(s) 0, 2, 15, or 100 mg/kg bw/day 7 days/week 100 mg/kg bw/day OECD Test Guideline 453 negative Digestive organs
Species Application Route Exposure time Dose Frequency of Treatment NOEL Method Result Target Organs	 Rat, females Oral 24 month(s) 0, 2, 15, or 100 mg/kg bw/day 7 days/week 2 mg/kg bw/day OECD Test Guideline 453 negative Digestive organs

according to Regulation (EC) No. 1907/2006

ARALDITE® CW 1312 GB

Version Revision Date: 2.0 19.06.2023

Date of last issue: 27.06.2019 Date of first issue: 02.08.2018

Print Date 01.07.2024

Reproductive toxicity

Test Type, Two generation study
 Test Type: Two-generation study Species: Rat, male and female Application Route: Oral Dose: 0, 50, 180, 540 or 750 milligram per kilogram Duration of Single Treatment: 238 d Frequency of Treatment: 1 daily General Toxicity - Parent: NOEL: 540 mg/kg body weight General Toxicity F1: NOEL: 750 mg/kg body weight Symptoms: No adverse effects Method: OECD Test Guideline 416 Result: No effects on fertility and early embryonic development were detected.
 Species: Rabbit, female Application Route: Dermal Dose: 0, 30, 100 or 300 milligram per kilogram Duration of Single Treatment: 28 d Frequency of Treatment: 1 daily General Toxicity Maternal: NOAEL: 30 mg/kg body weight Developmental Toxicity: NOAEL: 300 mg/kg body weight Method: Other guidelines Result: No teratogenic effects
Test Type: Pre-natal Species: Rabbit, female Application Route: Oral Dose: 0, 20, 60 or 180 milligram per kilogram Duration of Single Treatment: 13 d Frequency of Treatment: 1 daily General Toxicity Maternal: NOAEL: 60 mg/kg body weight Developmental Toxicity: NOAEL: 180 mg/kg body weight Method: OECD Test Guideline 414 Result: No teratogenic effects
Test Type: Pre-natal Species: Rat, female Application Route: Oral Dose: 0, 60, 180 and 540 milligram per kilogram Duration of Single Treatment: 10 d Frequency of Treatment: 1 daily General Toxicity Maternal: NOAEL: 180 mg/kg body weigh Developmental Toxicity: NOAEL: > 540 mg/kg body weight Method: OECD Test Guideline 414 Result: No teratogenic effects

STOT - repeated exposure No data available



Enriching lives through innovation

SDS Number: 400001007669

according to Regulation (EC) No. 1907/2006

ARALDITE® CW 1312 GB

Version F 2.0 1

Revision Date: 19.06.2023

SDS Number: 400001007669



Enriching lives through innovation

Date of last issue: 27.06.2019 Date of first issue: 02.08.2018

Print Date 01.07.2024

Repeated dose toxicity

Components:

2,2'-[(1-methylethylidene)bis(4,1-phenyleneoxymethylene)]bisoxirane:

Species NOAEL Application Route Exposure time Number of exposures Dose Method	:	Rat, male and female 50 mg/kg oral (gavage) 14 Weeks 7 d 0, 50, 250, 1000 mg/kg/day OECD Test Guideline 408
Species NOAEL Application Route Exposure time Number of exposures Dose Method	:	Rat, male and female >= 10 mg/kg Skin contact 13 Weeks 5 d 0, 10, 100, 1000 mg/kg/day OECD Test Guideline 411
Species NOAEL Application Route Exposure time Number of exposures Dose Method		Mouse, male 100 mg/kg Skin contact 13 Weeks 3 d 0, 1, 10, 100 mg/kg/day OECD Test Guideline 411

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

Further information

No data available

according to Regulation (EC) No. 1907/2006

ARALDITE® CW 1312 GB

Version 2.0 Revision Date: 19.06.2023

SDS Number: 400001007669

HUNTSMAN

Enriching lives through innovation

Date of last issue: 27.06.2019 Date of first issue: 02.08.2018

Print Date 01.07.2024

SECTION 12: Ecological information

12.1 Toxicity

<u>Components:</u>			
2,2'-[(1-methylethylidene)bis	s(4,	1-phenyleneoxymethylene)]bisoxirane:	
Toxicity to fish	:		
Toxicity to daphnia and other aquatic invertebrates	:	EC50 (Daphnia magna (Water flea)): 1,8 mg/l Exposure time: 48 h Test Type: static test Test substance: Fresh water Method: OECD Test Guideline 202	
Toxicity to algae/aquatic plants	:	EC50 : 11 mg/l Exposure time: 72 h Test Type: static test Test substance: Fresh water Method: EPA-660/3-75-009	
		NOEC : 4,2 mg/l Exposure time: 72 h Test Type: static test Test substance: Fresh water Method: EPA-660/3-75-009	
Toxicity to microorganisms	:	IC50 (activated sludge): > 100 mg/l Exposure time: 3 h Test Type: static test Test substance: Fresh water	
Toxicity to daphnia and other aquatic invertebrates (Chronic toxicity)	:	NOEC: 0,3 mg/l Exposure time: 21 d Species: Daphnia magna (Water flea) Test Type: semi-static test Test substance: Fresh water Method: OECD Test Guideline 211	
Ecotoxicology Assessment			
Chronic aquatic toxicity	:	Toxic to aquatic life with long lasting effects.	
Polypropyleneglycol diglycidylether:			
Toxicity to fish	:	LC50 (Danio rerio (zebra fish)): > 100 mg/l End point: mortality Exposure time: 96 h Test substance: Fresh water Method: Directive 67/548/EEC, Annex V, C.1.	
Toxicity to daphnia and other aquatic invertebrates	:	EC50 (Daphnia magna (Water flea)): > 320 mg/l End point: Immobilization	

according to Regulation (EC) No. 1907/2006

ARALDITE® CW 1312 GB

HUNTSMAN

2.0	Revision Date: 19.06.2023	SDS Number: 400001007669	Date of last issue: 27.06.2019 Date of first issue: 02.08.2018
			Print Date 01.07.202
			me: 24 h ance: Fresh water rective 67/548/EEC, Annex V, C.2.
Toxicity	y to microorganisms	Exposure t	vated sludge): > 100 mg/l me: 3 h Fresh water
12.2 Persis	stence and degrada	oility	
Compo	onents:		
2,2'-[(1	-methylethylidene)	bis(4,1-phenylend	eoxymethylene)]bisoxirane:
Biodeg	ıradability	Concentrat Result: Not Biodegrada Exposure t	activated sludge, non-adapted ion: 20 mg/l readily biodegradable. tion: 5 %
Stabilit	Stability in water :	pH: 4	n half life (DT50): 4,83 d (25 °C) ECD Test Guideline 111 Fresh water
		pH: 9	n half life (DT50): 7,1 d (25 °C) ECD Test Guideline 111 Fresh water
		pH: 7	n half life (DT50): 3,58 d (25 °C) ECD Test Guideline 111 Fresh water
Polypr	opyleneglycol digly	cidvlether	
	gradability	: Inoculum: S Concentrat Result: Not Biodegrada Related to: Exposure t	Dissolved organic carbon (DOC)
12.3 Bioaco	cumulative potentia	I	
Compo	onents:		
		bis(4,1-phenylend	eoxymethylene)]bisoxirane:
Bioacc	umulation		ration factor (BCF): 31 Does not bioaccumulate.
Partitio octano	n coefficient: n- I/water	: log Pow: 3, pH: 7,1	242 (25 °C)

according to Regulation (EC) No. 1907/2006

ARALDITE® CW 1312 GB

Version	Revision Date:	SDS Number:
2.0	19.06.2023	400001007669

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Date of last issue: 27.06.2019 Date of first issue: 02.08.2018

Print Date 01.07.2024

Method: OECD Test Guideline 117

12.4 Mobility in soil

Components:

2,2'-[(1-methylethylidene)bis(4,1-phenyleneoxymethylene)]bisoxirane:

:

Distribution among : Koc: 445 environmental compartments

12.5 Results of PBT and vPvB assessment

Product:

Assessment

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.

12.6 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

12.7 Other adverse effects

Product:

ironmental hazard cannot be excluded in the event of essional handling or disposal. Il to aquatic life with long lasting effects.
in to aquatic life with long lasting effects.

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

UNRTDG

: Not regulated as dangerous goods

according to Regulation (EC) No. 1907/2006

ARALDITE® CW 1312 GB

Vers 2.0		Revision Date: 19.06.2023	SI	DS Number: 00001007669	Date of last issue: 27.06.2019 Date of first issue: 02.08.2018	
					Print Date 01.07.2024	
	ADN		:	Not regulated as	dangerous goods	
	ADR		:	Not regulated as	dangerous goods	
	RID		:	Not regulated as	dangerous goods	
	IMDG		:	Not regulated as	dangerous goods	
	ΙΑΤΑ		:	Not regulated as	dangerous goods	
14.2	UN pro	oper shipping name				
	UNRT	DG	:	Not regulated as	dangerous goods	
	ADN		:	Not regulated as	dangerous goods	
	ADR		:	Not regulated as	dangerous goods	
	RID		:	Not regulated as	dangerous goods	
	IMDG		:	Not regulated as	dangerous goods	
	ΙΑΤΑ		:	Not regulated as	dangerous goods	
14.3	Transp	port hazard class(es)				
	ADN		:	Not regulated as	dangerous goods	
	ADR		:	Not regulated as	dangerous goods	
	RID		:	Not regulated as	dangerous goods	
	IMDG		:	Not regulated as	dangerous goods	
	ΙΑΤΑ		:	Not regulated as	dangerous goods	
14.4	Packir	ng group				
	ADN		:	Not regulated as	dangerous goods	
	ADR		:	Not regulated as	dangerous goods	
	RID		:	Not regulated as	dangerous goods	
	IMDG		:	Not regulated as	dangerous goods	
	IATA (Cargo)	:	Not regulated as	dangerous goods	
	IATA (Passenger)	:	Not regulated as	dangerous goods	
14.5 Environmental hazards						
Not regulated as dangerous goods						
14.6 Special precautions for user						
Not applicable						
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



according to Regulation (EC) No. 1907/2006

ARALDITE® CW 1312 GB

Version 2.0	Revision Date: 19.06.2023	SDS Number: 400001007669		of last issue: 27.06.2019 of first issue: 02.08.2018
				Print Date 01.07.2024
(Anne	əx XIV)			
Conc REAC the m	REACH - Candidate List of Substances of Very High Concern for Authorisation (Article 59). REACH - Restrictions on the manufacture, placing on the market and use of certain dangerous substances, mixtures and articles (Annex XVII)			 This product does not contain substances of very high concern (Regulation (EC) No 1907/2006 (REACH), Article 57). Conditions of restriction for the following entries should be considered: Number on list 75, 3
				If you intend to use this product as tattoo ink, please contact your vendor.
Europ	Seveso III: Directive 2012/18/EU of the European Parliament and of the Council on the control of major-accident hazards involving dangerous substances.			ot applicable
	pational Illnesses (R- 3, France)	: 51, 84, 25		

Other regulations:

Take note of Directive 92/85/EEC regarding maternity protection or stricter national regulations, where applicable.

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

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

DSL	: All components of this product are on the Canadian DSL
AIIC	: On the inventory, or in compliance with the inventory
NZIoC	: Not in compliance with the inventory
NZIoC	: On the inventory, or in compliance with the inventory
ENCS	: On the inventory, or in compliance with the inventory
KECI	: On the inventory, or in compliance with the inventory
PICCS	: On the inventory, or in compliance with the inventory

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according to Regulation (EC) No. 1907/2006

ARALDITE® CW 1312 GB

Version 2.0	Revision Date: 19.06.2023	SDS Number: 400001007669	Date of last issue: 27.06.2019 Date of first issue: 02.08.2018
			Print Date 01.07.2024
IECSC	2	: On the inventory	v, or in compliance with the inventory
TCSI		: On the inventory	y, or in compliance with the inventory
TSCA		: All substances li	sted 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						
H315 H317 H319 H411		Causes skin irritation. May cause an allergic skin reaction. Causes serious eye irritation. Toxic to aquatic life with long lasting effects.				
Full text of other abbreviations						
Aquatic Chronic Eye Irrit. Skin Irrit. Skin Sens. 2004/37/EC	:	Long-term (chronic) aquatic hazard Eye irritation Skin irritation Skin sensitisation Europe. Directive 2004/37/EC on the protection of workers from the risks related to exposure to carcinogens or mutagens at work				
FR VLE		France. Occupational Exposure Limits				
2004/37/EC / TWA FR VLE / VME		Long term exposure limit Time Weighted Average				
Further information						
Classification of the mixture: Classification procedure:						
Skin Irrit. 2	H3	15 Calculation method				
Eye Irrit. 2	H3	19 Calculation method				
Skin Sens. 1	H3	17 Calculation method				
Aquatic Chronic 3	H4	12 Calculation method				

20 / 21



according to Regulation (EC) No. 1907/2006

ARALDITE® CW 1312 GB

Version Revision Date: 2.0 19.06.2023

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Date of last issue: 27.06.2019 Date of first issue: 02.08.2018

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21/21

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