

Conforms to Regulation (EC) No. 1907/2006 (REACH), Annex II, as amended by Commission Regulation (EU) 2020/878

SAFETY DATA SHEET

FOR PROFESSIONAL and/or INDUSTRIAL USE ONLY

VERSATICTM Acid 10

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

1.1 Product identifier

Product name	:	VERSATIC TM Acid 10
SDS Number	:	V9113
Index number	:	Not available
EC number	:	248-093-9
CAS number	:	26896-20-8
REACH Registration number	:	01-2119449554-33-0001
Product type	:	Acid

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

Product use

Industrial use.

1.3 Details of the supplier of the safety data sheet

Manufacturer/Supplier/Importer	:	Hexion VAD B.V. Seattleweg 17 3195 ND Pernis - Rotterdam The Netherlands
Contact person Telephone 1.4	:	service@hexion.com General information +31 10 3136 500
Emergency telephone number Supplier Telephone number	:	CARECHEM24 +44 (0) 1235 239 670

SECTION 2: Hazards identification

2.1 Classification of the substance or mixture

Version: 4.0

Classification according to Regulation (EC) No. 1272/2008 [CLP/GHS]

Not classified.

See Section 16 for the full text of the H statements declared above.

2.2 Label elements

Signal word	:	No signal word.
Hazard statements	:	No known significant effects or critical hazards.
Precautionary statements		
Prevention	:	Not applicable.
Response	:	Not applicable.
Storage	:	Not applicable.
Disposal	:	Not applicable.
Hazardous ingredients	:	neodecanoic acid
Supplemental label elements	:	Not applicable.
2.3 Other hazards		
Substance meets the criteria for PBT according to Regulation (EC) No. 1907/2006, Annex XIII	:	No.
Substance meets the criteria for vPvB according to Regulation (EC) No. 1907/2006, Annex XIII	:	No.
Other hazards which do not result in classification	:	None known.

SECTION 3: Composition/information on ingredients

3.1 Substances

: UVCB

:

Product name

VERSATIC[™] Acid 10

Product/ingredient name	Identifiers	%	Classification	Specific Conc. Limits, M- factors and ATEs	Туре
neodecanoic acid	EC : 248-093-9 CAS : 26896-20-8	100	Not classified.	-	[1]

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See Section 16 for the full text of the H statements declared above.

There are no additional ingredients present which, within the current knowledge of the supplier, are classified and contribute to the classification of the substance and hence require reporting in this section.

Type

[1] Constituent

Occupational exposure limits, if available, are listed in Section 8.

SECTION 4: First aid measures

4.1 Description of first aid measures

Eye contact	:	Immediately flush eyes with plenty of water, occasionally lifting the upper and lower eyelids. Check for and remove any contact lenses. Get medical attention if irritation occurs.
Inhalation	:	Remove victim to fresh air and keep at rest in a position comfortable for breathing. Get medical attention if symptoms occur.
Skin contact	:	Flush contaminated skin with plenty of water. Remove contaminated clothing and shoes. Get medical attention if symptoms occur.
Ingestion	:	Wash out mouth with water. If material has been swallowed and the exposed person is conscious, give small quantities of water to drink. Do not induce vomiting unless directed to do so by medical personnel. Get medical attention if symptoms occur.
Protection of first aid personnel	:	No action shall be taken involving any personal risk or without suitable training.

4 3 3 4

4.2 Most important symptoms	and effects	s, both acute and delayed
Potential acute health effects		
Eye contact	:	No known significant effects or critical hazards.
Inhalation	:	No known significant effects or critical hazards.
Skin contact	:	No known significant effects or critical hazards.
Ingestion	:	No known significant effects or critical hazards.
Over-exposure signs/sympton	<u>ns</u>	
Eye contact	:	No specific data.
Inhalation	:	No specific data.
Skin contact	:	No specific data.
Ingestion	:	No specific data.
4.3 Indication of any immedia	te medical a	attention and special treatment needed
Notes to physician	:	Treat symptomatically. Contact poison treatment specialist immediately if large quantities have been ingested or inhaled.
Specific treatments	:	No specific treatment.

SECTION 5: Firefighting measures

5.1 Extinguishing media

Suitable extinguishing media

Use dry chemical, CO2, alcohol-resistant foam or water spray (fog).

:

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Unsuitable extinguishing media Do not use water jet. : 5.2 Special hazards arising from the substance or mixture Hazards from the substance or In a fire or if heated, a pressure increase will occur and the container : mixture may burst. Hazardous thermal Decomposition products may include the following materials: : carbon dioxide decomposition products carbon monoxide **5.3** Advice for firefighters Special protective actions for Promptly isolate the scene by removing all persons from the vicinity • fire-fighters of the incident if there is a fire. No action shall be taken involving any personal risk or without suitable training. Special protective equipment for Fire-fighters should wear appropriate protective equipment and self-: fire-fighters contained breathing apparatus (SCBA) with a full face-piece operated in positive pressure mode. Clothing for fire-fighters (including helmets, protective boots and gloves) conforming to European standard EN 469 will provide a basic level of protection for chemical incidents. **Additional information** Not available :

SECTION 6: Accidental release measures

6.1 Personal precautions, protective equipment and emergency procedures

For non-emergency personnel For emergency responders	:	No action shall be taken involving any personal risk or without suitable training. Evacuate surrounding areas. Keep unnecessary and unprotected personnel from entering. Do not touch or walk through spilled material. Put on appropriate personal protective equipment. If specialised clothing is required to deal with the spillage, take note of any information in Section 8 on suitable and unsuitable materials. See also the information in "For non-emergency personnel".
6.2 Environmental precautions	:	Avoid dispersal of spilled material and runoff and contact with soil, waterways, drains and sewers. Inform the relevant authorities if the product has caused environmental pollution (sewers, waterways, soil or air).
6.3 Methods and material for contain	inme	nt and cleaning up
Small spill Large spill	:	Stop leak if without risk. Move containers from spill area. Dilute with water and mop up if water-soluble. Alternatively, or if water- insoluble, absorb with an inert dry material and place in an appropriate waste disposal container. Dispose of via a licensed waste disposal contractor. Stop leak if without risk. Move containers from spill area. Prevent
		entry into sewers, water courses, basements or confined areas. Wash spillages into an effluent treatment plant or proceed as follows. Contain and collect spillage with non-combustible, absorbent material e.g. sand, earth, vermiculite or diatomaceous earth and place in container for disposal according to local regulations. Dispose of via a licensed waste disposal contractor.
6.4 Reference to other sections	:	See Section 1 for emergency contact information. See Section 8 for information on appropriate personal protective equipment. See Section 13 for additional waste treatment information.

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SECTION 7: Handling and storage

7.1 Precautions for safe handling

Protective measures	:	Put on appropriate personal protective equipment (see section 8 of SDS).
Advice on general occupational hygiene	:	Eating, drinking and smoking should be prohibited in areas where this material is handled, stored and processed. Workers should wash hands and face before eating, drinking and smoking. Remove contaminated clothing and protective equipment before entering eating areas. See also Section 8 for additional information on hygiene measures.

7.2 Conditions for safe storage, including any incompatibilities

Store in accordance with local regulations. Store in original container protected from direct sunlight in a dry, cool and well-ventilated area, away from incompatible materials (see section 10 of SDS) and food and drink. Keep container tightly closed and sealed until ready for use. Containers that have been opened must be carefully resealed and kept upright to prevent leakage. Do not store in unlabeled containers. Use appropriate containment to avoid environmental contamination.

7.3 Specific end use(s)

Recommendations	:	Not available
Industrial sector specific	:	Not available
solutions		

SECTION 8: Exposure controls/personal protection

:

8.1 Control parameters

Occupational exposure limits

No exposure limit value known.
Recommended monitoring
procedures

If this product contains ingredients with exposure limits, personal, workplace atmosphere or biological monitoring may be required to determine the effectiveness of the ventilation or other control measures and/or the necessity to use respiratory protective equipment. Reference should be made to monitoring standards, such as the following: European Standard EN 689 (Workplace atmospheres - Guidance for the assessment of exposure by inhalation to chemical agents for comparison with limit values and measurement strategy) European Standard EN 14042 (Workplace atmospheres - Guide for the application and use of procedures for the assessment of exposure to chemical and biological agents) European Standard EN 482 (Workplace atmospheres - General requirements for the performance of procedures for the measurement of chemical agents) Reference to national guidance documents for methods for the determination of hazardous substances will also be required.

DNELs/DMELs

Product/ingredie nt name	Туре	Exposure	Value	Population	Effects
neodecanoic acid	DNEL	Long term Dermal	7.41 mg/kg bw/day	Workers	Systemic
neodecanoic acid	DNEL	Long term	22.04 mg/m ³	Workers	Systemic

		Inhalation			
neodecanoic acid	DNEL	Long term	1.06 mg/kg	General	Systemic
		Dermal	bw/day	population	
neodecanoic acid	DNEL	Long term	6.52 mg/m ³	General	Systemic
		Inhalation	_	population	
neodecanoic acid	DNEL	Long term	1.88 mg/kg	General	Systemic
		Oral	bw/day	population	-

DNEL/DMEL Summary

: Not available

PNECs

Product/ingredient name	Туре	Compartment Detail	Value	Method Detail
neodecanoic acid	PNEC	Fresh water	0.478 mg/l	
neodecanoic acid	PNEC	Marine	0.0478 mg/l	
DNEC Commence	N.	4		

PNEC Summary : Not available Derived No-Effect Levels' (DNEL's) and Predicted No-Effect Concentrations' (PNEC's)

Explanatory note:

REACH requires manufacturers and importers to establish and report 'Derived No-Effect Levels' (DNEL's) for humans by inhalation, ingestion and dermal routes of exposure and 'Predicted No-Effect Concentrations' (PNEC's) for environmental exposure. DNEL's and PNEC's are established by the registrant without an official consultation process, and are not intended to be directly used for setting workplace or general population exposure limits. They are primarily used as input values in running Quantitative Risk Assessment models (like the ECETOC-TRA model).

Due to differences in calculation methodology the DNEL will tend to be lower (sometimes significantly) than any corresponding health-based OEL for that chemical substance. Further although DNEL's (and PNEC's) are an indication for setting risk reduction measures, it should be recognized that these limits do not have the same regulatory application as officially endorsed governmental OEL's.

8.2 Exposure controls

Appropriate engineering controls	:	No special ventilation requirements. Good general ventilation should be sufficient to control worker exposure to airborne contaminants. If this product contains ingredients with exposure limits, use process enclosures, local exhaust ventilation or other engineering controls to keep worker exposure below any recommended or statutory limits.
Individual protection measures		
Hygiene measures Eye/face protection	:	Wash hands, forearms and face thoroughly after handling chemical products, before eating, smoking and using the lavatory and at the end of the working period. Appropriate techniques should be used to remove potentially contaminated clothing. Wash contaminated clothing before reusing. Ensure that eyewash stations and safety showers are close to the workstation location. Safety eyewear complying with an approved standard should be used when a risk assessment indicates this is necessary to avoid exposure to liquid splashes, mists, gases or dusts. If contact is possible, the following protection should be worn, unless the assessment indicates a higher degree of protection: safety glasses with side-shields.
Skin protection		
Hand protection	:	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.

		Material: 730 Camatril Minimum break through time: 480 min
		Material: 898 Butoject Minimum break through time: 480 min Producer: This recommendation is valid only for our Product as delivered. If this product will be mixed with other substances you need to contact a supplier of CE approved protective gloves (e.g. KCL GmbH, D-36124 Eichenzell, Tel. 0049 (0) 6659 87300, Fax. 0049 (0) 6659 87155, email: vertrieb@kcl.de).
Body protection	:	Personal protective equipment for the body should be selected based on the task being performed and the risks involved and should be approved by a specialist before handling this product.
Other skin protection	:	Appropriate footwear and any additional skin protection measures should be selected based on the task being performed and the risks involved and should be approved by a specialist before handling this product.
Respiratory protection	:	Based on the hazard and potential for exposure, select a respirator that meets the appropriate standard or certification. Respirators must be used according to a respiratory protection program to ensure proper fitting, training, and other important aspects of use.
Environmental exposure controls	:	Emissions from ventilation or work process equipment should be checked to ensure they comply with the requirements of environmental protection legislation. In some cases, fume scrubbers, filters or engineering modifications to the process equipment will be necessary to reduce emissions to acceptable levels.

SECTION 9: Physical and chemical properties

9.1 Information on basic physical and chemical properties

Appearance

Physical state Color	:	Liquid Colorless/Colourless
Odor Odor threshold	:	strong Not available
рН	:	Not available
Melting point/freezing point	:	Less than -30 °C
Initial boiling point and boiling	:	270 - 280 °C
range Flash point	:	129 °C
Evaporation rate	:	Not available
Upper/lower flammability or explosive limits	:	Lower: Not available Upper: Not available
Vapor pressure	:	< 1 Pa @ 20 °C
Vapor density	:	5.9 [Air = 1]
Relative density	:	Not available

Density	:	910 kg/m3
Solubility(ies)	:	Not available (not measured)
Solubility in water	:	@ 25 °C Negligible
Partition coefficient: n- octanol/water	:	Not available (not measured)
Auto-ignition temperature	:	Not available
Decomposition temperature	:	Not available (not measured)
Viscosity	:	Dynamic: Not available
		Kinematic: 45 mm2/s @ 20 °C
Explosive properties	:	Not available (not measured)
Oxidizing properties	:	Not available (not measured)
Particle characteristics		
Median particle size	:	Not applicable.

9.2 Other information

No additional information.

SECTION 10: Stability and reactivity

10.1 Reactivity	:	Stable under normal conditions.
10.2 Chemical stability	:	The product is stable.
10.3 Possibility of hazardous reactions	:	Under normal conditions of storage and use, hazardous reactions will not occur.
10.4 Conditions to avoid	:	No specific data.
10.5 Incompatible materials	:	No specific data.
10.6 Hazardous decomposition products	:	Under normal conditions of storage and use, hazardous decomposition products should not be produced.

SECTION 11: Toxicological information

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

Acute toxicity

Product/ingredient name	Result	Species	Dose	Exposure
neodecanoic acid				
	LD50 Oral	Rat - male and female	2,066 mg/kg OECD-Guideline 401 (Acute Oral Toxicity)	-
	LD50 Oral	Rat	2,066 mg/kg	-

			OECD-Guideline			
			401 (Acute Oral			
			Toxicity)			
	LC50 Inhalation	Rat - Male	3 mg/l 403	6 h		
			Acute Inhalation			
			Toxicity			
	LD50 Dermal	Rat - male and	> 3,640 mg/kg	-		
		female				
Remarks - Dermal:	In this study male and female rats were exposed to 4 ml/kg (3640 mg/kg)					
	neodecanoic acid via an occluded dermal patch for 24 hours. After 24 hours, the					
	patch was removed and clinical observations were made once daily for 9 days.					
	There were no deaths observed in this study and there were no signs of a					
	toxicity response. It is concluded that the LD50 is greater than 3640 mg/kg.					
	LD50 Dermal	Rat	> 3,640 mg/kg	-		

Conclusion/Summary

: Not available

Acute toxicity estimates

Product/ingredient name	Oral	Dermal	Inhalation (gases)	Inhalation (vapors)	Inhalation (dusts and mists)
VERSATIC [™] Acid 10	2,066 mg/kg	N/A	N/A	N/A	N/A

Irritation/Corrosion

Product/ingredient name	Result	Species	Score	Exposure	Observation
neodecanoic acid	Skin -	Rabbit	0.72	4 hrs	24 - 72 hrs
	Erythema/Eschar				
	Skin - Edema	Rabbit	0	4 hrs	24 - 72 hrs
	eyes - Cornea	Rabbit	0		1 - 72 hrs
	opacity 405 Acute				
	Eye				
	Irritation/Corrosion				
	eyes - Iris lesion	Rabbit	0		1 - 72 hrs
	405 Acute Eye				
	Irritation/Corrosion				
	eyes - Redness of	Rabbit	0.67		1 - 72 hrs
	the conjunctivae				
	405 Acute Eye				
	Irritation/Corrosion				
	eyes - Edema of	Rabbit	0.33		1 - 72 hrs
	the conjunctivae				
	405 Acute Eye				
	Irritation/Corrosion				

Conclusion/Summary

Skin	: Not available
eyes	: Not available
Respiratory	: Not available

Sensitization

Product/ingredient name	Route of exposure	Species	Result			
neodecanoic acid	Skin	Guinea pig	Not sensitizing			
Remarks:	In this study, neodecanoic acid was examined for skin sensitization potential in					
	the guinea pig maximization procedure of Magnusson and Kligman.					
	A preliminary screen was carried out to determine the concentrations of test					
	material to be used for intradermal induction, topical induction, and topical					

	challenge. Two male and female guinea pigs were used for each test
	concentration.
	Groups of ten male and ten female guinea pigs were used for the test and a
	further five males and five females as controls.
	Induction was accomplished in two stages.
	1) Intradermal injection
	Two rows of three injections were made, one on each side of the midline in the
	shorn skin of the shoulder region.
	2) Topical application
	One week after the intradermal injections, the same area was clipped free from
	hair. A 4x4 cm patch of filter paper was soaked in a solution of the test material
	and placed over the injection sites and covered with an occlusive dressing. The
	dressing was left in place for 48 hours.
	The challenge procedure was carried out two weeks after topical induction.
	Challenge was accomplished by topical application of the test material to the
	flank of animals via an occluded patch. The challenge lasted 24 hours.
	Immediately after the challenge and then again at 24 and 48 hours later each
	animals was examined for signs of skin sensitization
	At no point was there any avidence of skin sensitization produced by
	At no point was there any evidence of skin sensitization produced by
<u> </u>	
Conclusion/Summary	
Skin	: Not available

Skin Respiratory

Not available Not available :

Mutagenicity

Product/ingredient name	Test	Experiment	Result			
neodecanoic acid	-	Subject: See Remarks	Negative			
Remarks:	The test substance Versatic 10 (neodecanoic acid) was examined for its					
	potential to induce structural chromosome aberrations in cultured human					
	lymphocytes in both the ab	sence and presence of a meta	bolic activation system			
	(S9 mix), in compliance wi	th OECD guideline 473.				
	Two independent chromoso	ome aberration tests were cor	nducted in both the			
	absence and presence of S9	. In the absence of S9, cells v	were exposed to the test			
	substance continuously for 24 or 48 hours. In the presence of the S9, cells were					
	exposed to the test substance for 3 hours and harvested at 24 or 48 hours later.					
	The choice for the highest concentrations scored was based on toxicity. The test					
	substance was dissolved in DMSO.					
	In neither chromosome aberration assay, Versatic 10 did not induce a					
	statistically significant increase in the percentage of cells with structural					
	chromosome aberrations at any of the concentrations and time points analyzed.					
	The positive controls gave appropriate responses.					
	It is concluded that Versation	c 10 is not clastogenic under	the conditions used in			
	this study.					
Conclusion/Summary	: Not available					

: Not available

Carcinogenicity

Product/ingredient name	Result	Species	Dose	Exposure	
neodecanoic acid	Negative	See Remarks			
Remarks:	Justification for data waiving:				
	Testing for carcinogenicity does not appear scientifically necessary. The data				
	generated in the repeated dose dermal toxicity test are adequate for the purposes				
	of classification and labeling and indicate there is no systemic toxicity.				
	Neoacids do not bioaccumulate and are readily excreted. Neodecanoic acid does				
	not have any structural alerts for carcinogenicity (ToxTree v1.5) and was				
	negative in several tests that assess genetic toxicity. Additionally, neodecanoic				
	acid has a low pote	ntial for acute toxicit	у.		
Conclusion/Summary	Not ava	ilable			

Conclusion/Summary

Reproductive toxicity

Product/ingredient name	Maternal	Fertility	Developmen	Species	Dose	Exposure
	toxicity		t toxin			
neodecanoic acid	Negative	Negative	Negative	Rat	Oral	-
Remarks:	This study wa	s conducted to	evaluate the eff	ects of long-ter	m ingestion of 1	neodecanoic
	acid on reproc	luction in albin	o rats. Neodeca	noic acid was a	dministered in t	he diet at
	levels of 100,	500, and 1500	ppm fed to the	rats through two	parental and to	o two-litter
	filial generation	ons. Following	nine weeks of d	lietary administ	ration to the F2	B weanlings
	designated as the third parental generation, the study was terminated. There was no					
	evidence at any test level of an adverse effect on the survival, appearance, behavior,					
	body weight gain, and food consumption of the parental generations; on the reproductive					
	performance of the parents reflected by the various indices; or on the growth,					
	appearance, and behavior of the offspring. Gross and macroscopic pathological findings					
	revealed no ev	vidence of a con	mpound-related	effect at any of	the dietary leve	els.
Conclusion/Summary	: 1	Not available				

<u>Teratogenicity</u>

Product/ingredient name	Result	Species	Dose	Exposure		
neodecanoic acid	Negative - Oral	Rat	-	-		
Remarks:	In this study, pregn	ant rats, n=22 per do	se, were treated by or	ral gavage to 50,		
	250, 600 or 800 mg	g/kg/day Neoheptanoi	ic acid during gestati	on days 6-15. On		
	gestation day 21, the dams were euthanized and the pups were examined for					
	signs of developme	ental toxicity. Under t	he conditions of the	experimental		
	methods, the test m	naterial produced mat	ernal toxicity at dose	levels of 600 and		
	800 mg/kg with ma	800 mg/kg with maternal lethality at 800 mg/kg. The test material was severely				
	embryotoxic at 800) mg/kg with less that	n 20% of embryos su	rviving. Offspring		
	of the 800 mg/kg g	roup had reduced boo	ly weight, reduced ci	own-rump		
	distance, displayed	variations signifying	delayed development	nt, and a significant		
	percentage (25%) v	were malformed. In th	ne 600 mg/kg group,	there were an		
	increase number of dams with 3 or more resorptions. Offspring of the 600					
	mg/kg group displayed significant incidences of major (hydrocephalus) and					
	minor (knobby or angular ribs, extra lumbar vertebrae) malformations but					
	showed few signs of delayed development and were not runted.					
	There was no statis	stically significant evi	dence of maternal to	xicity at dose		
	levels of 50 or 250	mg/kg. There was a s	slight, but not statisti	cally significant,		
	increase in embryonic resorption noted for the 250 mg/kg group. There was no					
	statistically significant evidence of developmental toxicity at doses for 50 or					
	250 mg/kg. The NOAEL for maternal toxicity is 600 mg/kg and the NOAEL for					
	developmental toxi	icity is 250 mg/kg.				
Conclusion/Summary	: Not ava	ailable				

Specific target organ toxicity (single exposure) Not available

Specific target organ toxicity (repeated exposure) Not available

Aspiration hazard	
Not available	

Information on likely routes of : Not available exposure

Potential acute health effects

Eye contact	:	No known significant effects or critical hazards.
Inhalation	:	No known significant effects or critical hazards.

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Skin contact	:	No known significant effects or critical hazards.
Ingestion	:	No known significant effects or critical hazards.

Symptoms related to the physical, chemical and toxicological characteristics

Eye contact	:	No specific data.
Inhalation	:	No specific data.
Skin contact	:	No specific data.
Ingestion	:	No specific data.

Delayed and immediate effects as well as chronic effects from short and long-term exposure

Short term exposure

Potential immediate effects	:	Not available
Potential delayed effects	:	Not available
Long term exposure		

Potential immediate effects	:	Not available
Potential delayed effects	:	Not available

Potential chronic health effects

Product/ingredient name	Result	Species	Dose	Exposure
neodecanoic acid	NOAEL Oral	Rat	100 mg/kg/d	90 days 7 days
			Repeated dose	per week
			408 Repeated	
			Dose 90-Day	
			Oral Toxicity	
			Study in Rodents	
Conclusion/Summary	: Not av	ailable		
General	: No kno	own significant effect	ts or critical hazards.	
Carcinogenicity	: No kno	own significant effect	ts or critical hazards.	
Mutagenicity	: No kno	own significant effect	ts or critical hazards.	
Reproductive toxicity	: No kno	own significant effect	ts or critical hazards.	
1.2. Information on other haz	ards			
11.2.1 Endocrine disrupting p	roperties :	Not available		

11.2.2 Other information :

: Not available

SECTION 12: Ecological information

12.1 Toxicity

Product/ingredient name	Result	Species	Exposure
neodecanoic acid			
	Acute LC50 > 100 mg/l -	Rainbow trout, donaldson	96 h
		trout	
	Acute NOEC=No Observed	Fish	30 d
	Effect Concentration 1.6 mg/l		
	Fresh water		
	Acute EC50 > 100 mg/l Fresh	Daphnia	48 h
	water		
	Acute EC50 > 100 mg/l Fresh	Daphnia	48 h
	water		
	Acute Inhibition concentration	Alga	72 h

to 50% of test organisms > 100		
mg/l 201 Alga, Growth		
Inhibition Test		
Chronic NOEC=No Observed	Rainbow trout, donaldson	-
Effect Concentration > 2.22	trout	
mg/l Fresh water 305		
Bioconcentration: Flow-through		
Fish Test		
Chronic NOEC=No Observed	Daphnia	-
Effect Concentration 4.78 mg/l	_	
Fresh water other methods		

Conclusion/Summary

: Not available

12.2 Persistence and degradability

Product/ingredient name	Test	Result	Dose	Inoculum	
neodecanoic acid	301F Ready	11 % - The	-	Activated sludge	
	Biodegradability	product is not		_	
	- Manometric	readily			
	Respirometry	biodegradable			
	Test	28 d			
Remarks:	Neodecanoic acid was not inherently biodegradable under the conditions of the				
	study.				
Conclusion/Summary	: Not ava	ailable			

12.3 Bioaccumulative potential

Product/ingredient name	LogPow	BCF	Potential
neodecanoic acid	2.1	< 225	low

12.4 Mobility in soil

Soil/water partition coefficient (KOC)	:	Not available
Mobility	:	Not available

12.5 Results of PBT and vPvB assessment

Product/ingredient name	PBT	Р	В	Т	vPvB	vP	vB
neodecanoic acid	No	N/A	No	No	No	N/A	No
12.6 Endocrine disrupting p	oroperties	: No	t available				
12.7 Other adverse effects		: No No	: No known significant effects or critical hazards. No known significant effects or critical hazards.				

SECTION 13: Disposal considerations

13.1 Waste treatment methods

Product

Methods of disposal	:	The generation of waste should be avoided or minimized wherever
		possible. Disposal of this product, solutions and any by-products
		should at all times comply with the requirements of environmental

Hazardous waste	:	protection and waste disposal legislation and any regional local authority requirements. Dispose of surplus and non-recyclable products via a licensed waste disposal contractor. Waste should not be disposed of untreated to the sewer unless fully compliant with the requirements of all authorities with jurisdiction. Within the present knowledge of the supplier, this product is not regarded as hazardous waste, as defined by EU Directive 2008/98/EC.
Packaging		
Methods of disposal	:	The generation of waste should be avoided or minimized wherever possible. Waste packaging should be recycled. Incineration or landfill should only be considered when recycling is not feasible.
Special precautions	:	This material and its container must be disposed of in a safe way. Empty containers or liners may retain some product residues. Avoid dispersal of spilled material and runoff and contact with soil, waterways, drains and sewers.

SECTION 14: Transport information

Regulatory information	14.1. UN number	14.2. UN pr	oper shipping name	14.3. Transport hazard class(es)	14.4. Packing group
ADR/ADN		Non-regulate	ed		
RID		Non-regulated	l		
ADN		Non-regulate	ed		
ICAO/IATA		Non-regulate	ed		
IMO/IMDG		Non-regulate	ed		
14.5. Environm	iental hazard	ls			
Environmentally	/ hazardous a	nd/or Marine	Pollutant :	No.	
14.6 Special pre	cautions for a	user :	Transport within user's pro- containers that are upright transporting the product kn or spillage.	emises: always transpo and secure. Ensure tha now what to do in the e	rt in closed at persons event of an accident

SECTION 15: Regulatory information

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

<u>EU Regulation (EC) No. 1907/2006 (REACH)</u> Annex XIV - List of substances subject to authorization Conforms to Regulation (EC) No. 1907/2006 (REACH), Annex II, as amended by Regulation (EU) No. 2015/830 VERSATIC[™] Acid 10 Page:15/16

Annex XIV None required.	
Substances of very high concern None required.	
Annex XVII - Restrictions on the manufacture, placing on the market and use of certain dangerous substances, mixtures and articles	: Not applicable.
Other EU regulations	
REACH Status	: The substance(s) in this product has (have) been Registered, or are exempted from registration, according to Regulation (EC) No. 1907/2006 (REACH).
Prior Informed Consent (PIC) (64 None required.	<u>9/2012/EU)</u>
<u>Seveso Directive</u> This product is not controlled under	the Seveso Directive.
National regulations	
International regulations	
International lists : Austral Canada Japan i China i Korea i New Z Philipp United Taiwar Thailar Vietnar	ia inventory (AIIC) This material is listed or exempted. inventory This material is listed or exempted. inventory This material is listed or exempted. inventory (IECSC) This material is listed or exempted. ealand Inventory (NZIOC) This material is listed or exempted. ines inventory (PICCS) This material is listed or exempted. States inventory (TSCA 8b) This material is active or exempted. inventory (TCSI) This material is listed or exempted. inventory (TCSI) This material is listed or exempted. inventory (NOTIOC) This material is listed or exempted. inventory (TCSI) This material is listed or exempted. inventory (NOTIOC) This material is listed or exempted. inventory (TCSI) This material is listed or exempted. inventory Not determined.
15.2 Chemical Safety Assessment	: Not available

SECTION 16: Other information

Abbreviations and acronyms	:	ATE = Acute Toxicity Estimate
		CLP = Classification, Labelling and Packaging Regulation
		[Regulation (EC) No. 1272/2008]
		DMEL = Derived Minimal Effect Level
		DNEL = Derived No Effect Level
		EUH statement = CLP-specific Hazard statement
		N/A = Not available
		PBT = Persistent, Bioaccumulative and Toxic
		PNEC = Predicted No Effect Concentration

RRN = REACH Registration Number SGG = Segregation Group vPvB = Very Persistent and Very Bioaccumulative

Procedure used to derive the classification according to Regulation (EC) No. 1272/2008 [CLP/GHS] Not classified.

Full text of abbreviated H statements

Not applicable.

Full text of classifications [CLP/GHS]

Not applicable.

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