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

1.1 Product identifier Trade name	:	Sitagliptin / Metformin Extended Release Formulation
1.2 Relevant identified uses of	the s	substance or mixture and uses advised against
Use of the Sub- stance/Mixture	:	Pharmaceutical
Recommended restrictions on use	:	Not applicable
1.3 Details of the supplier of th	e saf	fety data sheet
Company	:	MSD Kilsheelan Clonmel Tipperary, IE
Telephone	:	353-51-601000
E-mail address of person responsible for the SDS	:	EHSDATASTEWARD@msd.com
1.4 Emergency telephone num	ber	

+1-908-423-6000

SECTION 2: Hazards identification

2.1 Classification of the substance or mixture

Classification (REGULATION (EC) No 1272/2008)Acute toxicity, Category 4H302: Harmful if swallowed.

2.2 Label elements

Labelling (REGULATION (EC) No 1272/2008)

Hazard pictograms	:	
Signal word	:	Warning
Hazard statements	:	H302 Harmful if swallowed.
Precautionary statements	:	Prevention:



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			n thoroughly after handling. t, drink or smoke when using this product.
			330 IF SWALLOWED: Call a POISON if you feel unwell. Rinse mouth.

Hazardous components which must be listed on the label: metformin hydrochloride

2.3 Other hazards

This substance/mixture contains no components considered to be either persistent, bioaccumulative and toxic (PBT), or very persistent and very bioaccumulative (vPvB) at levels of 0.1% or higher.

Ecological information: The substance/mixture does not contain components considered to have endocrine disrupting properties according to REACH Article 57(f) or Commission Delegated regulation (EU) 2017/2100 or Commission Regulation (EU) 2018/605 at levels of 0.1% or higher.

Toxicological information: The substance/mixture does not contain components considered to have endocrine disrupting properties according to REACH Article 57(f) or Commission Delegated regulation (EU) 2017/2100 or Commission Regulation (EU) 2018/605 at levels of 0.1% or higher.

Dust contact with the eyes can lead to mechanical irritation. Contact with dust can cause mechanical irritation or drying of the skin. May form explosive dust-air mixture during processing, handling or other means.

SECTION 3: Composition/information on ingredients

3.2 Mixtures

Components

Chemical name	CAS-No. EC-No. Index-No. Registration number	Classification	Concentration (% w/w)
metformin hydrochloride	1115-70-4 214-230-6	Acute Tox. 4; H302	>= 50 - < 70
Sitagliptin	654671-77-9	Eye Irrit. 2; H319	>= 1 - < 10

For explanation of abbreviations see section 16.

SECTION 4: First aid measures

4.1 Description of first aid measures

General advice : In the case of accident or if you feel unwell, seek medical ad-

Commission Regulation (EU) 2020/878



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			vice immediately. When symptoms advice.	persist or in all cases of doubt seek medical		
Protection of first-aiders		:	and use the recor	First Aid responders should pay attention to self-protection, and use the recommended personal protective equipment when the potential for exposure exists (see section 8).		
lf inha	aled	:	If inhaled, remove Get medical atter	e to fresh air. tion if symptoms occur.		
In cas	e of skin contact	:	Wash with water and soap. Get medical attention if symptoms occur.			
In cas	In case of eye contact		If in eyes, rinse well with water. Get medical attention if irritation develops and persists.			
If swallowed		:	If swallowed, DO NOT induce vomiting unless directed to do so by medical personnel. Get medical attention. Rinse mouth thoroughly with water. Never give anything by mouth to an unconscious person.			
4.2 Most i	mportant symptoms a	and	effects, both acute	e and delayed		
Risks		:	Harmful if swallow	ved.		
			Contact with dust can cause mechanical irritation or the skin. Dust contact with the eyes can lead to mechanical irr			
4.3 Indica	tion of any immediate	me	dical attention and	special treatment needed		
Treat	ment	:	Treat symptomati	cally and supportively.		

SECTION 5: Firefighting measures

5.1 Extinguishing media		
Suitable extinguishing media	:	Water spray Alcohol-resistant foam Carbon dioxide (CO2) Dry chemical
Unsuitable extinguishing media	:	None known.

5.2 Special hazards arising from the substance or mixture

Specific hazards during fire-	:	Avoid generating dust; fine dust dispersed in air in sufficient
fighting		concentrations, and in the presence of an ignition source is a
		potential dust explosion hazard.



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	Hazard ucts	lous combustion prod-	:	Exposure to comb Carbon oxides Metal oxides Nitrogen oxides (1	oustion products may be a hazard to health.
5.3 A	Advice	for firefighters		Silicon oxides	
	Specia for firef	l protective equipment ighters	:		e, wear self-contained breathing apparatus. ective equipment.
	Specifi ods	c extinguishing meth-	:	cumstances and t Use water spray t	measures that are appropriate to local cir- he surrounding environment. o cool unopened containers. ged containers from fire area if it is safe to do

SECTION 6: Accidental release measures

6.1 Personal precautions, protective equipment and emergency procedures

Personal precautions	:	Use personal protective equipment. Follow safe handling advice (see section 7) and personal pro- tective equipment recommendations (see section 8).
6.2 Environmental precautions		
Environmental precautions	:	Avoid release to the environment. Prevent further leakage or spillage if safe to do so. Retain and dispose of contaminated wash water. Local authorities should be advised if significant spillages cannot be contained.

6.3 Methods and material for containment and cleaning up

Methods for cleaning up	 Sweep up or vacuum up spillage and collect in suitable container for disposal. Avoid dispersal of dust in the air (i.e., clearing dust surfaces with compressed air). Dust deposits should not be allowed to accumulate on surfaces, as these may form an explosive mixture if they are released into the atmosphere in sufficient concentration. Local or national regulations may apply to releases and disposal of this material, as well as those materials and items employed in the cleanup of releases. You will need to determine which regulations are applicable. Sections 13 and 15 of this SDS provide information regarding certain local or national requirements.
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6.4 Reference to other sections

See sections: 7, 8, 11, 12 and 13.

SECTION 7: Handling and storage

7.1 Precautions for safe handling

	5		
Technical measures	:	Static electricity may accumulate and ignite suspended dust causing an explosion. Provide adequate precautions, such as electrical grounding and bonding, or inert atmospheres.	
Local/Total ventilation		Use only with adequate ventilation.	
Advice on safe handling	:	Do not breathe dust.	
Advice on sale handling	•	Do not swallow.	
		Avoid contact with eyes.	
		Avoid prolonged or repeated contact with skin.	
		Wash skin thoroughly after handling.	
		Handle in accordance with good industrial hygiene and safety practice, based on the results of the workplace exposure assessment	
		Minimize dust generation and accumulation.	
		Keep container closed when not in use.	
		Keep away from heat and sources of ignition.	
		Take precautionary measures against static discharges.	
		Do not eat, drink or smoke when using this product.	
		Take care to prevent spills, waste and minimize release to the environment.	
Hygiene measures	:	If exposure to chemical is likely during typical use, provide eye flushing systems and safety showers close to the working place. When using do not eat, drink or smoke. Wash contami- nated clothing before re-use.	
		The effective operation of a facility should include review of	
		engineering controls, proper personal protective equipment, appropriate degowning and decontamination procedures, industrial hygiene monitoring, medical surveillance and the use of administrative controls.	
7.2 Conditions for safe storage	, incl	luding any incompatibilities	
Requirements for storage areas and containers	:	Keep in properly labelled containers. Store in accordance with the particular national regulations.	
Advice on common storage	:	Do not store with the following product types: Strong oxidizing agents	
7.3 Specific end use(s)			
Specific use(s)	:	No data available	
• • • • • • • • • • • • • • • • • • • •			



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SECTION 8: Exposure controls/personal protection

8.1 Control parameters

Occupational Exposure Limits

Dust

5 mg/m3 Value type (Form of exposure): TWA (respirable dust) Basis: FOR-2011-12-06-1358

10 mg/m3 Value type (Form of exposure): TWA (total dust) Basis: FOR-2011-12-06-1358

Components	CAS-No.	Value type (Form of exposure)	Control parameters	Basis
metformin hydro- chloride	1115-70-4	TWA	1 mg/m3 (OEB 1)	Internal
Sitagliptin	654671-77- 9	TWA	0.5 mg/m3 (OEB 2)	Internal

8.2 Exposure controls

Engineering measures

Use feasible engineering controls to minimize exposure to compound. All engineering controls should be implemented by facility design and operated in accordance with GMP principles to protect products, workers, and the environment.

Personal protective equipment

Eye/face protection	:	Wear safety glasses with side shields or goggles. If the work environment or activity involves dusty conditions, mists or aerosols, wear the appropriate goggles. Wear a faceshield or other full face protection if there is a potential for direct contact to the face with dusts, mists, or aerosols.
Hand protection Material	:	Chemical-resistant gloves
Skin and body protection Respiratory protection	:	Work uniform or laboratory coat. If adequate local exhaust ventilation is not available or expo- sure assessment demonstrates exposures outside the rec- ommended guidelines, use respiratory protection. Equipment should conform to NS EN 143
Filter type	:	Particulates type (P)

SECTION 9: Physical and chemical properties

9.1 Information on basic physical and chemical properties

Physical state : powder



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C	Colour		:	blue green	
C	Ddour		:	No data available)
C	Ddour T	hreshold	:	No data available)
Ν	Melting	point/freezing point	:	No data available)
	nitial bo ange	piling point and boiling	:	No data available	
F	lamma	ability (solid, gas)	:	May form explosi dling or other me	ve dust-air mixture during processing, han- ans.
F	lamma	ability (liquids)	:	No data available)
		explosion limit / Upper bility limit	:	No data available	
		explosion limit / Lower bility limit	:	No data available	
F	-lash p	pint	:	Not applicable	
A	Auto-igr	nition temperature	:	No data available)
C	Decomp	oosition temperature	:	No data available)
p	эΗ		:	No data available)
V	/iscosit/ Visc	y osity, kinematic	:	Not applicable	
S	Solubilit Wate	y(ies) er solubility	:	No data available	
	Partitior	n coefficient: n- Water	:	Not applicable	
V	/apour	pressure	:	Not applicable	
F	Relative	edensity	:	No data available)
C	Density		:	No data available	9
F	Relative	e vapour density	:	Not applicable	
F		characteristics cle size	:	No data available	9



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9.2 Other information

Explosives	:	Not explosive
Oxidizing properties	:	The substance or mixture is not classified as oxidizing.
Evaporation rate	:	Not applicable
Molecular weight	:	No data available

SECTION 10: Stability and reactivity

10.1 Reactivity

Not classified as a reactivity hazard.

10.2 Chemical stability

Stable under normal conditions.

10.3 Possibility of hazardous reactions

Hazardous reactions	: May form explosive dust-air mixture during processing, han- dling or other means. Can react with strong oxidizing agents.
10.4 Conditions to avoid Conditions to avoid	: Heat, flames and sparks. Avoid dust formation.

10.5 Incompatible materials

Materials to avoid : Oxidizing agents

10.6 Hazardous decomposition products

No hazardous decomposition products are known.

SECTION 11: Toxicological information

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

Informati exposure	on on likely routes of	:	Inhalation Skin contact Ingestion Eye contact	
Acute to	xicity			
Harmful i	f swallowed.			
– – – – –				

Product:

Acute oral toxicity

: Acute toxicity estimate: 1.588 mg/kg Method: Calculation method



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	<u>Comp</u>	onents:			
	metfo	rmin hydrochloride:			
	Acute	oral toxicity	:	LD50 (Rat): 1.000) mg/kg
				LD50 (Mouse): 1.	450 - 3.500 mg/kg
				LD50 (Monkey): 4	163 mg/kg
				LD50 (Rabbit): 35	50 mg/kg
				LD50 (Guinea pig	ı): 500 mg/kg
	Oite all				
	Sitagli Acute	oral toxicity	:	LD50 (Rat): > 3.0	00 mg/kg
				LD50 (Mouse): 3.	000 mg/kg
	Skin corrosion/irritation Not classified based on available information.				
	<u>Comp</u>	onents:			
		rmin hydrochloride:		D 11 %	
	Specie Result		:	Rabbit Mild skin irritation	
	Sitagl	iptin:			
	Specie	es	:	Rabbit	
	Metho Result		:	Draize Test No skin irritation	
	Serious eye damage/eye irritation Not classified based on available information.				
		onents:			
	Specie	rmin hydrochloride:	:	Rabbit	
	Result		:	Mild eye irritation	
	Sitagl	iptin:			
	Specie Metho		:	Rabbit Droizo Toot	
	Result		:	Draize Test Irritating to eyes.	



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Respiratory or skin sensitisation

Skin sensitisation

Not classified based on available information.

Respiratory sensitisation

Not classified based on available information.

Components:

Sitagliptin:

Test Type	:	Local lymph node assay (LLNA)
Species	:	Mouse
Method	:	OECD Test Guideline 429
Result	:	Not a skin sensitizer.

Germ cell mutagenicity

Not classified based on available information.

Components:

metformin hydrochloride:		
Genotoxicity in vitro		Test Type: Bacterial reverse mutation assay (AMES) Result: negative
		Test Type: in vitro assay Test system: mouse lymphoma cells Result: negative
		Test Type: Chromosomal aberration Test system: Human lymphocytes Result: negative
Genotoxicity in vivo	:	Test Type: Micronucleus test Species: Mouse Application Route: Oral Result: negative
Sitagliptin:		
Genotoxicity in vitro	:	Test Type: Ames test Result: negative
		Test Type: Chromosome aberration test in vitro Test system: Chinese hamster ovary cells Result: negative
		Test Type: DNA damage and repair, unscheduled DNA syn- thesis in mammalian cells (in vitro) Test system: rat hepatocytes Result: negative



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Gen	otoxicity in vivo	: Test Type: Mi Species: Mou Application Re Result: negati	oute: Oral
	cinogenicity classified based on avail	able information.	
Con	nponents:		
met	formin hydrochloride:		
Spe Exp Dos Res	osure time e	: Mouse : 91 weeks : 1500 mg/kg b : negative	ody weight
	lication Route osure time e	: Rat, male : Oral : 104 weeks : 900 mg/kg bo : negative	dy weight
Exp LOA Res Tarç	lication Route osure time \EL	 Rat, female Oral 104 weeks 900 mg/kg bo negative Uterus (includ) The mechanis mans. 	
Sita	gliptin:		
Spe App	cies lication Route osure time	: Mouse : Oral : 2 Years : negative	
Exp Res Tarç Ren	lication Route osure time ult get Organs narks	0	icity observed in testing
Care mer	cinogenicity - Assess- It	: Weight of evic cinogen	lence does not support classification as a car-

Reproductive toxicity

Not classified based on available information.



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	<u>Comp</u>	onents:			
	metfor	min hydrochloride:			
		on fertility	:	Test Type: Fertilit Species: Rat Application Route Fertility: NOAEL: Result: No effects	: Oral 600 mg/kg body weight
	Effects ment	on foetal develop-	:	Test Type: Develo Species: Rat Application Route Developmental To Result: No teratog	: Oral oxicity: NOAEL: 600 mg/kg body weight
				Species: Rabbit Application Route	icity: NOAEL: 140 mg/kg body weight
	Sitagli	ptin:			
	-	on fertility	:	Species: Rat Application Route Fertility: NOAEL F	y/early embryonic development : Oral Parent: 1.000 mg/kg body weight sting did not show any effects on fertility.
	Effects ment	on foetal develop-	:	Species: Rat Application Route Teratogenicity: LC Result: Embryoto spring were detec Test Type: Embry Species: Rabbit	o-foetal development : Oral DAEL: 250 mg/kg body weight kic effects and adverse effects on the off- ted., No teratogenic effects ro-foetal development DAEL: 125 mg/kg body weight
				Result: No teratog	
		 single exposure ssified based on available 	able	information.	
		- repeated exposure ssified based on availa	able	information.	
	Repea	ted dose toxicity			
	Compo	onents:			
	metfor	min hydrochloride:			

according to Regulation (EC) No. 1907/2006, as amended by Commission Regulation (EU) 2020/878



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	EL cation Route sure time	: Rat : 125 mg/kg : Oral : 1 year : No significant :	adverse effects were reported
	EL cation Route sure time	: Rabbit : 100 mg/kg : Oral : 1 Year : No significant :	adverse effects were reported
	EL cation Route sure time	: Dog : 50 mg/kg : Subcutaneous : 2 year : No significant :	adverse effects were reported
Sitag	liptin:		
Expo	EL	: Mouse : 500 mg/kg : 1.000 mg/kg : Oral : > 2 yr : Kidney	
Expo	EL	: Rat : 500 mg/kg : 1.000 mg/kg : Oral : 14 Weeks : Liver, Kidney,	Heart, Teeth
Expo Targe	EL EL cation Route sure time et Organs otoms	 Dog 10 mg/kg 50 mg/kg Oral 53 Weeks Central nervou Loss of balance The mechanise mans. 	
Expo Targe	EL EL cation Route sure time et Organs otoms	: Loss of balance	e, Central nervous system e m or mode of action may not be relevant in hu-



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		mans.	
Specie NOAE		: Monkey : 100 mg/kg	
	ation Route	: Oral	
	sure time	: 14 Weeks	
Rema	rks	: No significant	adverse effects were reported

Aspiration toxicity

Not classified based on available information.

11.2 Information on other hazards

Endocrine disrupting properties

Product:

Assessment	:	The substance/mixture does not contain components consid- ered 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

Components:

metformin hydrochloride:		
Skin contact	:	Remarks: May irritate skin.
Eye contact	:	Remarks: May irritate eyes.
Ingestion	:	Symptoms: Diarrhoea, Nausea, Vomiting, Gastrointestinal discomfort, flatulence, asthenia, Fatigue, Headache
Sitagliptin:		
Inhalation	:	Symptoms: upper respiratory tract infection, pharyngitis, Headache
Ingestion	:	Symptoms: upper respiratory tract infection, nasopharyngitis, Headache, Nausea, Abdominal pain, Diarrhoea

SECTION 12: Ecological information

12.1 Toxicity

Components:

metformin hydrochloride:

Toxicity to algae/aquatic plants	EC50 (Pseudokirchneriella subcapitata (green algae)): > 100 mg/l
	Exposure time: 72 h Method: OECD Test Guideline 201

NOEC (Pseudokirchneriella subcapitata (green algae)): 100



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				mg/l Exposure time: 72 Method: OECD Te	
Τ	oxicity	to microorganisms	:	EC50 : > 1.000 m Exposure time: 3 Test Type: Respir Method: OECD Te	h ation inhibition
	oxicity ity)	to fish (Chronic tox-	:	NOEC: 10 mg/l Exposure time: 33 Species: Pimepha Method: OECD Te	ales promelas (fathead minnow)
a		to daphnia and other invertebrates (Chron- ty)	:	NOEC: 40 mg/l Exposure time: 21 Species: Daphnia Method: OECD Te	magna (Water flea)
S	itaglip	otin:			
		to fish	:	LC50 (Pimephales Exposure time: 96 Method: OECD Te	
		to daphnia and other invertebrates	:	EC50 (Daphnia m Exposure time: 48 Method: OECD Te	
	oxicity lants	to algae/aquatic	:	EC50 (Pseudokiro mg/l Exposure time: 96 Method: OECD Te	
				NOEC (Pseudokir mg/l Exposure time: 96 Method: OECD Te	
T	oxicity	to microorganisms	:	EC50 : > 150 mg/ Exposure time: 3 Test Type: Respir Method: OECD Te	h ation inhibition
				NOEC : 150 mg/l Exposure time: 3 Test Type: Respir	
	oxicity ity)	to fish (Chronic tox-	:	NOEC: 9,2 mg/l Exposure time: 33 Species: Pimepha Method: OECD Te	ales promelas (fathead minnow)



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		ty to daphnia and other c invertebrates (Chron- city)	:		1 d a magna (Water flea) est Guideline 211
12.	2 Persis	stence and degradabil	ity		
	<u>Comp</u>	onents:			
		rmin hydrochloride: gradability	:	Result: rapidly de Biodegradation: Exposure time: 2	50 %
	Sitagl	iptin:			
	Biode	gradability	:	Result: not rapidly Biodegradation: Exposure time: 24 Method: OECD T	39,7 %
	Stabili	ty in water	:	pH: 7 Hydrolysis: 50 % Method: OECD T	(401 d) est Guideline 111
12.3 Bioaccumulative potential					
	<u>Comp</u>	onents:			
	Partitic	rmin hydrochloride: on coefficient: n- ol/water	:	log Pow: -2	
	Sitagl	iptin:			
		on coefficient: n- bl/water	:	log Pow: -0,03	
12.	4 Mobil	ity in soil			
	<u>Comp</u>	onents:			
	metfo	rmin hydrochloride:			
		ution among environ- I compartments	:		est Guideline 106
		iptin: ution among environ- I compartments	:	log Koc: 4,37	



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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

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

No data available

SECTION 13: Disposal considerations

13.1 Waste treatment methods

Product	:	Dispose of in accordance with local regulations.
		According to the European Waste Catalogue, Waste Codes
		are not product specific, but application specific.
		Waste codes should be assigned by the user, preferably in
		discussion with the waste disposal authorities.
		Do not dispose of waste into sewer.
Contaminated packaging	:	Empty containers should be taken to an approved waste han-
		dling site for recycling or disposal.
		If not otherwise specified: Dispose of as unused product.

SECTION 14: Transport information

14.1 UN number or ID number ADN : Not regulated as a dangerous good ADR : Not regulated as a dangerous good RID Not regulated as a dangerous good : IMDG : Not regulated as a dangerous good ΙΑΤΑ Not regulated as a dangerous good : 14.2 UN proper shipping name ADN Not regulated as a dangerous good :

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ADR		:	Not regulated as	a dangerous good		
RID		:	Not regulated as	a dangerous good		
IMDG		:	Not regulated as a dangerous good			
ΙΑΤΑ		:	Not regulated as a dangerous good			
14.3 Trans	sport hazard class(es))				
ADN		:	Not regulated as	a dangerous good		
ADR		:	Not regulated as	a dangerous good		
RID		:	Not regulated as	a dangerous good		
IMDG		:	Not regulated as	a dangerous good		
ΙΑΤΑ		:	Not regulated as	a dangerous good		
14.4 Packing group						
ADN		:	Not regulated as	a dangerous good		
ADR		:	Not regulated as	a dangerous good		
RID		:	Not regulated as	a dangerous good		
IMDG		:	Not regulated as	a dangerous good		
ΙΑΤΑ	(Cargo)	:	Not regulated as	a dangerous good		
ΙΑΤΑ	(Passenger)	:	Not regulated as	a dangerous good		
14.5 Envir	onmental hazards					
Not regulated as a dangerous good						
•	ial precautions for use oplicable	er				
14.7 Marit	ime transport in bulk	acco	ording to IMO inst	ruments		
Rema	rks	:	Not applicable for	r product as supplied.		

SECTION 15: Regulatory information

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

ons of restriction for the fol- entries should be considered: on list 75 tend to use this product as
on list 7

If you intend to use this product as tattoo ink, please contact your vendor.

Substance(s) or mixture(s) are listed here according to their appearance in the regulation, irrespective of their use/purpose or the conditions of the Commission Regulation (EU) 2020/878



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				restriction. Please refer to the condi- tions in corresponding Regulation to determine whether an entry is appli- cable to the placing on the market or not.
	H - Candidate List of S	ubstances of Very High rticle 59).	า	: Not applicable
	H - List of substances	subject to authorisation	1	: Not applicable
Regula	,	09 on substances that o	de-	: Not applicable
	ation (EU) 2019/1021 c	on persistent organic po	ollu-	: Not applicable
Regula ment a	ation (EC) No 649/2012	2 of the European Parli ning the export and imp		: Not applicable
Seves	o III: Directive 2012/18	/EU of the European P ving dangerous substa Not applicable		ent and of the Council on the control of

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

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

15.2 Chemical safety assessment

A Chemical Safety Assessment has not been carried out.

SECTION 16: Other information

Other information	:	Items where changes have been made to the previous version are highlighted in the body of this document by two vertical lines.

Full text of H-Statements

H302	:	Harmful if swallowed.
H319	:	Causes serious eye irritation.

Full text of other abbreviations

Acute Tox.	:	Acute toxicity
Eye Irrit.	:	Eye irritation
FOR-2011-12-06-1358	:	Norway. Occupational Exposure limits
FOR-2011-12-06-1358 /	:	Long term exposure limit
TWA		0

ADN - European Agreement concerning the International Carriage of Dangerous Goods by Inland Waterways; ADR - Agreement concerning the International Carriage of Dangerous Goods by Road; AIIC - Australian Inventory of Industrial Chemicals; ASTM - American Society for the Test-



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ing of Materials; bw - Body weight; CLP - Classification Labelling Packaging Regulation; Regulation (EC) No 1272/2008; CMR - Carcinogen, Mutagen or Reproductive Toxicant; DIN - Standard of the German Institute for Standardisation; DSL - Domestic Substances List (Canada); ECHA -European Chemicals Agency; EC-Number - European Community number; ECx - Concentration associated with x% response; ELx - Loading rate associated with x% response; EmS - Emergency Schedule; ENCS - Existing and New Chemical Substances (Japan); ErCx - Concentration associated with x% growth rate response; GHS - Globally Harmonized System; GLP - Good Laboratory Practice; IARC - International Agency for Research on Cancer; IATA - International Air Transport Association; IBC - International Code for the Construction and Equipment of Ships carrying Dangerous Chemicals in Bulk; IC50 - Half maximal inhibitory concentration; ICAO - International Civil Aviation Organization; IECSC - Inventory of Existing Chemical Substances in China; IMDG - International Maritime Dangerous Goods; IMO - International Maritime Organization; ISHL - Industrial Safety and Health Law (Japan); ISO - International Organisation for Standardization; KECI - Korea Existing Chemicals Inventory; LC50 - Lethal Concentration to 50 % of a test population; LD50 - Lethal Dose to 50% of a test population (Median Lethal Dose); MARPOL - International Convention for the Prevention of Pollution from Ships; n.o.s. - Not Otherwise Specified; NO(A)EC - No Observed (Adverse) Effect Concentration; NO(A)EL - No Observed (Adverse) Effect Level; NOELR - No Observable Effect Loading Rate; NZIoC - New Zealand Inventory of Chemicals; OECD - Organization for Economic Co-operation and Development; OPPTS - Office of Chemical Safety and Pollution Prevention; PBT - Persistent, Bioaccumulative and Toxic substance; PICCS - Philippines Inventory of Chemicals and Chemical Substances; (Q)SAR - (Quantitative) Structure Activity Relationship; REACH - Regulation (EC) No 1907/2006 of the European Parliament and of the Council concerning the Registration, Evaluation, Authorisation and Restriction of Chemicals; RID - Regulations concerning the International Carriage of Dangerous Goods by Rail; SADT - Self-Accelerating Decomposition Temperature; SDS - Safety Data Sheet; SVHC - Substance of very high concern; TCSI - Taiwan Chemical Substance Inventory; TECI -Thailand Existing Chemicals Inventory; TSCA - Toxic Substances Control Act (United States); UN - United Nations; UNRTDG - United Nations Recommendations on the Transport of Dangerous Goods; vPvB - Very Persistent and Very Bioaccumulative

Further information

Sources of key data used to :	Internal technical data, data from raw material SDSs, OECD
compile the Safety Data	eChem Portal search results and European Chemicals Agen-
Sheet	cy, http://echa.europa.eu/

Classification of the mixture:

Classification procedure:

Acute Tox. 4

H302

Calculation method

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



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