SAFETY DATA SHEET

Based upon Regulation (EC) No 1907/2006, as amended by Regulation (EU) No 2020/878



SEAL & BOND SIL 25

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

1.1. Product identifier

: SEAL & BOND SIL 25 Product name **Registration number REACH** Product type REACH : Mixture

: Not applicable (mixture)

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

1.2.1 Relevant identified uses Sealing compound

1.2.2 Uses advised against

No uses advised against known

1.3. Details of the supplier of the safety data sheet

Supplier of the safety data sheet

Novatio* Industrielaan 5B B-2250 Olen +32 14 25 76 40 **▲** +32 14 22 02 66 info@novatio.be *NOVATIO is a registered trademark of Novatech International N.V.

Manufacturer of the product

Novatech International N.V. Industrielaan 5B B-2250 Olen +32 14 85 97 37 **i ⊟** +32 14 85 97 38 info@novatech.be

1.4. Emergency telephone number

24h/24h (Telephone advice: English, French, German, Dutch) : +32 14 58 45 45 (BIG)

SECTION 2: Hazards identification

2.1. Classification of the substance or mixture

Not classified as dangerous according to the criteria of Regulation (EC) No 1272/2008

2.2. Label elements

Not classified as dangerous according to the criteria of Regulation (EC) No 1272/2008

Supplemental information EU

H208	Contains: butan-2-one O,O',O''-(methylsilylidyne)trioxime; 2-butanone oxime. May produce an allergic reaction.
H211	Warning! Hazardous respirable droplets may be formed when sprayed. Do not breathe spray or mist.
H210	Safety data sheet available on request.

2.3. Other hazards

EU EU

No other hazards known

SECTION 3: Composition/information on ingredients

3.1. Substances

Not applicable

3.2. Mixtures

Name REACH Registration No	CAS No EC No	Conc. (C)	Classification according to CLP	Note	lRemark	M-factors and ATE
Created by: Brandweerinformatiecentrum voo Technische Schoolstraat 43 A, B-2440 Geel http://www.big.be © BIG vzw	r gevaarlijke stoffer	n vzw (BIG)		tion date: 2001-(revision: 2021-0		16239-019-en
Reason for revision: 15 Revision number: 0401			BIG nun	nber: 35139		& % 1 / 15

	S	EAL & I	BOND SIL 2	.5		
titanium dioxide; [in powder form containing 1 % or more of particles with aerodynamic diameter ≤ 10 μm] 01-2119489379-17	13463-67-7 236-675-5	C≥1%	Carc. 2; H351	(1)(2)	Constituent	
butan-2-one 0,0',0''-(methylsilylidyne)trioxime	22984-54-9 245-366-4	0.1% <c<1%< td=""><td>Skin Sens. 1; H317 Skin Irrit. 2; H315 Eye Irrit. 2; H319</td><td>(1)(10)</td><td>Constituent</td><td></td></c<1%<>	Skin Sens. 1; H317 Skin Irrit. 2; H315 Eye Irrit. 2; H319	(1)(10)	Constituent	
2-butanone oxime	96-29-7 202-496-6	0.1% <c<1%< td=""><td>Carc. 2; H351 Skin Sens. 1; H317 Acute Tox. 4; H312 Eye Dam. 1; H318</td><td>(1)(10)</td><td>Constituent</td><td>ATE oral: 100 mg/kg ATE dermal: 1100 mg/kg</td></c<1%<>	Carc. 2; H351 Skin Sens. 1; H317 Acute Tox. 4; H312 Eye Dam. 1; H318	(1)(10)	Constituent	ATE oral: 100 mg/kg ATE dermal: 1100 mg/kg
SECTION 4: First aid measur 4.1. Description of first aid measu						
General: Observe (own) safety. If possible number 112. Treat symptoms st symptoms.	, approach vic			• •		
After inhalation: Remove victim into fresh air. In o	case of respira	tory problems,	consult a doctor/medica	al service.		
After skin contact: If possible, wipe up/dry remove doctor/medical service.	chemical. The	n rinse/shower	immediately with (lukew	varm) water. If irri	itation persists, con	sult a
After eye contact: Rinse immediately with (lukewa doctor/medical service.	rm) water. Ren	nove contact le	nses, if present and eas	y to do. Continue	rinsing. If irritation	persists, consult a
After ingestion: Rinse mouth with water. If you f	eel unwell, cor	nsult a doctor/r	nedical service. Do not v	vait for symptom	s to occur to consul	t Poison Center.
4.2. Most important symptoms ar	nd effects, bo	th acute and	delayed			
4.2.1 Acute symptoms After inhalation: No effects known.						
After skin contact: No effects known.						
After eye contact: No effects known.						

No effects known. After ingestion: No effects known.

4.2.2 Delayed symptoms

No effects known.

4.3. Indication of any immediate medical attention and special treatment needed

If applicable and available it will be listed below.

SECTION 5: Firefighting measures

5.1. Extinguishing media

5.1.1 Suitable extinguishing media:

Small fire: Quick-acting ABC powder extinguisher, Quick-acting BC powder extinguisher, Quick-acting class B foam extinguisher, Quick-acting CO2 extinguisher. Major fire: Class B foam (not alcohol-resistant).

5.1.2 Unsuitable extinguishing media:

Small fire: Water (quick-acting extinguisher, reel); risk of puddle expansion. Major fire: Water; risk of puddle expansion.

5.2. Special hazards arising from the substance or mixture

Upon combustion: formation of CO, CO2 and small quantities of nitrous vapours.

5.3. Advice for firefighters

5.3.1 Instructions:

No specific fire-fighting instructions required.

5.3.2 Special protective equipment for fire-fighters:

Gloves (EN 374). Protective clothing (EN 14605 or EN 13034). Heat/fire exposure: self-contained breathing apparatus (EN 136 + EN 137).

Reason for revision: 15

SECTION 6: Accidental release measures

6.1. Personal precautions, protective equipment and emergency procedures

No naked flames.

- 6.1.1 Protective equipment for non-emergency personnel
- See section 8.2
- 6.1.2 Protective equipment for emergency responders Gloves (EN 374). Protective clothing (EN 14605 or EN 13034).

Suitable protective clothing

See section 8.2

6.2. Environmental precautions

Contain released product.

6.3. Methods and material for containment and cleaning up

Solid spill: cover with absorbent material. Scoop solid spill into closing containers. Clean contaminated surfaces with an excess of water. Wash clothing and equipment after handling.

6.4. Reference to other sections

See section 13.

SECTION 7: Handling and storage

The information in this section is a general description. If applicable and available, exposure scenarios are attached in annex. Always use the relevant exposure scenarios that correspond to your identified use.

7.1. Precautions for safe handling

Keep away from naked flames/heat. Observe strict hygiene. Keep container tightly closed.

7.2. Conditions for safe storage, including any incompatibilities

7.2.1 Safe storage requirements:

Meet the legal requirements. Store in a cool area. Store in a dry area. Keep container in a well-ventilated place.

7.2.2 Keep away from:

Heat sources, water/moisture.

7.2.3 Suitable packaging material: Synthetic material. metal.

7.2.4 Non suitable packaging material:

No data available

7.3. Specific end use(s)

If applicable and available, exposure scenarios are attached in annex. See information supplied by the manufacturer.

SECTION 8: Exposure controls/personal protection

8.1. Control parameters

8.1.1 Occupational exposure

a) Occupational exposure limit values If limit values are applicable and available these will be listed below.

Belgium Titane (dioxyde de)

Titane (dioxyde de)	Time-weighted average exposure limit 8 h	10 mg/m ³
France		
Titane (dioxyde de), en Ti	Time-weighted average exposure limit 8 h (VL: Valeur non réglementaire indicative)	10 mg/m³
Germany		
Butanonoxim	Time-weighted average exposure limit 8 h (TRGS 900)	0.3 ppm
	Time-weighted average exposure limit 8 h (TRGS 900)	1 mg/m³
υκ		
Titanium dioxide respirable	Time-weighted average exposure limit 8 h (Workplace exposure limit (EH40/2005))	4 mg/m³
Titanium dioxide total inhalable	Time-weighted average exposure limit 8 h (Workplace exposure limit (EH40/2005))	10 mg/m ³
USA (TLV-ACGIH)		-
Titanium dioxide	Time-weighted average exposure limit 8 h (TLV - Adopted Value)	10 mg/m ³
 <u>b) National biological limit values</u> If limit values are applicable and available these v USA (BEI-ACGIH) 	vill be listed below.	
or revision: 15	Publication date: 2001-03-15	
	Date of revision: 2021-04-27	
		2

1.2 Sampling methods Product name TiO2 TiO2 1.3 Applicable limit values when using If limit values are applicable and a 1.4 Threshold values DNEL/DMEL - Workers butan-2-one O,O',O''-(methylsilylidyne Effect level (DNEL/DMEL) DNEL 2-butanone oxime Effect level (DNEL/DMEL) DNEL DNEL	available these e <u>)trioxime</u> Type Long-term syste		Number 7302 7304		
TiO2 TiO2 TiO2 1.3 Applicable limit values when using If limit values are applicable and a 1.4 Threshold values <u>DNEL/DMEL - Workers</u> <u>butan-2-one O,O',O''-(methylsilylidyne</u> <u>Effect level (DNEL/DMEL)</u> <u>DNEL</u> <u>2-butanone oxime</u> <u>Effect level (DNEL/DMEL)</u>	available these e <u>)trioxime</u> Type Long-term syste	NIOSH NIOSH r mixture as intended will be listed below.	7302 7304		
TiO2 I.3 Applicable limit values when using If limit values are applicable and a I.4 Threshold values DNEL/DMEL - Workers butan-2-one O,O',O''-(methylsilylidyne Effect level (DNEL/DMEL) DNEL 2-butanone oxime Effect level (DNEL/DMEL)	available these e <u>)trioxime</u> Type Long-term syste	NIOSH r mixture as intended will be listed below.	7304		
1.3 Applicable limit values when using If limit values are applicable and a 1.4 Threshold values <u>DNEL/DMEL - Workers butan-2-one O,O',O''-(methylsilylidynee Effect level (DNEL/DMEL) DNEL 2-butanone oxime Effect level (DNEL/DMEL) </u>	available these e <u>)trioxime</u> Type Long-term syste	r mixture as intended will be listed below.			
If limit values are applicable and a 1.4 Threshold values DNEL/DMEL - Workers butan-2-one O,O',O''-(methylsilylidyne Effect level (DNEL/DMEL) DNEL 2-butanone oxime Effect level (DNEL/DMEL)	available these e <u>)trioxime</u> Type Long-term syste	will be listed below.	L		
Effect level (DNEL/DMEL) DNEL 2-butanone oxime Effect level (DNEL/DMEL)	Type Long-term syste	mic effects inhalation			
DNEL <u>2-butanone oxime</u> Effect level (DNEL/DMEL)	Long-term syste	mic effects inhalation	Value		Remark
2-butanone oxime Effect level (DNEL/DMEL)			1.02 mg/m ³	3	
Effect level (DNEL/DMEL)	- 0	mic effects dermal	0.145 mg/k		
			0,	0 , . ,	
DNEL	Туре		Value		Remark
	Long-term syste	mic effects inhalation	28 μg/m³		
	Long-term local	effects inhalation	0.9 mg/m ³		
	Long-term syste	mic effects dermal	4 μg/kg bw	/day	
DNEL/DMEL - General population	\				
butan-2-one O,O',O''-(methylsilylidyne Effect level (DNEL/DMEL)			halis		Demont
	Туре	unin offente in balation	Value 0.25 mg/m ²	3	Remark
DNEL		mic effects inhalation	0.25 mg/m 0.072 mg/k		
	· · ·	mic effects oral 0.072 mg/kg			
2-butanone oxime	Long-term syste		0.072 118/ K	g Dw/uay	
Effect level (DNEL/DMEL)	Туре		Value		Remark
DNEL	<i></i>	mic effects inhalation	4.82 μg/m ³		
		effects inhalation	0.43 mg/m		
	Long-term syste	mic effects oral	1.6 μg/kg b	w/day	
PNEC butan-2-one O,O',O''-(methylsilylidyne	<u>)trioxime</u>				
Compartments		Value		Remark	
Fresh water		0.018 mg/l			
Marine water		0.002 mg/l			
STP		3.9 mg/l			
Fresh water sediment		557.543 mg/kg sediment dw			
Marine water sediment		55.754 mg/kg sediment dw			
Soil		65.63 mg/kg soil dw			
Oral		3.22 mg/kg food			
2-butanone oxime		Value		Remark	
Compartments		0.256 mg/l		Remark	
Fresh water Marine water		0.026 mg/l		+	
Fresh water (intermittent releases)		0.118 mg/l			
Marine water (intermittent releases))	0.012 mg/l			
STP	1	177 mg/l			
Fresh water sediment		1.012 mg/kg sediment dw		+	
Marine water sediment		0.101 mg/kg sediment dw			
Soil		0.052 mg/kg soil dw			
1.5 Control banding				1	

relevant exposure scenarios that correspond to your identified use. 8.2.1 Appropriate engineering controls

Keep away from naked flames/heat. Carry operations in the open/under local exhaust/ventilation or with respiratory protection.

8.2.2 Individual protection measures, such as personal protective equipment

Observe strict hygiene. Do not eat, drink or smoke during work.

a) Respiratory protection:

Respiratory protection not required in normal conditions. Mist formation: aerosol mask with filter type P3.

b) Hand protection:

Materials	Measured	Thickness	Protection index	Remark
	breakthrough time			
butyl rubber		> 0.7 mm		Good resistance
viton		> 0.7 mm		Good resistance

Safety glasses (EN 166).

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d) Skin protection:

Protective clothing (EN 14605 or EN 13034).

8.2.3 Environmental exposure controls:

See sections 6.2, 6.3 and 13

SECTION 9: Physical and chemical properties

9.1. Information on basic physical and chemical properties

Physical form	Paste
Viscosity	Viscous
Odour	Characteristic odour
Odour threshold	No data available in the literature
Colour	Variable in colour, depending on the composition
Particle size	Not applicable (liquid)
Explosion limits	No data available in the literature
Flammability	Not classified as flammable
Log Kow	Not applicable (mixture)
Dynamic viscosity	No data available in the literature
Kinematic viscosity	No data available in the literature
Melting point	No data available in the literature
Boiling point	No data available in the literature
Relative vapour density	Not applicable
Vapour pressure	No data available in the literature
Solubility	Water ; insoluble
Relative density	1.4 ; 20 °C
Absolute density	1400 kg/m³ ; 20 °C
Decomposition temperature	No data available in the literature
Auto-ignition temperature	No data available in the literature
Flash point	No data available in the literature
рН	Not applicable (non-soluble in water)

9.2. Other information

No data available

SECTION 10: Stability and reactivity

10.1. Reactivity

Heating increases the fire hazard.

10.2. Chemical stability

Stable under normal conditions.

10.3. Possibility of hazardous reactions

No data available.

10.4. Conditions to avoid

Precautionary measures

Keep away from naked flames/heat.

10.5. Incompatible materials

Water/moisture.

10.6. Hazardous decomposition products

Upon combustion: formation of CO, CO2 and small quantities of nitrous vapours.

SECTION 11: Toxicological information

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

11.1.1 Test results

Acute toxicity

SEAL & BOND SIL 25

No (test)data on the mixture available Judgement is based on the relevant ingredients

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Revision number: 0401

Route of exposure	Parameter	Method	Value	Exposure time	Species	Value determination	Remark
Oral	LD50	OECD 401	> 2000 mg/kg bw		Rat (male / female)	Experimental value	
Dermal						Data waiving	
Inhalation (dust)	LC50	OECD 403	> 5.09 mg/l	4 h	Rat (male)	Experimental value	
an-2-one 0,0',0''-(me	thylsilylidyn	e)trioxime				•	
Route of exposure	Parameter	Method	Value	Exposure time	Species	Value determination	Remark
Oral	LD50	OECD 401	2463 mg/kg bw		Rat (male / female)	Experimental value	
Dermal	LD50	OECD 402	> 2000 mg/kg bw	24 h	Rat (male / female)	Experimental value	
Inhalation						Data waiving	
utanone oxime				-			
Route of exposure	Parameter	Method	Value	Exposure time	Species	Value determination	Remark
Oral	LD50	Equivalent to OECD 401	2326 mg/kg bw		Rat (male)	Experimental value	
Dermal	LD50	Equivalent to OECD 402	> 1000 mg/kg bw	24 h	Rabbit (male / female)	Experimental value	
Inhalation (vapours)	LC50	Equivalent to OECD 403	> 4.83 mg/l air	4 h	Rat (male / female)	Experimental value	

Not classified for acute toxicity

Corrosion/irritation

SEAL & BOND SIL 25

No (test)data on the mixture available

Judgement is based on the relevant ingredients <u>titanium dioxide;</u> [in powder form containing 1 % or more of particles with aerodynamic diameter $\le 10 \mu$ m]

Route of exposure	Result	Method	Exposure time	Time point	Species	Value determination	Remark
Eye	Not irritating	OECD 405		1; 24; 48; 72 hours	Rabbit	Experimental value	
Skin	Not irritating	Equivalent to OECD 404	4 h	48 hours	Rabbit	Experimental value	
tan-2-one 0,0',0''-(methylsilylidyne)	<u>trioxime</u>					
Route of exposure	Result	Method	Exposure time	Time point	Species	Value determination	Remark
Еуе	Irritating	Equivalent to OECD 405		24; 48; 72 hours	Rabbit	Experimental value	
Skin	Irritating; category 2					Literature study	
outanone oxime							
Route of exposure	Result	Method	Exposure time	Time point	Species	Value	Remark

Route of exposure	Result	Method	Exposure time	Time point	Species	Value	Remark
						determination	
Eye	Serious eye	Equivalent to			Rabbit	Experimental	Single treatment
	damage	OECD 405				value	
Skin	Irritating	Equivalent to EPA	4 h	1; 24; 48; 72 hours	Rabbit	Experimental	
		OPPTS 870.2500				value	

Classification of this substance according to Annex VI is debatable as it does not correspond to the conclusion from the test

Conclusion

Not classified as irritating to the skin

Not classified as irritating to the eyes

Not classified as irritating to the respiratory system

Respiratory or skin sensitisation

SEAL & BOND SIL 25

No (test)data on the mixture available

Judgement is based on the relevant ingredients titanium dioxide; [in powder form containing 1 % or more of particles with aerodynamic diameter \leq 10 μ m]

Route of exposure	Result	Method	Exposure time	Observation time	Species	Value determination	Remark
				point			
Skin	Not sensitizing	Equivalent to OECD 429			Mouse (female)	Experimental value	
Inhalation (dust)	Not sensitizing				Mouse (female)	Experimental value	

Reason for revision: 15

Route of exposure	Result	Method	Expo	sure time	Observation time	Species	Value determination	n Remark
					point			
Skin	Sensitizing	Equivalent 406	to OECD			Guinea pig (female)	Experimental value	
2-butanone oxime								
Route of exposure	Result	Method	Ехро	sure time	Observation time point	Species	Value determination	n Remark
Skin	Sensitizing	Equivalent 406	to OECD			Guinea pig (female)	Experimental value	
onclusion						•		
Not classified as sensi	-	ation						
Not classified as sensi	tizing for skin							
ic target organ toxici	ty							
L & BOND SIL 25								
o (test)data on the mi	xture availabl	2						
udgement is based o	n the relevant	ingredients						
itanium dioxide; [in p		-					-	
Route of exposure	e Parameter	Method	Value	Organ	Effect	Exposure time	Species	Value determinatio
Oral (stomach	NOAEL	OECD 408	> 1000 mg/kg		No effect	90 day(s)	Rat (male /	Experimenta
tube)			bw/day				female)	value
Dermal								Data waiving
outan-2-one 0,0',0"-								
Route of exposure	e Parameter	Method	Value	Organ	Effect	Exposure time	Species	Value determinatio
Oral (stomach	NOEL	Subacute	4 mg/kg	Blood	No effect	28 day(s)	Rat (male /	Experimenta
tube)	NOLL	toxicity test	bw/day	biood	No chect	20 44 (0)	female)	value
Oral (drinking	NOAEL	Equivalent to	25 mg/kg	Blood	No effect	13 week(s)	Rat (male /	Experimenta
water)		OECD 408	bw/day - 30				female)	value
			mg/kg bw/day					
Classification of the	his substance i	s debatable as it	does not corre	spond to the co	nclusion from the	test		
-butanone oxime								
-butanone oxime Route of exposure	e Parameter	Method	Value	Organ	Effect	Exposure time	Species	Value
	e Parameter	Method	Value	Organ	Effect	Exposure time	Species	
Route of exposure Oral (drinking	Parameter	EPA OPPTS	25 mg/kg	Organ Blood	Effect No effect	Exposure time 13 week(s)	Rat (male /	determination Experimenta
Route of exposure			25 mg/kg bw/day - 30	Blood				determinatio
Route of exposure Oral (drinking water)	NOAEL	EPA OPPTS 870.3100	25 mg/kg bw/day - 30 mg/kg bw/day	Blood	No effect	13 week(s)	Rat (male / female)	determination Experimentation value
Route of exposure Oral (drinking		EPA OPPTS	25 mg/kg bw/day - 30	Blood			Rat (male / female)	determination Experimentation value
Route of exposure Oral (drinking water) Oral (stomach	NOAEL	EPA OPPTS 870.3100 EPA OTS	25 mg/kg bw/day - 30 mg/kg bw/day 125 mg/kg	Blood / Central	No effect No effect	13 week(s) 13 weeks (5 days / week)	Rat (male / female) / Rat (male / female)	determination Experimentation value Experimentation
Route of exposure Oral (drinking water) Oral (stomach tube) Oral (stomach	NOAEL	EPA OPPTS 870.3100 EPA OTS 798.6050 EPA OTS	25 mg/kg bw/day - 30 mg/kg bw/day 125 mg/kg bw/day 40 mg/kg	Blood / Central nervous	No effect No effect Change in the	13 week(s) 13 weeks (5 days / week) 13 weeks (5 days /	Rat (male / female) / Rat (male / female) / Rat (male /	determination Experimentaria value Experimentaria value Experimentaria value
Route of exposure Oral (drinking water) Oral (stomach tube)	NOAEL NOAEL	EPA OPPTS 870.3100 EPA OTS 798.6050	25 mg/kg bw/day - 30 mg/kg bw/day 125 mg/kg bw/day	Blood Central nervous system	No effect No effect Change in the haemogramm	13 week(s) 13 weeks (5 days / week) 13 weeks (5 days /	Rat (male / female) / Rat (male / female)	determination Experimenta value Experimenta value
Route of exposure Oral (drinking water) Oral (stomach tube) Oral (stomach	NOAEL NOAEL	EPA OPPTS 870.3100 EPA OTS 798.6050 EPA OTS	25 mg/kg bw/day - 30 mg/kg bw/day 125 mg/kg bw/day 40 mg/kg	Blood Central nervous system	No effect No effect Change in the	13 week(s) 13 weeks (5 days / week) 13 weeks (5 days /	Rat (male / female) / Rat (male / female) / Rat (male /	determination Experimentaria value Experimentaria value Experimentaria value
Route of exposure Oral (drinking water) Oral (stomach tube) Oral (stomach	NOAEL NOAEL	EPA OPPTS 870.3100 EPA OTS 798.6050 EPA OTS	25 mg/kg bw/day - 30 mg/kg bw/day 125 mg/kg bw/day 40 mg/kg	Blood Central nervous system	No effect No effect Change in the haemogramm e/blood	13 week(s) 13 weeks (5 days / week) 13 weeks (5 days /	Rat (male / female) / Rat (male / female) / Rat (male /	determination Experimentation Value Experimentation value Experimentation value
Route of exposure Oral (drinking water) Oral (stomach tube) Oral (stomach tube) Dermal Inhalation	NOAEL NOAEL	EPA OPPTS 870.3100 EPA OTS 798.6050 EPA OTS 798.6050 Equivalent to	25 mg/kg bw/day - 30 mg/kg bw/day 125 mg/kg bw/day 40 mg/kg	Blood Central nervous system	No effect No effect Change in the haemogramm e/blood	13 week(s) 13 weeks (5 days / week) 13 weeks (5 days / week) 4 weeks (6h / day,	Rat (male / female) / Rat (male / female) / Rat (male / female) 5 Rat (male /	determination Experimentary Value Experimentary Value Experimentary Value Data waiving Experimentary
Route of exposure Oral (drinking water) Oral (stomach tube) Oral (stomach tube) Dermal Inhalation (vapours)	NOAEL NOAEL LOAEL	EPA OPPTS 870.3100 EPA OTS 798.6050 EPA OTS 798.6050	25 mg/kg bw/day - 30 mg/kg bw/day 125 mg/kg bw/day 40 mg/kg bw/day	Blood Central nervous system Blood	No effect No effect Change in the haemogramm e/blood composition	13 week(s) 13 weeks (5 days / week) 13 weeks (5 days / week)	Rat (male / female) / Rat (male / female) / Rat (male / female)	determination Experimentation Value Experimentation Value Experimentation Value Data waiving
Route of exposure Oral (drinking water) Oral (stomach tube) Oral (stomach tube) Dermal Inhalation (vapours) nclusion	NOAEL NOAEL LOAEL NOAEL NOAEL	EPA OPPTS 870.3100 EPA OTS 798.6050 EPA OTS 798.6050 Equivalent to OECD 412	25 mg/kg bw/day - 30 mg/kg bw/day 125 mg/kg bw/day 40 mg/kg bw/day	Blood Central nervous system Blood	No effect No effect Change in the haemogramm e/blood composition	13 week(s) 13 weeks (5 days / week) 13 weeks (5 days / week) 4 weeks (6h / day,	Rat (male / female) / Rat (male / female) / Rat (male / female) 5 Rat (male /	determination Experimentation Value Experimentation Value Experimentation Data waiving Experimentation
Route of exposure Oral (drinking water) Oral (stomach tube) Oral (stomach tube) Dermal Inhalation (vapours) nclusion Not classified for subo	NOAEL NOAEL LOAEL NOAEL NOAEL	EPA OPPTS 870.3100 EPA OTS 798.6050 EPA OTS 798.6050 Equivalent to OECD 412	25 mg/kg bw/day - 30 mg/kg bw/day 125 mg/kg bw/day 40 mg/kg bw/day	Blood Central nervous system Blood	No effect No effect Change in the haemogramm e/blood composition	13 week(s) 13 weeks (5 days / week) 13 weeks (5 days / week) 4 weeks (6h / day,	Rat (male / female) / Rat (male / female) / Rat (male / female) 5 Rat (male /	determination Experimentary Value Experimentary Value Experimentary Value Data waiving Experimentary
Oral (drinking water) Oral (stomach tube) Oral (stomach tube) Dermal Inhalation	NOAEL NOAEL LOAEL NOAEL NOAEL	EPA OPPTS 870.3100 EPA OTS 798.6050 EPA OTS 798.6050 Equivalent to OECD 412	25 mg/kg bw/day - 30 mg/kg bw/day 125 mg/kg bw/day 40 mg/kg bw/day	Blood Central nervous system Blood	No effect No effect Change in the haemogramm e/blood composition	13 week(s) 13 weeks (5 days / week) 13 weeks (5 days / week) 4 weeks (6h / day,	Rat (male / female) / Rat (male / female) / Rat (male / female) 5 Rat (male /	determination Experimentation Value Experimentation value Experimentation Value Data waiving Experimentation
Route of exposure Oral (drinking water) Oral (stomach tube) Oral (stomach tube) Dermal Inhalation (vapours) nclusion Not classified for subo	NOAEL NOAEL LOAEL NOAEL NOAEL	EPA OPPTS 870.3100 EPA OTS 798.6050 EPA OTS 798.6050 Equivalent to OECD 412	25 mg/kg bw/day - 30 mg/kg bw/day 125 mg/kg bw/day 40 mg/kg bw/day	Blood Central nervous system Blood	No effect No effect Change in the haemogramm e/blood composition	13 week(s) 13 weeks (5 days / week) 13 weeks (5 days / week) 4 weeks (6h / day,	Rat (male / female) / Rat (male / female) / Rat (male / female) 5 Rat (male /	determination Experimentation Value Experimentation value Experimentation Value Data waiving Experimentation
Route of exposure Oral (drinking water) Oral (stomach tube) Oral (stomach tube) Dermal Inhalation (vapours) nclusion Not classified for subo genicity (in vitro)	NOAEL NOAEL LOAEL NOAEL NOAEL NOAEC	EPA OPPTS 870.3100 EPA OTS 798.6050 EPA OTS 798.6050 Equivalent to OECD 412	25 mg/kg bw/day - 30 mg/kg bw/day 125 mg/kg bw/day 40 mg/kg bw/day	Blood Central nervous system Blood	No effect No effect Change in the haemogramm e/blood composition	13 week(s) 13 weeks (5 days / week) 13 weeks (5 days / week) 4 weeks (6h / day,	Rat (male / female) / Rat (male / female) / Rat (male / female) 5 Rat (male /	determination Experimentation Value Experimentation value Experimentation Value Data waiving Experimentation
Route of exposure Oral (drinking water) Oral (stomach tube) Oral (stomach tube) Dermal Inhalation (vapours) nclusion Not classified for subo genicity (in vitro) L & BOND SIL 25 No (test)data on the is	NOAEL NOAEL LOAEL NOAEC NOAEC	EPA OPPTS 870.3100 EPA OTS 798.6050 EPA OTS 798.6050 Equivalent to OECD 412	25 mg/kg bw/day - 30 mg/kg bw/day 125 mg/kg bw/day 40 mg/kg bw/day 90 mg/m ³ air	Blood Central nervous system Blood Blood	No effect No effect Change in the haemogramm e/blood composition No effect	13 week(s) 13 weeks (5 days / week) 13 weeks (5 days / week) 4 weeks (6h / day, days / week)	Rat (male / female) / Rat (male / female) / Rat (male / female) 5 Rat (male /	determination Experimentation Value Experimentation value Experimentation Value Data waiving Experimentation
Route of exposure Oral (drinking water) Oral (stomach tube) Oral (stomach tube) Dermal Inhalation (vapours) nclusion Not classified for subo genicity (in vitro) L & BOND SIL 25 No (test)data on the in Judgement is based of itanium dioxide; [in p	NOAEL NOAEL NOAEL LOAEL NOAEC NOAEC	EPA OPPTS 870.3100 EPA OTS 798.6050 EPA OTS 798.6050 Equivalent to OECD 412	25 mg/kg bw/day - 30 mg/kg bw/day 125 mg/kg bw/day 40 mg/kg bw/day 90 mg/m ³ air	Es with aerody	No effect No effect Change in the haemogramm e/blood composition No effect	13 week(s) 13 weeks (5 days / week) 13 weeks (5 days / week) 4 weeks (6h / day, days / week) 10 μm]	Rat (male / female) Rat (male / female) Rat (male / female) S Rat (male / female) S Rat (male / female)	determination Experimentation value Experimentation value Experimentation value Data waiving Experimentation value
Route of exposure Oral (drinking water) Oral (stomach tube) Oral (stomach tube) Dermal Inhalation (vapours) nclusion Not classified for suborgenicity (in vitro) L & BOND SIL 25 No (test)data on the point of the suborgeneration of the sub	NOAEL NOAEL NOAEL LOAEL NOAEC NOAEC	EPA OPPTS 870.3100 EPA OTS 798.6050 EPA OTS 798.6050 Equivalent to OECD 412	25 mg/kg bw/day - 30 mg/kg bw/day 125 mg/kg bw/day 40 mg/kg bw/day 90 mg/m ³ air	Es with aerody	No effect No effect Change in the haemogramm e/blood composition No effect No effect Effect	13 week(s) 13 weeks (5 days / week) 13 weeks (5 days / week) 4 weeks (6h / day, days / week) <u>10 µm]</u> <u>Valu</u>	Rat (male / female) / Rat (male / female) / Rat (male / female) 5 Rat (male / female) 5 Rat (male / female) 9	determination Experimentation Value Experimentation value Experimentation Value Data waiving Experimentation
Route of exposure Oral (drinking water) Oral (stomach tube) Oral (stomach tube) Dermal Inhalation (vapours) nclusion Not classified for subo genicity (in vitro) L & BOND SIL 25 No (test)data on the in ludgement is based of itanium dioxide; [in p	NOAEL NOAEL NOAEL LOAEL NOAEC NOAEC	EPA OPPTS 870.3100 EPA OTS 798.6050 EPA OTS 798.6050 Equivalent to OECD 412	25 mg/kg bw/day - 30 mg/kg bw/day 125 mg/kg bw/day 40 mg/kg bw/day 90 mg/m ³ air	Es with aerody	No effect No effect Change in the haemogramm e/blood composition No effect No effect Effect	13 week(s) 13 weeks (5 days / week) 13 weeks (5 days / week) 4 weeks (6h / day, days / week) <u>10 µm]</u> <u>Valu</u>	Rat (male / female) Rat (male / female) Rat (male / female) S Rat (male / female) S Rat (male / female)	determination Experimentation value Experimentation value Experimentation value Data waiving Experimentation value

М

Result	Method	Test substrate	Effect	Value determination	Remark
Negative with metabolic activation, negative without metabolic activation	OECD 473	Chinese hamster ovary (CHO)		Experimental value	
Negative with metabolic activation, negative without metabolic activation	OECD 471	Bacteria (S.typhimurium)		Experimental value	

Reason for revision: 15

Publication date: 2001-03-15 Date of revision: 2021-04-27

Revision number: 0401

Result	Method	Test substrate	Effect	Value determination	Remark
Negative with metabolic activation, negative without metabolic activation	Equivalent to OECD 476	CHL/IU cells	No effect	Experimental value	
Positive with metabolic activation, positive without metabolic activation	Equivalent to OECD 476	Mouse (lymphoma L5178Y cells)	No effect	Read-across	
Negative with metabolic activation, negative without metabolic activation	Equivalent to OECD 473	Chinese hamster ovary (CHO)	No effect	Experimental value	
utanone oxime		•		•	
Result	Method	Test substrate	Effect	Value determination	Remark
Ambiguous	Equivalent to OECD 476	Mouse (lymphoma L5178Y cells)		Experimental value	
Negative with metabolic activation, negative without metabolic activation	OECD 471	Bacteria (S.typhimurium)		Experimental value	
Negative with metabolic activation, negative without metabolic activation	OECD 482	Rat liver cells		Experimental value	

Mutagenicity (in vivo)

SEAL & BOND SIL 25

No (test)data on the mixture available

Judgement is based on the relevant ingredients

<u>titanium dioxide;</u> [in powder form containing 1 % or more of particles with aerodynamic diameter \leq 10 μ m]

	Result	Method	Exposure time	Test substrate	Organ	Value determination
	Negative (Oral (stomach tube))	OECD 474		Mouse (male / female)		Experimental value
but	an-2-one 0,0',0"-(methylsilylidyne)tri	<u>oxime</u>				
	Result	Method	Exposure time	Test substrate	Organ	Value determination
	Negative (Oral (drinking water))	Equivalent to OECD 474	13 weeks (daily)	Mouse (male / female)		Experimental value
<u>2-b</u>	utanone oxime					
	Result	Method	Exposure time	Test substrate	Organ	Value determination
	Negative (Oral (diet))	Equivalent to OECD 477	3 day(s)	Drosophila melanogaster (male)		Experimental value
	Negative (Oral (stomach tube))	Equivalent to OECD 475		Rat (male / female)		Experimental value

Conclusion

Not classified for mutagenic or genotoxic toxicity

Carcinogenicity

SEAL & BOND SIL 25

No (test)data on the mixture available

Judgement is based on the relevant ingredients

The classification as a carcinogen by inhalation applies only to mixtures in powder form containing 1 % or more of titanium dioxide which is in the form of or incorporated in particles with aerodynamic diameter \leq 10 μ m.

titanium dioxide; [in powder form containing 1 % or more of particles with aerodynamic diameter \leq 10 μ m]

Route of exposure	Parameter	Method	Value	Exposure time	Species	Effect	Organ	Value determination
Inhalation (dust)	NOAEC	OECD 453	0,	104 weeks (6h / day, 5 days / week)	Rat (male / female)	No carcinogenic effect	Lungs	Experimental value
Oral (diet)		Carcinogenic toxicity study	50000 ppm	103 weeks (7 days / week)	Rat (male / female)	No carcinogenic effect		Experimental value

2-butanone oxime

Route of exposure	Parameter	Method	Value	Exposure time	Species	Effect	Organ	Value determination
Inhalation (vapours)	LOAEC	EPA OTS 798.3300	0,	26 weeks (6h / day, 5 days / week)	Rat (male)	Neoplastic effects		Experimental value

Conclusion

Not classified for carcinogenicity

Reproductive toxicity

Reason for revision: 15

SEAL & BOND SIL 25

No (test)data on the mixture available

Judgement is based on the relevant ingredients titanium dioxide; [in powder form containing 1 % or more of particles with aerodynamic diameter ≤ 10 µm]

	Parameter	Method	Value	Exposure time	Species	Effect	- 0.	Value determination
Developmental toxicity (Oral (stomach tube))	NOAEL	OECD 414	1000 mg/kg bw/day	2 weeks (7 days / week)	Rat	No effect		Experimental value
Maternal toxicity (Oral (stomach tube))	NOAEL	OECD 414	1000 mg/kg bw/day	2 weeks (7 days / week)	Rat	No effect		Experimental value
utan-2-one 0,0',0"-(meth	ylsilylidyne)trioxi	me	-		-			-
	Parameter	Method	Value	Exposure time	Species	Effect	- 0.	Value determination

	Parameter	Method	Value	Exposu

Developmental toxicity (Oral (stomach tube))	NOAEL	EPA OTS 798.4900	> 600 mg/kg bw/day	10 days (gestation, daily)	Rat	No effect	Experimental value
Maternal toxicity (Oral (stomach tube))	LOAEL	EPA OTS 798.4900	60 mg/kg bw/day	10 days (gestation, daily)		Spleen enlargement/af fection	 Experimental value
Effects on fertility (Oral (stomach tube))	NOAEL	Equivalent to OECD 416	> 200 mg/kg bw/day	13 weeks (5 days / week) - 19 weeks (5 days / week)	Rat (male / female)	No effect	Experimental value

2-butanone oxime

	Parameter	Method	Value	Exposure time	Species	Effect	- 0-	Value determination
Developmental toxicity (Oral (stomach tube))	NOAEL (F1)	OECD 414	> 600 mg/kg bw/day	10 day(s)	Rat	No effect		Experimental value
Maternal toxicity (Oral (stomach tube))	LOAEL	OECD 414	60 mg/kg bw/day	10 days (gestation, daily)	Rat	Maternal toxicity		Experimental value
Effects on fertility (Oral (stomach tube))	NOAEL (F1)	EPA TSCA testing guidelines	10 mg/kg bw/day	≥ 200 day(s)	Rat (male / female)	No effect		Experimental value

Conclusion Not classified for reprotoxic or developmental toxicity

Toxicity other effects

SEAL & BOND SIL 25

No (test)data on the mixture available

Chronic effects from short and long-term exposure

SEAL & BOND SIL 25

Skin rash/inflammation.

11.2. Information on other hazards

No evidence of endocrine disrupting properties

SECTION 12: Ecological information

12.1. Toxicity

SEAL & BOND SIL 25

No (test)data on the mixture available

Judgement of the mixture is based on the relevant ingredients

titanium dioxide; [in powder form containing 1 % or more of particles with aerodynamic diameter \leq 10 µm]

	Parameter	Method	Value	Duration	Species	Test design	Fresh/salt	Value determination
							water	
Acute toxicity fishes	LC50		> 1000 mg/l		Pisces		Fresh water	
Acute toxicity crustacea	EC50		> 1000 mg/l		Invertebrata		Fresh water	
Toxicity algae and other aquatic plants	EC50	OECD 201	> 100 mg/l			Static system	Fresh water	Experimental value; Growth rate
	NOEC	OECD 201	≥ 100 mg/l			Static system		Experimental value; Growth rate

	Parameter	Method	Value	Duration	Species	Test design	Fresh/salt water	Value determinatio
Acute toxicity fishes	LC50	OECD 203	> 100 mg/l	96 h	Oryzias latipes	Semi-static system	Fresh water	Experimental value; GLP
Acute toxicity crustacea	EC50	OECD 202	201 mg/l	48 h	Daphnia magna	Static system	Fresh water	Experimental value; Locomotor effect
Toxicity algae and other aquatic plants	ErC50	OECD 201	16 mg/l	72 h	Pseudokirchneri ella subcapitata	Static system	Fresh water	Experimental value; GLP
	NOEC	OECD 201	2.6 mg/l	72 h	Pseudokirchneri ella subcapitata	Static system	Fresh water	Experimental value; Growth rate
Long-term toxicity fish	NOEC	OECD 204	50 mg/l	14 day(s)	Oryzias latipes	Flow- through system	Fresh water	Experimental value; GLP
Long-term toxicity aquatic crustacea	NOEC	OECD 211	≥ 100 mg/l	21 day(s)	Daphnia magna	Semi-static system	Fresh water	Experimental value; GLP
Toxicity aquatic micro- organisms	EC50	OECD 209	> 300 mg/l	3 h	Activated sludge	Static system	Fresh water	Experimental value; GLP
<u>butanone oxime</u>								
	Parameter	Method	Value	Duration	Species	Test design	Fresh/salt water	Value determination
Acute toxicity fishes	LC50	OECD 203	> 100 mg/l	96 h	Oryzias latipes	Semi-static system	Fresh water	Experimental value; Nominal concentration
Acute toxicity crustacea	EC50	OECD 202	201 mg/l	48 h	Daphnia magna	Static system	Fresh water	Experimental value; Locomotor effect
Toxicity algae and other aquatic plants	ErC50	OECD 201	11.8 mg/l	72 h	Selenastrum capricornutum	Static system	Fresh water	Experimental value; Nominal concentration
	NOEC	OECD 201	2.56 mg/l	72 h	Selenastrum capricornutum	Static system	Fresh water	Experimental value; Growth rate
Long-term toxicity fish	NOEC	OECD 204	≥ 100 mg/l	14 day(s)	Oryzias latipes	Flow- through system	Fresh water	Experimental value; Lethal
Long-term toxicity aquatic crustacea	NOEC	OECD 211	≥ 100 mg/l	21 day(s)	Daphnia magna	Semi-static system	Fresh water	Experimental value; Reproduction
Toxicity aquatic micro- organisms	EC50		281 mg/l	17 h	Pseudomonas putida	Static system	Fresh water	Experimental value; Growth

Conclusion

Not classified as dangerous for the environment according to the criteria of Regulation (EC) No 1272/2008

12.2. Persistence and degradability

butan-2-one 0,0',0"-(methylsilylidyne)trioxime

Method	Value	Duration	Value determination
OECD 301C	20 % - 28 %	28 day(s)	Experimental value
alf-life water (t1/2 water)			
Method	Value	Primary	Value determination
		degradation/mineralisation	
EU Method C.7	< 1 h; GLP	Primary degradation	Experimental value

Biodegradation water

Method	Value	Duration	Value determination
Equivalent to OECD 301C	14.5 % - 27 %	21 day(s)	Experimental value

Conclusion

Water

Contains non readily biodegradable component(s)

12.3. Bioaccumulative potential

SEAL & BOND SIL 25

Log Kow

Method	Remark	Value	Temperature	Value determination
	Not applicable (mixture)			

<u>titanium dioxide;</u> [in powder form containing 1 % or more of particles with aerodynamic diameter \leq 10 μ m]

Log Kow

Method	Remark	Value	Temperature	Value determination
	No data available			

Reason for revision: 15

butan-2-one O,O',O''-(methylsilylidyne)trioxime

Parameter	Method		Value	Duration	Species		Value determination
BCF		0	0.5 - 5.8	6 week(s)	Cyprinu	s carpio	Experimental value
og Kow	·						
Method		Remark		Value		Temperature	Value determination
				0.36			Experimental value
butanone oxime							
BCF fishes							
	Method	N	Value	Duration	Species		Value determination
BCF fishes	Method OECD 305		Value 0.5 - 5.8; GLP		Species Cyprinus		
BCF fishes Parameter				Duration	·		Value determination
BCF fishes Parameter BCF				Duration	·		Value determination
BCF fishes Parameter BCF Log Kow	OECD 305			Duration 42 day(s)	·	s carpio	Value determination Experimental value

12.4. Mobility in soil

butan-2-one O,O',O"-(methylsilylidyne)trioxime

(log) Koc			
Parameter	Method	Value	Value determination
log Koc	SRC PCKOCWIN v2.0	5.481	Calculated value
<u>2-butanone oxime</u>			
(log) Koc			
Parameter	Method	Value	Value determination
log Koc	SRC PCKOCWIN v2.0	0.55	QSAR

Conclusion

Contains component(s) that adsorb(s) into the soil

Contains component(s) with potential for mobility in the soil

12.5. Results of PBT and vPvB assessment

Does not contain component(s) that meet(s) the criteria of PBT and/or vPvB as listed in Annex XIII of Regulation (EC) No 1907/2006.

12.6. Endocrine disrupting properties

No evidence of endocrine disrupting properties

12.7. Other adverse effects

SEAL & BOND SIL 25 Greenhouse gases None of the known components is included in the list of fluorinated greenhouse gases (Regulation (EU) No 517/2014) Ozone-depleting potential (ODP) Not classified as dangerous for the ozone layer (Regulation (EC) No 1005/2009) Groundwater Groundwater pollutant

2-butanone oxime Groundwater Groundwater pollutant

SECTION 13: Disposal considerations

The information in this section is a general description. If applicable and available, exposure scenarios are attached in annex. Always use the relevant exposure scenarios that correspond to your identified use.

13.1. Waste treatment methods

13.1.1 Provisions relating to waste

European Union

Can be considered as non hazardous waste according to Directive 2008/98/EC, as amended by Regulation (EU) No 1357/2014 and Regulation (EU) No 2017/997.

Waste material code (Directive 2008/98/EC, Decision 2000/0532/EC).

08 04 10 (wastes from MFSU of adhesives and sealants (including waterproofing products): waste adhesives and sealants other than those mentioned in 08 04 09). Depending on branch of industry and production process, also other waste codes may be applicable.

13.1.2 Disposal methods

Remove waste in accordance with local and/or national regulations. Dispose of the small quantities as household waste. Do not discharge into drains or the environment. Dispose of at authorized waste collection point.

13.1.3 Packaging/Container

European Union

Waste material code packaging (Directive 2008/98/EC).

- 15 01 01 (paper and cardboard packaging).
- 15 01 02 (plastic packaging).
- 15 01 04 (metallic packaging).

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SECTION 14: Transport information

Road (ADR), Rail (RID), Inland waterways (ADN), Sea (IMDG/IMSBC), Air (ICAO-TI/IATA-DGR)

14. <u>1. UN number</u>		
Transport	Not subject	
14.2. UN proper shipping name		
14.3. Transport hazard class(es)		
Hazard identification number		
Class		
Classification code		
14.4. Packing group		
Packing group		
Labels		
14.5. Environmental hazards		
Environmentally hazardous substance mark	no	
14.6. Special precautions for user		
Special provisions		
Limited quantities		
14.7. Maritime transport in bulk according to IMO instruments	5	
Annex II of MARPOL 73/78	Not applicable, based on available data	

SECTION 15: Regulatory information

15.1. Safety, health and environmental regulations/legislation specific for the substance or mixture <u>European legislation:</u>

VOC content Directive 2010/75/EU

VOC content	Remark
0.0 %	

REACH Annex XVII - Restriction

Contains component(s) subject to restrictions of Annex XVII of Regulation (EC) No 1907/2006: restrictions on the manufacture, placing on the market and use of certain dangerous substances, mixtures and articles.

	Designation of the substance, of the group of	Conditions of restriction
	substances or of the mixture	
butan-2-one O,O',O''-(methylsilylidyne) trioxime 2-butanone oxime	Liquid substances or nixtures fulfilling the criteria for any of the following hazard classes or categories set out in Annex I to Regulation (EC) No 1272/2008: (a) hazard classes 2.1 to 2.4, 2.6 and 2.7, 2.8 types A and B, 2.9, 2.10, 2.12, 2.13 categories 1 and 2, 2.14 categories 1 and 2, 2.15 types A to F; (b) hazard classes 3.1 to 3.6, 3.7 adverse effects on sexual function and fertility or on development, 3.8 effects other than narcotic effects, 3.9 and 3.10; (c) hazard class 4.1; (d) hazard class 5.1.	 Shall not be used in: ornamental articles intended to produce light or colour effects by means of different phases, for example in ornamental lamps and ashtrays, tricks and jokes, games for one or more participants, or any article intended to be used as such, even v ornamental aspects, Articles not complying with paragraph 1 shall not be placed on the market. Shall not be placed on the market if they contain a colouring agent, unless required for fiscal reasons, or perfume, or both, if they:
· 2-butanone oxime	Substances falling within one or more of the following points: (a) substances classified as any of the following in Part 3 of Annex VI to Regulation (EC) No 1272/2008: — carcinogen category 1A, 1B or 2, or germ cell mutagen category 1A, 1B or 2 2, but excluding any such substances classified due to effects only following exposure by inhalation — reproductive toxicant category 1A, 1B or 2 but excluding any such substances classified due to effects only following exposure by inhalation — skin sensitiser category 1, 1A or 1B — skin corrosive category 1, 1A, 1B or 1C or	 Shall not be placed on the market in mixtures for use for tattooing purposes, and mixt containing any such substances shall not be used for tattooing purposes, after 4 January 2022 if the substance or substances in question is or are present in the following circumstances: (a) in the case of a substance classified in Part 3 of Annex VI to Regulation (EC) No 1272/2008 as carcinogen category 1A, 1B or 2, or germ cell mutagen category 1A, 1B or 2 the substance is present in the mixture in a concentration equal to or greater than 0,00005 9 weight; (b) in the case of a substance classified in Part 3 of Annex VI to Regulation (EC) No 1272/2008 as reproductive toxicant category 1A, 1B or 2, the substance is present in the mixture in a concentration equal to or greater than 0,001 % by weight; (c) in the case of a substance classified in Part 3 of Annex VI to Regulation (EC) No 1272/2008 as sin sensitiser category 1A, 1B or 1B, the substance is present in the mixture in a concentration equal to or greater than 0,001 % by weight; (d) in the case of a substance classified in Part 3 of Annex VI to Regulation (EC) No 1272/2008 as skin sensitiser category 1, 1A or 1B, the substance is present in the mixture in a concentration equal to or greater than 0,001 % by weight; (d) in the case of a substance classified in Part 3 of Annex VI to Regulation (EC) No 1272/2008 as skin corrosive category 1, 1A, 1B or 1C or skin irritant category 2, or as
son for revision: 15		Publication date: 2001-03-15 Date of revision: 2021-04-27

Revision number: 0401

JLAL & DO	
skin irritant category 2 — serious eye damage category 1 or eye irritant category 2 (b) substances listed in Annex II to Regulation (EC) No 1223/2009 for which a condition is specified in at least one of the columns g, h and i of the table in that Annex (d) substances listed in Appendix 13 to this Annex. The ancillary requirements in paragraphs 7 and 8 of column 2 of this entry apply to all mixtures for use for tattooing purposes, whether or not they contain a substance falling within points (a) to (d) of this column of this entry.	 serious eye damage category 1 or eye inflant category 2, the substance is present in the mixture in a concentration equal to or greater than: (i) 0.15 Works, the substance is used solely as a pH regulator; (ii) 0.01 S Works, the inflant cases; (ii) in the case of a substance listed in Annex II to Regulation (EC) No 1223/2009 (f) the substance is present in the mixture in a concentration equal to or greater than 0,00005 K by weight; (ii) The substance is present in the mixture in a concentration equal to or greater than 0,00005 K by weight; (iii) The substance is present in the mixture in a concentration equal to or greater than 0,00005 K by weight; (iii) The substance is present in the mixture in a concentration equal to or greater than 0,00005 K by weight; (iii) The substance is present in the mixture in a concentration, or in some other way, that does not accord with the condition is specified in column h (Maximum Concentration in ready for use preparation) or column i(Other) of the table in Annex IV to Regulation (EC) No 1223/2009, the substance is present in the mixture in a concentration, our is some other way, that does not accord with the condition is specified in that abstance in that Appendix. 2. For the purposes of this entry use of a mixture "for tation gruposes" means injection or introduction on the appendix 13 to this Annex, IV to autostance interation equal to or greater than the concentration limit is a domination in that Appendix. 3. If a substance intel Appendix 13 alis within more than os of points, mixer bidding and micro-pigmentation), with the aim of making a mark or design on his or her body. 3. If a substance intel (In Appendix 13 alis within one or more of points of that substance. 4. Sway of deregation, paragraph 1 shall not appendix 13 alis within one or more of points of that substance. 4. Sway of deregation, paragraph 1 shall not appendix 13 alis sith an one of points

Reason for revision: 15

Publication date: 2001-03-15 Date of revision: 2021-04-27

Revision number: 0401

		SEAL &	BOND SIL 25	
			9. This entry does not apply to substances that are gases at tempera pressure of 101,3 kPa, or generate a vapour pressure of more than 3 of 50 °C, with the exception of formaldehyde (CAS No 50-00-0, EC N 10. This entry does not apply to the placing on the market of a mixtu purposes, or to the use of a mixture for tattooing purposes, when pl exclusively as a medical device or an accessory to a medical device, Regulation (EU) 2017/745, or when used exclusively as a medical device or an accessory to a medical device Regulation (EU) 2017/745 and of this Regulation shall apply cumulat	300 kPa at temperatur o 200-001-8). ure for use for tattooir laced on the market within the meaning of vice or an accessory t market or use may m ce, the requirements of
<u>National legislation E</u> _SEAL & BOND SIL				
No data availab National legislation 1				
SEAL & BOND SIL	. 25	i		
Waterbezwaarli 2-butanone oxime	,	Z (1); Algemene Beoordelingsmet	hodiek (ABM)	
SZW - Lijst van kankerverwekke	ende stoffen	butanonoxim; Listed in SZW-list o	f carcinogenic substances	
National legislation F SEAL & BOND SIL No data availab	. 25			
titanium dioxide;	[in powder form	i	es with aerodynamic diameter ≤ 10 μm]	
Catégorie cancé	erogène	Titane (dioxyde de), en Ti; C2		
National legislation (SEAL & BOND SIL				
WGK	lin nowdar farm		Umgang mit wassergefährdenden Stoffen (AwSV) - 18. April 2017	
titanium dioxide; TA-Luft	In powaer form	5.2.1	es with aerodynamic diameter ≤ 10 μm]	
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DMEL	Derived Minimal Effect Level
DNEL	Derived No Effect Level
EC50	Effect Concentration 50 %
ErC50	EC50 in terms of reduction of growth rate
LC50	Lethal Concentration 50 %
LD50	Lethal Dose 50 %
NOAEL	No Observed Adverse Effect Level
NOEC	No Observed Effect Concentration
OECD	Organisation for Economic Co-operation and Development
PBT	Persistent, Bioaccumulative & Toxic
PNEC	Predicted No Effect Concentration
STP	Sludge Treatment Process
vPvB	very Persistent & very Bioaccumulative

The information in this safety data sheet is based on data and samples provided to BIG. The sheet was written to the best of our ability and according to the state of knowledge at that time. The safety data sheet only constitutes a guideline for the safe handling, use, consumption, storage, transport and disposal of the substances/preparations/mixtures mentioned under point 1. New safety data sheets are written from time to time. Only the most recent versions may be used. Unless indicated otherwise word for word on the safety data sheet, the information does not apply to substances/preparations/mixtures in purer form, mixed with other substances or in processes. The safety data sheet offers no quality specification for the substances/preparations/mixtures in question. Compliance with the instructions in this safety data sheet does not release the user from the obligation to take all measures dictated by common sense, regulations and recommendations or which are necessary and/or useful based on the real applicable circumstances. BIG does not guarantee the accuracy or exhaustiveness of the information provided and cannot be held liable for any changes by third parties. This safety data sheet is only to be used within the European Union, Switzerland, Iceland, Norway and Liechtenstein. Any use outside of this area is at your own risk. Use of this safety data sheet is subject to the licence and liability limiting conditions as stated in your BIG licence agreement or when this is failing the general conditions of BIG. All intellectual property rights to this sheet are the property of BIG and its distribution and reproduction are limited. Consult the mentioned agreement/conditions for details.

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