according to Regulation (EC) No. 1907/2006 (REACH)



Silicone oil TT 3, dried, low viscosity, 3 cSt

article number: **1952** Version: **3.0 en** Replaces version of: 24.10.2022 Version: (2)

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

1952

613-156-5

63148-62-9

Polydimethylsiloxane

Laboratory chemical

1.1 Product identifier

Identification of the substance

Article number

Registration number (REACH)

EC number

CAS number

Alternative name(s)

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

Relevant identified uses:

Uses advised against:

Laboratory and analytical use Do not use for private purposes (household).

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The substance does not require registration according to Regulation (EC) No 1907/2006 [REACH].

Food, drink and animal feedingstuffs.

1.3 Details of the supplier of the safety data sheet

Carl Roth GmbH + Co. KG Schoemperlenstr. 3-5 D-76185 Karlsruhe Germany

Telephone:+49 (0) 721 - 56 06 0 **Telefax:** +49 (0) 721 - 56 06 149 **e-mail:** sicherheit@carlroth.de **Website:** www.carlroth.de

Competent person responsible for the safety data Department Health, Safety and Environment sheet:

e-mail (competent person):

sicherheit@carlroth.de

1.4 Emergency telephone number

SECTION 2: Hazards identification

2.1 Classification of the substance or mixture

Classification according to Regulation (EC) No 1272/2008 (CLP)

This substance does not meet the criteria for classification in accordance with Regulation No 1272/ 2008/EC.

2.2 Label elements

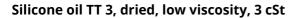
Labelling according to Regulation (EC) No 1272/2008 (CLP)

not required

2.3 Other hazards

Special danger of slipping by leaking/spilling product.

date of compilation: 07.04.2021 Revision: 03.03.2024 according to Regulation (EC) No. 1907/2006 (REACH)





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Results of PBT and vPvB assessment

The substance was identified as a PBT (persistent, bioaccumulative and toxic). The substance was identified as a vPvB (very persistent and very bioaccumulative). Non-classified PBT substance. Non-classified vPvB substance.

Endocrine disrupting properties

The substance has an endocrine disrupting potential.

SECTION 3: Composition/information on ingredients

3.1 Substances

Name of substance	Silicone oil
Molecular formula	(C₂H₀OSi)n
CAS No	63148-62-9
EC No	613-156-5

Impurities/additives/constituents:

Name of substance	Identifier	Wt%
Dodecamethylcyclohexasiloxane	CAS No 540-97-6	0,1 – 3
	EC No 208-762-8	
Decamethylcyclopentasiloxane	CAS No 541-02-6	0,1 - 3
	EC No 208-764-9	
Octamethylcyclotetrasiloxane	CAS No 556-67-2	0,1 – 1
	EC No 209-136-7	
	Index No 014-018-00-1	

Remarks

For full text of abbreviations: see SECTION 16

SECTION 4: First aid measures

4.1 Description of first aid measures



General notes

No special measures are necessary.

Following inhalation

Provide fresh air.

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Following skin contact

Wash with plenty of soap and water.

Following eye contact

Rinse cautiously with water for several minutes.

Following ingestion

Rinse mouth. Call a doctor if you feel unwell.

- **4.2** Most important symptoms and effects, both acute and delayed Symptoms and effects are not known to date.
- **4.3 Indication of any immediate medical attention and special treatment needed** none

SECTION 5: Firefighting measures

5.1 Extinguishing media



Suitable extinguishing media

co-ordinate firefighting measures to the fire surroundings! water spray, dry extinguishing powder, BC-powder, carbon dioxide (CO₂)

Unsuitable extinguishing media

water jet

5.2 Special hazards arising from the substance or mixture

Combustible.

Hazardous combustion products

In case of fire may be liberated: Carbon monoxide (CO), Carbon dioxide (CO₂)

5.3 Advice for firefighters

In case of fire and/or explosion do not breathe fumes. Fight fire with normal precautions from a reasonable distance. Wear self-contained breathing apparatus.

SECTION 6: Accidental release measures

6.1 Personal precautions, protective equipment and emergency procedures



For non-emergency personnel

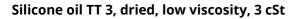
Special danger of slipping by leaking/spilling product.

6.2 Environmental precautions

Keep away from drains, surface and ground water. Retain contaminated washing water and dispose of it.

6.3 Methods and material for containment and cleaning up

according to Regulation (EC) No. 1907/2006 (REACH)





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Advice on how to contain a spill

Covering of drains.

Other information relating to spills and releases

Place in appropriate containers for disposal.

6.4 Reference to other sections

Hazardous combustion products: see section 5. Personal protective equipment: see section 8. Incompatible materials: see section 10. Disposal considerations: see section 13.

SECTION 7: Handling and storage

7.1 Precautions for safe handling

Provision of sufficient ventilation.

Advice on general occupational hygiene

Keep away from food, drink and animal feedingstuffs.

7.2 Conditions for safe storage, including any incompatibilities

Keep container tightly closed.

Incompatible substances or mixtures

Observe hints for combined storage.

Consideration of other advice:

Specific designs for storage rooms or vessels

Recommended storage temperature: 15 - 25 °C

7.3 Specific end use(s)

No information available.

SECTION 8: Exposure controls/personal protection

8.1 Control parameters

National limit values

Occupational exposure limit values (Workplace Exposure Limits)

This information is not available.

Relevant DNELs of components							
Name of sub- stance	CAS No	End- point	Threshol d level	Protection goal, route of exposure	Used in	Exposure time	
Dodecamethylcyclo- hexasiloxane	540-97-6	DNEL	11 mg/m³	human, inhalat- ory	worker (industry)	chronic - systemic effects	
Dodecamethylcyclo- hexasiloxane	540-97-6	DNEL	1,22 mg/ m³	human, inhalat- ory	worker (industry)	chronic - local ef- fects	
Dodecamethylcyclo- hexasiloxane	540-97-6	DNEL	6,1 mg/m³	human, inhalat- ory	worker (industry)	acute - local ef- fects	
Decamethylcyclo- pentasiloxane	541-02-6	DNEL	97,3 mg/ m³	human, inhalat- ory	worker (industry)	chronic - systemic effects	
Decamethylcyclo- pentasiloxane	541-02-6	DNEL	97,3 mg/ m³	human, inhalat- ory	worker (industry)	acute - systemic effects	

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Relevant DNELs						
Name of sub- stance	CAS No	End- point	Threshol d level	Protection goal, route of exposure	Used in	Exposure time
Decamethylcyclo- pentasiloxane	541-02-6	DNEL	24,2 mg/ m ³	human, inhalat- ory	worker (industry)	chronic - local ef fects
Decamethylcyclo- pentasiloxane	541-02-6	DNEL	24,2 mg/ m ³	human, inhalat- ory	worker (industry)	acute - local ef- fects
Octamethylcyclotet- rasiloxane	556-67-2	DNEL	73 mg/m ³	human, inhalat- ory	worker (industry)	chronic - system effects
Octamethylcyclotet- rasiloxane	556-67-2	DNEL	73 mg/m ³	human, inhalat- ory	worker (industry)	acute - systemi effects
Octamethylcyclotet- rasiloxane	556-67-2	DNEL	73 mg/m ³	human, inhalat- ory	worker (industry)	chronic - local e fects
Octamethylcyclotet- rasiloxane	556-67-2	DNEL	73 mg/m ³	human, inhalat- ory	worker (industry)	acute - local ef fects
Relevant PNECs	of compone	ents				
Name of sub- stance	CAS No	End- point	Threshol d level	Organism	Environmental compartment	Exposure tim
Dodecamethylcyclo- hexasiloxane	540-97-6	PNEC	1 ^{mg} / _l	aquatic organ- isms	sewage treatment plant (STP)	short-term (sing instance)
Dodecamethylcyclo- hexasiloxane	540-97-6	PNEC	13 ^{mg} / _{kg}	aquatic organ- isms	freshwater sedi- ment	short-term (sing instance)
Dodecamethylcyclo-	540-97-6	PNFC	1.3 ^{mg} /kg	aquatic organ-	marine sediment	short-term (sinc

hexasiloxane	570 570	TNEC	15 / Kg	isms	ment	instance)
Dodecamethylcyclo- hexasiloxane	540-97-6	PNEC	1,3 ^{mg} / _{kg}	aquatic organ- isms	marine sediment	short-term (single instance)
Dodecamethylcyclo- hexasiloxane	540-97-6	PNEC	3,77 ^{mg} / _{kg}	terrestrial organ- isms	soil	short-term (single instance)
Decamethylcyclo- pentasiloxane	541-02-6	PNEC	1,2 ^{µg} / _l	aquatic organ- isms	freshwater	short-term (single instance)
Decamethylcyclo- pentasiloxane	541-02-6	PNEC	0,12 ^{µg} / _l	aquatic organ- isms	marine water	short-term (single instance)
Decamethylcyclo- pentasiloxane	541-02-6	PNEC	10 ^{mg} / _l	aquatic organ- isms	sewage treatment plant (STP)	short-term (single instance)
Decamethylcyclo- pentasiloxane	541-02-6	PNEC	11 ^{mg} / _{kg}	aquatic organ- isms	freshwater sedi- ment	short-term (single instance)
Decamethylcyclo- pentasiloxane	541-02-6	PNEC	1,1 ^{mg} / _{kg}	aquatic organ- isms	marine sediment	short-term (single instance)
Decamethylcyclo- pentasiloxane	541-02-6	PNEC	1,27 ^{mg} / _{kg}	terrestrial organ- isms	soil	short-term (single instance)
Octamethylcyclotet- rasiloxane	556-67-2	PNEC	1,5 ^{µg} / _l	aquatic organ- isms	freshwater	short-term (single instance)
Octamethylcyclotet- rasiloxane	556-67-2	PNEC	0,15 ^{µg} / _l	aquatic organ- isms	marine water	short-term (single instance)
Octamethylcyclotet- rasiloxane	556-67-2	PNEC	10 ^{mg} / _l	aquatic organ- isms	sewage treatment plant (STP)	short-term (single instance)
Octamethylcyclotet- rasiloxane	556-67-2	PNEC	3 ^{mg} / _{kg}	aquatic organ- isms	freshwater sedi- ment	short-term (single instance)
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Relevant PNECs of components								
Name of sub- stance	CAS No	End- point	Threshol d level	Organism	Environmental compartment	Exposure time		
Octamethylcyclotet- rasiloxane	556-67-2	PNEC	0,3 ^{mg} / _{kg}	aquatic organ- isms	marine sediment	short-term (single instance)		
Octamethylcyclotet- rasiloxane	556-67-2	PNEC	0,54 ^{mg} / _{kg}	terrestrial organ- isms	soil	short-term (single instance)		

8.2 Exposure controls

Individual protection measures (personal protective equipment)

Eye/face protection



Use safety goggle with side protection.

Skin protection



hand protection

Wear suitable gloves. Chemical protection gloves are suitable, which are tested according to EN 374.

• type of material

NBR (Nitrile rubber)

material thickness

>0,11 mm

• breakthrough times of the glove material

>480 minutes (permeation: level 6)

• other protection measures

Take recovery periods for skin regeneration. Preventive skin protection (barrier creams/ointments) is recommended.

Respiratory protection



Respiratory protection necessary at: Aerosol or mist formation. Type: A (against organic gases and vapours with a boiling point of > 65 $^{\circ}$ C, colour code: Brown). Usually no personal respirative protection necessary.

Environmental exposure controls

Keep away from drains, surface and ground water.

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SECTION 9: Physical and chemical properties

Information on basic physical and chemical properties 9.1 Physical state liquid Form viscous Colour colourless Odour odourless Melting point/freezing point not determined Boiling point or initial boiling point and boiling not determined range Flammability this material is combustible, but will not ignite readily Lower and upper explosion limit not determined >62 °C Flash point Auto-ignition temperature not determined >150 °C Decomposition temperature not determined pH (value) 2,7 - 3,3 ^{mm²}/_s at 25 °C Kinematic viscosity Solubility(ies) Water solubility (The study does not need to be conducted because the substance is known to be insoluble in water)

Solubility in hydrocarbons, aliphatic	soluble
Solubility in hydrocarbons, aromatic	soluble
Solubility in ethylene glycol	practically insoluble
Solubility in ethyl acetate	soluble
Solubility in n-butyl acetate	soluble
Solubility in toluene	soluble
Solubility in trichloroethylene	soluble
Solubility in methanol	practically insoluble
Solubility in trichloromethane (chloroform)	soluble
Partition coefficient	
Partition coefficient n-octanol/water (log value):	this information is not available
Vapour pressure	not determined
Density and/or relative density	

0,9 – 0,91 ^g/_{cm³} at 25 °C

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Density

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	Relative vapour density	Information on this property is not available.
	Particle characteristics	not relevant (liquid)
	Other safety parameters	
	Oxidising properties	none
9.2	Other information	
	Information with regard to physical hazard classes:	hazard classes acc. to GHS (physical hazards): not relevant
	Other safety characteristics:	
	Temperature class (EU, acc. to ATEX)	T2 Maximum permissible surface temperature on the equipment: 300°C

SECTION 10: Stability and reactivity

10.1 Reactivity

This material is not reactive under normal ambient conditions.

If heated

Vapours may form explosive mixtures with air.

10.2 Chemical stability

The material is stable under normal ambient and anticipated storage and handling conditions of temperature and pressure.

10.3 Possibility of hazardous reactions

Violent reaction with: strong oxidiser

10.4 Conditions to avoid

Keep away from heat. Decompostion takes place from temperatures above: >150 °C.

10.5 Incompatible materials

There is no additional information.

10.6 Hazardous decomposition products

Hazardous combustion products: see section 5.

SECTION 11: Toxicological information

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

Classification according to GHS (1272/2008/EC, CLP)

This substance does not meet the criteria for classification in accordance with Regulation No 1272/ 2008/EC.

Acute toxicity

Shall not be classified as acutely toxic.

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Acute toxicity								
Exposure route	Endpoint	Value	Speci	Species		ethod	Source	
oral	LD50	>5.000 ^{mg} / _{kg}	rat				TOXNET	
dermal	LD50	>2.000 ^{mg} / _{kg}	rabb	it			TOXNET	
Acute toxicity of components								
Name of su	ıbstance	CAS No	Exposure route	Endp	ooint	Value	Species	
Dodecamethylcyc	lohexasiloxane	540-97-6	oral	LD	50	>2.000 ^{mg} /µ	_{kg} rat	
Dodecamethylcyc	lohexasiloxane	540-97-6	dermal	LD50		>2.000 ^{mg} /µ	_{(g} rat	
Decamethylcyclo	pentasiloxane	541-02-6	oral	LD	50	>5.000 ^{mg} /µ	_{kg} rat	
Decamethylcyclo	pentasiloxane	541-02-6	inhalation: dust/mist	LC	50	8,67 ^{mg} / _l /4	h rat	
Decamethylcyclo	ylcyclopentasiloxane 541-02-6 dermal LD50		LD50 >2		_{(g} rabbit			
Octamethylcyclotetrasiloxane		556-67-2	oral	LD	50	>4.800 ^{mg} /µ	_{kg} rat	
Octamethylcyclo	tetrasiloxane	556-67-2	inhalation: dust/mist	LC	50	36 ^{mg} / _l /4h	rat	

Skin corrosion/irritation

Shall not be classified as corrosive/irritant to skin.

Serious eye damage/eye irritation

Shall not be classified as seriously damaging to the eye or eye irritant.

Respiratory or skin sensitisation

Shall not be classified as a respiratory or skin sensitiser.

Germ cell mutagenicity

Shall not be classified as germ cell mutagenic.

Carcinogenicity

Shall not be classified as carcinogenic.

Reproductive toxicity

Shall not be classified as a reproductive toxicant.

Specific target organ toxicity - single exposure

Shall not be classified as a specific target organ toxicant (single exposure).

Specific target organ toxicity - repeated exposure

Shall not be classified as a specific target organ toxicant (repeated exposure).

Aspiration hazard

Shall not be classified as presenting an aspiration hazard.

Symptoms related to the physical, chemical and toxicological characteristics

• If swallowed

Data are not available.

according to Regulation (EC) No. 1907/2006 (REACH)



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• If in eyes

Data are not available.

If inhaled

Data are not available.

• If on skin

Data are not available.

• Other information

Health effects are not known. This information is based upon the present state of our knowledge.

11.2 Endocrine disrupting properties

Endocrine disrupting chemicals (EDC)						
Name of substance	CAS No	Combined cat- egory	Human health category	Wildlife cat- egory		
Octamethylcyclotetrasiloxane	556-67-2	CAT1	CAT1	CAT3b		

Legend

CAT1 Category 1 - evidence of endocrine disruption in at least one species using intact animals CAT3b Category 3b - no evidence of endocrine disruption or no data available

11.3 Information on other hazards

There is no additional information.

SECTION 12: Ecological information

12.1 Toxicity

Shall not be classified as hazardous to the aquatic environment.

Aquatic toxicity (acute) of components								
Name of sub- stance	CAS No	Endpoint	Value	Species	Exposure time			
Dodecamethylcyclo- hexasiloxane	540-97-6	ErC50	>2 ^{µg} / _l	algae	72 h			
Decamethylcyclo- pentasiloxane	541-02-6	LC50	>16 ^{µg} / _l	fish	96 h			
Decamethylcyclo- pentasiloxane	541-02-6	EC50	>2,9 ^{µg} / _l	aquatic invertebrates	48 h			
Octamethylcyclotet- rasiloxane	556-67-2	LC50	>22 ^{µg} / _l	fish	96 h			
Octamethylcyclotet- rasiloxane	556-67-2	EC50	>15 ^{µg} / _l	aquatic invertebrates	48 h			
Octamethylcyclotet- rasiloxane	556-67-2	ErC50	>22 ^{µg} / _l	algae	96 h			

according to Regulation (EC) No. 1907/2006 (REACH)



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Aquatic toxicity (chronic) of components								
Name of sub- stance	CAS No	Endpoint	Value	Species	Exposure time			
Dodecamethylcyclo- hexasiloxane	540-97-6	EC50	>100 ^{mg} / _l	microorganisms	3 h			
Decamethylcyclo- pentasiloxane	541-02-6	EC50	>15 ^{µg} / _l	aquatic invertebrates	21 d			
Octamethylcyclotet- rasiloxane	556-67-2	EC50	>15 ^{µg} / _l	aquatic invertebrates	21 d			

12.2 Persistence and degradability

Biodegradation

Not readily biodegradable.

Degradability of components							
Name of substance	CAS No	Process	Degrada- tion rate	Time	Method	Source	
Dodecamethyl- cyclohexasilox- ane	540-97-6	carbon dioxide generation	4,47 %	28 d		ECHA	
Decamethyl- cyclopentas- iloxane	541-02-6	carbon dioxide generation	0,14 %	28 d		ECHA	
Octamethylcyc- lotetrasiloxane	556-67-2	carbon dioxide generation	3,7 %	29 d		ECHA	

12.3 Bioaccumulative potential

Data are not available.

Bioaccumulative potential of components							
Name of substance	CAS No	BCF	Log KOW	BOD5/COD			
Dodecamethylcyclohexasiloxane	540-97-6	1.160	8,87 (23,6 °C)				
Decamethylcyclopentasiloxane	541-02-6	7.060	8,023 (25,3 °C)				
Octamethylcyclotetrasiloxane	556-67-2	12.400	6,488 (25,1 °C)				

12.4 Mobility in soil

Data are not available.

12.5 Results of PBT and vPvB assessment

The substance was identified as a PBT (persistent, bioaccumulative and toxic). The substance was identified as a vPvB (very persistent and very bioaccumulative).

12.6 Endocrine disrupting properties

according to Regulation (EC) No. 1907/2006 (REACH)



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Endocrine disrupting chemicals (EDC)						
Name of substance	CAS No	Combined cat- egory	Human health category	Wildlife cat- egory		
Octamethylcyclotetrasiloxane	556-67-2	CAT1	CAT1	CAT3b		

Legend

CAT1 Category 1 - evidence of endocrine disruption in at least one species using intact animals

CAT3b Category 3b - no evidence of endocrine disruption or no data available

12.7 Other adverse effects

Data are not available.

SECTION 13: Disposal considerations

13.1 Waste treatment methods



Consult the appropriate local waste disposal expert about waste disposal.

Sewage disposal-relevant information

Do not empty into drains.

Waste treatment of containers/packagings

Handle contaminated packages in the same way as the substance itself. Completely emptied packages can be recycled.

13.2 Relevant provisions relating to waste

The allocation of waste identity numbers/waste descriptions must be carried out according to the EEC, specific to the industry and process.

Properties of waste which render it hazardous

HP 10 toxic for reproduction

HP 14 ecotoxic

13.3 Remarks

Waste shall be separated into the categories that can be handled separately by the local or national waste management facilities. Please consider the relevant national or regional provisions. Non-contaminated packages may be recycled.

SECTION 14: Transport information

- 14.1 UN number or ID number
- 14.2 UN proper shipping name
- 14.3 Transport hazard class(es)
- 14.4 Packing group
- 14.5 Environmental hazards

not subject to transport regulations not assigned

none

not assigned

non-environmentally hazardous acc. to the dangerous goods regulations

14.6 Special precautions for user

There is no additional information.

14.7 Maritime transport in bulk according to IMO instruments

according to Regulation (EC) No. 1907/2006 (REACH)





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The cargo is not intended to be carried in bulk.

14.8 Information for each of the UN Model Regulations

> International Maritime Dangerous Goods Code (IMDG) - Additional information Not subject to IMDG.

International Civil Aviation Organization (ICAO-IATA/DGR) - Additional information Not subject to ICAO-IATA.

SECTION 15: Regulatory information

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

Relevant provisions of the European Union (EU)

Restrictions according to REACH, Annex XVII

Dangerous substances with restrictions (REACH, Annex XVII)					
Name of substance	Name acc. to inventory	CAS No	Restriction	No	
Dodecamethylcyclohexasiloxane	this product meets the criteria for classification in accordance with Reg- ulation No 1272/2008/EC		R3	3	
Decamethylcyclopentasiloxane	decamethylcyclopentasiloxane	541-02-6	R70	70	
Decamethylcyclopentasiloxane	this product meets the criteria for classification in accordance with Reg- ulation No 1272/2008/EC		R3	3	
Octamethylcyclotetrasiloxane	octamethylcyclotetrasiloxane	556-67-2	R70	70	
Octamethylcyclotetrasiloxane	this product meets the criteria for classification in accordance with Reg- ulation No 1272/2008/EC		R3	3	
Octamethylcyclotetrasiloxane	flammable / pyrophoric		R40	40	
Octamethylcyclotetrasiloxane	substances in tattoo inks and perman- ent make-up		R75	75	

Legend R3

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 with 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

can be used as fuel in decorative oil lamps for supply to the general public, and

present an aspiration hazard and are labelled with H304.
4. Decorative oil lamps for supply to the general public shall not be placed on the market unless they conform to the European Standard on Decorative oil lamps (EN 14059) adopted by the European Committee for Standardisation

(CEN). 5. Without prejudice to the implementation of other Union provisions relating to the classification, labelling and packaging of substances and mixtures, suppliers shall ensure, before the placing on the market, that the following requirements are met:

(a) lamp oils, labelled with H304, intended for supply to the general public are visibly, legibly and indelibly marked as follows: "Keep lamps filled with this liquid out of the reach of children"; and, by 1 December 2010, "Just a sip of lamp oil – or even sucking the wick of lamps – may lead to life-threatening lung damage";
(b) grill lighter fluids, labelled with H304, intended for supply to the general public are legibly and indelibly marked by 1 December 2010 as follows: 'Just a sip of grill lighter fluid may lead to life threatening lung damage';
(c) lamps oils and grill lighters, labelled with H304, intended for supply to the general public are packaged in black on a containers not exceeding 1 litre by 1 December 2010;

opaque containers not exceeding 1 litre by 1 December 2010.';

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Legend R40

- 1. Shall not be used, as substance or as mixtures in aerosol dispensers where these aerosol dispensers are intended for supply to the general public for entertainment and decorative purposes such as the following: metallic glitter intended mainly for decoration,
- artificial snow and frost,
- 'whoopee' cushions,
- silly string aerosols,

- imitation excrement, horns for parties, decorative flakes and foams,
- artificial cobwebs,
- stink bombs.

2. Without prejudice to the application of other Community provisions on the classification, packaging and labelling of substances, suppliers shall ensure before the placing on the market that the packaging of aerosol dispensers referred to above is marked visibly, legibly and indelibly with: (For professional users only.

3. By way of derogation, paragraphs 1 and 2 shall not apply to the aerosol dispensers referred to Article 8 (1a) of Council Directive 75/324/EEC (2).

4. The aerosol dispensers referred to in paragraphs 1 and 2 shall not be placed on the market unless they conform to

 The durements indicated.
 Shall not be placed on the market in wash-off cosmetic products in a concentration equal to or greater than 0,1 % by weight of either substance, after 31 January 2020.
 For the purposes of this entry, "wash-off cosmetic products" means cosmetic products as defined in Article 2(1)(a) of Regulation (EC) No 1223/2009 that, under normal conditions of use, are washed off with water after application. R70

according to Regulation (EC) No. 1907/2006 (REACH)

Silicone oil TT 3, dried, low viscosity, 3 cSt



article number: 1952

 stances shall not be used for tattooing purposes, after 4 January 2022 if the substance or substances in question 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 categing or greater than 0.0000 % by weight; (b) in the case of a substance classified in Part 3 of Annex VI to Regulation (EC) No 1272/2008 as carcinogen categing or greater than 0.0000 % by weight; (c) in the case of a substance classified in Part 3 of Annex VI to Regulation (EC) No 1272/2008 as skin corrosive category 1. As or 16 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. As or 16 or skin intrant category 2, or as serious yee damage category 1 or eye irritant category 2, th stores is present in the mixture in a concentration equal to or greater than 0.001 % by weight; (e) in the case of a substance listed in Annex II to Regulation (EC) No 1222/2008 as skin corrosive category 1. 1A, 1B or 1C or skin irritant category 2, or as serious equilor; (f) in the case of a substance listed in Annex II to Regulation; (g) in the case of a substance listed in Annex II to Regulation; (f) in the case of a substance listed in Annex II to Regulation; (f) in the case of a substance listed in Annex II to Regulation; (f) in the case of a substance listed in Annex II to Regulation; (f) in the case of a substance listed in Annex VI to Regulation; (f) in the case of a substance listed in Appendix 13 to this Annex, the substance in action or not read of the following kinds is specified in column g mixture in a concentration equal to or greater than 0.00005 % by weight; (f) Thot to be used in products? (f) Thot to be used in products? (f) Thot to be u	Legend	
 (a) In the case of a substance classified in Part 3 of Annex VI to Regulation (EC) No 1272/2008 as cranogen category 14, 16 or 2, or ear call or or greater than 0,0005 shy leng Part 3 of Annex VI to Regulation (EC) No 1272/2008 as skin sensitiser category 14, 16 or 2, the substance lassified in Part 3 of Annex VI to Regulation (EC) No 1272/2008 as skin sensitiser category 14, 16 or 2, the substance classified in Part 3 of Annex VI to Regulation (EC) No 1272/2008 as skin sensitiser category 14, 16 or 2, the substance classified in Part 3 of Annex VI to Regulation (EC) No 1272/2008 as skin sensitiser category 1. A or 16, the case of a substance classified in Part 3 of Annex VI to Regulation (EC) No 1272/2008 as skin corrosive category 1. A respective to the mixture in a concentration equal to or greater than 0,001 % by (0) in the case of a substance classified in Part 3 of Annex VI to Regulation (EC) No 1272/2008 as skin corrosive category 1, 14, 18 or 1C or skin intriant category 2, or as serious eye damage category 1 or eye intriant category 2, th substance is present in the mixture in a concentration equal to or greater than 0,00005 % by weight. (f) in the case of a substance losued a condition of one or more of the following kinds is specified in column 9 (Product type, Body parts) of the table in Annex VI to Regulation (EC) No 1223/2009 (the substance is present in the 0,00005 % by weight: (f) in the case of a substance losued an omotion septified in column h Maximum concentration in ready for (g) in the case of a substance in some other way, that does not acceding weight: (f) "Not to be used in eye products"; (f) "Not to be used in eye product	R75	 Shall not be placed on the market in mixtures for use for tattooing purposes, and mixtures containing any such stances shall not be used for tattooing purposes, after 4 January 2022 if the substance or substances in question is are present in the following circumstances:
 (b) in the case of a substance classified in Part 3 of Annex VI to Regulation (EC) No 1272/2008 as reproductive toxin experiment memory in the substance is present in the mixture in a concentration equal to or greater than 0.001 % in very the present of the mixture in a concentration equal to or greater than 0.001 % present in the mixture in a concentration equal to or greater than 0.001 % present in the substance is present in the mixture in a concentration equal to or greater than 0.001 % present in the substance is ubstance classified in Part 3 of Annex VI to Regulation (EC) No 1272/2008 as sin corrosive cater equary 1.1 A; 1B or 1C or sin infrant category 2, or as serious eye damage category 1 or eye irritant category 2, the case of a substance listed in Annex II to Regulation (EC) No 1223/2009 (*1), the substance is used solely as a pH regulator; (i) 0.10 % by weight; in all other cases; (ii) not case of a substance listed in Annex II to Regulation (EC) No 1223/2009 (*1), the substance is present in the mixture in a concentration equal to or greater than 0.00005 % by weight; (ii) Two to be used in products applied on mucous membranes; (iii) Two to be used in products applied on mucous membranes; (iii) Two to be used in products applied on mucous membranes; (iii) Two to be used in products applied on mucous membranes; (iii) Two to be used in products applied on mucous membranes; (iii) Two to be used in products membranes; (iii) Two to be used in products applied on mucous membranes; (iii) Two to be used in products applied on mucous membranes; (iii) Two to be used in products applied on mucous membranes; (iii) Two to be used in products applied on mucous membranes; (iii) Two to be used in products applied on mucous membranes; (iii) Two to be used in products applied on mucous membranes; (iii) Two to be used in products applied on mucous membranes; (iiii) Tw		(a) in the case of a substance classified in Part 3 of Annex VI to Regulation (EC) No 1272/2008 as carcinogen catego 1A, 1B or 2, or germ cell mutagen category 1A, 1B or 2, the substance is present in the mixture in a concentration
 (c) in the case of a substance lossified in Part 3 of Annex VI to Regulation (EC) No 1272/2008 as skin sensitiser category 2, no service service		(b) in the case of a substance classified in Part 3 of Annex VI to Regulation (EC) No 1272/2008 as reproductive toxic category 1A, 1B or 2, the substance is present in the mixture in a concentration equal to or greater than 0,001 % b
 (d) in the case of a substance classified in Part 3 of Annex VI to Regulation (EC) No 1222/2008 as skin corrosive cat a egory 1, rh substance is present in the mixture in a concentration gale to or greater than: (i) O i the substance is used solely as a pull equilator. (i) O i the substance is used solely as a pull equilator. (i) In the case of a substance listed in Annex II to Regulation (EC) No 1223/2009 (*1), the substance is present in the mixture in a concentration equal to or greater than 0.0005 % by weight; (i) in the case of a substance for which a condition of one or more of the following kinds is specified in column g (Product type, Body parts) of the table in AnneX IV to Regulation (EC) No 1223/2009, the substance is present in the mixture in a concentration due to or greater than 0.0005 % by weight; (ii) Thot to be used in eye products?; (jii) The case of a substance for which a condition is specified in column h (Maximum concentration in ready for preparation) or olumn i (Other) of the table in Annex. If to Regulation (EC) No 1223/2009, the substance is present in that case of a substance is incered to a substance or every that deso not accord with the condition specified in that condition appetified in that condition appetified in that condition specified in that condition is present in the inxiture in a concentration in this specified in column h (Maximum concentration intro a present skin, mucous membrane er eyeball, by any process or procedure (including procedures commony) referred to as permanent make. Up, cosmetic tattooing, micro-bidding and micro-pigmentation. With the amaking a mark or design on his or her body. 3. If a substance or conternation in the points in question shall apply to that substance. If a substance listed in		(c) in the case of a substance classified in Part 3 of Annex VI to Regulation (EC) No 1272/2008 as skin sensitiser cat- egory 1, 1A or 1B, the substance is present in the mixture in a concentration equal to or greater than 0,001 % by
 (e) in the case of a substance listed in Annex II to Regulation (EC) No 1223/2009 (*1), the substance is present in timixture in a concentration equal to or greater than 0,00005 % by weight; (f) in the case of a substance for which a condition of one or more of the following kinds is specified in column g (Product type, Body parats) of the table in Annex IV to Regulation (EC) No 1223/2009, the substance is present in the mixture in a concentration products applied on mucous membranes"; (ii) "Not to be used in eye products?"; (ji) in the case of a substance for which a condition is specified in column h (Maximum concentration in ready for preparation) or column i (Other) of the table in Annex IV to Regulation (EC) No 1223/2009, the substance is present the mixture in a concentration, or is some other way, that does not accord with the condition specified in that coll (h) in the case of a substance listed in Appendix 13 to this Annex, the substance is present in the mixture in a concentration or is more other way, that does not accord with the condition specified in that coll (h) in the case of a substance listed in Appendix 13 to this Annex, the substance is present in the mixture in a concentration in this operfield for that substance in that Appendix. 2. For the purposes of this entry use of a mixture "for tatiooing micro-blading and micro-pigmentation), with the a making a mark or design on his or her body. 3. If a substance not listed in Appendix 13 alls within more than one of points (a) to (g) of paragraph 1, the stricter concentration inmit specified of after 4 January 2021 to classify or re-classify 2 stance such that the substance then becomes caught by point (a), (b), (c) (c) of paragraph 1 of this entry, or suct that ithe fallowing information is farter the date referred to in paragraph 1 of this entry, or suct that ithe fallowing information is farter the date referred to in paragraph 1 of this entry, or suct at a cost with the fallowing substance		 (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 serious eye damage category 1 or eye irritant category 2, the substance is present in the mixture in a concentration equal to or greater than: (i) 0,1 % by weight, if the substance is used solely as a pH regulator;
 (ii) "Not to be used in products applied on muccus membranes"; (iii) "Not to be used in eye products"; (g) in the case of a substance for which a condition is specified in column h (Maximum concentration in repertation) or column i (Other) of the table in Annex IV to Regulation (EC) No 1223/2009, the substance is present the mixture in a concentration, or in some other way, that does not accord with the condition specified in that column i (Other) of the table in Annex IV to Regulation (EC) No 1223/2009, the substance listed in Appendix. To this Annex, the substance is present in the mixture in a concentration limit specified for that substance in that Appendix. To read the appendix Sim, mucculos membrane or eyeball, by any process or proceedure (including procedures commonly referred to as permanent make-up, cosmetic tattooing, micro-blading and micro-pigmentation), with the a making a mark or design on his or her body. 3. If a substance not listed in Appendix 13 falls within more than one of points (a) to (g) of paragraph 1, the stricted concentration limit lial down in the points in question shall apply to that substance. If a substance listed in Appendix 13 falls within more than one of points (a) to (g) of paragraph 1, the stricted concentration limit Sid CM is 10 to (g) of paragraph 1, the stricted concentration limit Sid CM is 10 to (g) of paragraph 1, the stricted concentration limit sid down in busistance. 4. By way of derogation, paragraph 1 shall not apply to the following substance sum 14 January 2023; (a) Pigment Gueen 7 (c) 124260, EC No 205-632-7472M is 1328-5340, c) (d) of paragraph 1 of this entry, or sug that it then falls within a different one of those points from the one within which it fell previously, and the date of application of that new or revised classification. 6. If Annex II or Annex IV to Regulation (EC) No 1223/2009 is amended after 4 January 2021 to lassify reading a first the date referred to in paragraph 1 of this entry, or		 (e) in the case of a substance listed in Annex II to Regulation (EC) No 1223/2009 (*1), the substance is present in th mixture in a concentration equal to or greater than 0,00005 % by weight; (f) in the case of a substance for which a condition of one or more of the following kinds is specified in column g (Product type, Body parts) of the table in Annex IV to Regulation (EC) No 1223/2009, the substance is present in the mixture in a concentration equal to or greater than 0,00005 % by weight:
 (g) in the case of a substance for which a condition is specified in column h (Maximum concentration in ready for preparation) or column i (Other) of the table in Annex / Yt to Regulation (EC) No 1223/2009, the substance is present the mixture in a concentration, or in some other way, that does not accord with the condition specified in that col (h) in the case of a substance its present in the mixture in a contration equal to or greater than the concentration limit specified for that substance is present in the mixture is a contration equal to or greater than the concentration limit specified for that substance in that Appendix. 2. For the purposes of this entry use of a mixture "for tationig, micro-blading and micro-pigmentation), with the a making a mark or design on his or her body. 3. If a substance not listed in Appendix 13 d bits within more than one of points (a) to (g) of paragraph 1, the stricte concentration limit laid down in phenolix in (uogion shall apply to that substance). If a substance listed in Appendix 10 (g) of paragraph 1, the concentration limit alid down in point (h) of a paragraph 1, the strict of a substance listed in Appendix 13 d bits and the Ar-14-48); (a) Pigment Blue 153 (C1 Ar160, EC No 20-565-7. (CS No 13-28-53-6). 5. If Part 3 of Annex VI to Regulation (EC) No 122/2/2008 is amended after 4 January 2021 to classify or re-classify 2 stance such that the substance then becomes caught by point (a), (b), (c) or (a) of paragraph 1 of this entry, or su that it then falls within a different one of those points from the one within which it fell previously, and the date of plication of that new or revised classification. 6. If Annex II to Annex IV to Regulation (EC) No 1223/2008 is amended after 4 January 2021 to classify or change theil in dynerol within a the different one of those points from the one within which it fell previously, and the amendment shall, for the purposes of applying this entry to that sub		(ii) "Not to be used in products applied on mucous membranes";
 3. If a substance not listed in Appendix 13 falls within more than one of points (a) to (g) of paragraph 1, the stricture concentration limit laid down in the points in question shall apply to that substance. 4. By way of derogation, paragraph 1 shall not apply to the following substances until 4 January 2023: (a) Pigment Blue 153: (Cl 74160, EC No 205-685-1, CAS No 1472-14-8); (b) Pigment Green 7 (Cl 74260, EC No 215-524-7, CAS No 1328-53-6). 5. If Part 3 of Annex VI to Regulation (EC) No 1272/2008 is amended after 4 January 2021 to classify or re-classify a stance such that the substance then becomes caught by point (a), (b), (c) or (d) of paragraph 1 of this entry, or suc that it then falls within a different one of those points from the one within which it fell previously, and the date of plication of that new or revised classification is after the date referred to in paragraph 1 or, as the case may be, p graph 4 of this entry. Not amendment shall, for the purposes of applying this entry to that substance, be treated taking effect on the date or application of that new or revised classification. 6. If Annex II or Annex IV to Regulation (EC) No 1223/2009 is amended after 4 January 2021 to list or change the li of a substance such that the substance the becomes caught by point (e), (f) or (g) of paragraph 1 of this entry. or such that it selfect after the date referred to in paragraph 1 of this entry. 7. Suppliers placing a mixture on the market for use for tatosoing purposes shall ensure that, after 4 January 2021 to list or change the market for use in tatoso or permanent make-up"; (b) a reference number to uniquely identify the batch; (c) the list of ingredients in accordance with the nomenclature established in the glossary of common ingredient name, by with strate of the soft required to be regarded as ingredient withis Regulation; (d) the statement "Mixture for use in tattoos or permanent make-up"; (e) the list of		(g) in the case of a substance for which a condition is specified in column h (Maximum concentration in ready for u preparation) or column i (Other) of the table in Annex IV to Regulation (EC) No 1223/2009, the substance is present the mixture in a concentration, or in some other way, that does not accord with the condition specified in that colu (h) in the case of a substance listed in Appendix 13 to this Annex, the substance is present in the mixture in a concentration equal to or greater than the concentration limit specified for that substance in that Appendix. 2. For the purposes of this entry use of a mixture "for tattooing purposes" means injection or introduction of the rure into a person's skin, mucous membrane or eyeball, by any process or procedure (including procedures commonly referred to as permanent make-up, cosmetic tattooing, micro-blading and micro-pigmentation), with the aid
 4. By way of derogation, paragraph 1 shall not apply to the following substances until 4 January 2023: (a) Pigment Blue 153: (C1 74160, EC No 205-685-1, CAS No 147-14-8); (b) Pigment Green 7 (C1 74260, EC No 215-524-7, CAS No 1328-53-6). 5. If Part 3 of Annex VI to Regulation (EC) No 1272/2008 is amended after 4 January 2021 to classify or re-classify a stance such that the substance then becomes caught by point (a), (b), (c) or (d) of paragraph 1 or, as the case may be, p graph 4 of this entry, that amendment shall, for the purposes of applying this entry to that substance, be treated taking effect on the date of application of that new or revised classification. 6. If Annex II or Annex IV to Regulation (EC) No 1223/2009 is amended after 4 January 2021 to list or change the li of a substance such that the substance then becomes caught by point (e), (f) or (g) of paragraph 1 of this entry, on such that it then falls within a different one of those points from the one within which it fell previously, and the amendment takes effect after the date referred to in paragraph 1 or, as the case may be, paragraph 4 of this entry, on such that it then falls within a different one of those points from the one within which it fell previously, and the amendment shall, for the purposes of applying this entry to that substance, be treated as taking effect from date falling 18 months after entry into force of the act by which that amendment was made. 7. Suppliers placing a mixture on the market for use for tattooing purposes shall ensure that, after 4 January 2022 mixture is marked with the following information: (a) the statement "Mixture for use in tattoos or permanent make-up"; (b) a reference number to uniquely identify the batch; (c) the list of ingredients in accordance with the nomenclature established in the glossary of common ingredient names pursuant to Article 33 of Regulation (EC) No 1223/2009, or in the absence of a common ingredient		3. If a substance not listed in Appendix 13 falls within more than one of points (a) to (g) of paragraph 1, the stricte concentration limit laid down in the points in question shall apply to that substance. If a substance listed in Apper 13 also falls within one or more of points (a) to (g) of paragraph 1, the concentration limit laid down in point (h) of
 (b) Pigment Green 7 (CI 74260, EC No 213-524-7, CAS No 1328-53-6). 5. If Part 3 of Annex VI to Regulation (EC) No 1272/2008 is amended after 4 January 2021 to classify or re-classify a stance such that the substance then becomes caught by point (a), (b), (c) or (d) of paragraph 1 of this entry, or such that it then falls within a different one of those points from the one within which it fell previously, and the date of plication of that new or revised classification is after the date referred to in paragraph 1 or, as the case may be, p graph 4 of this entry, that amendment shall, for the purposes of applying this entry to that substance, be treated taking effect on the date of application of that new or revised classification. 6. If Annex II or Annex IV to Regulation (EC) No 1223/2009 is amended after 4 January 2021 to list or change the li of a substance such that the substance then becomes caught by point (e), (f) of (g) of paragraph 1 of this entry, or such that it then falls within a different one of those points from the one within which it fell previously, and the amendment takes effect after the date referred to in paragraph 1 or, as the case may be, paragraph 4 of this entry or such that it then falls months after entry into fore of the act by which that amendment was made. 7. Suppliers placing a mixture on the market for use for tattooing purposes shall ensure that, after 4 January 2021 mixture is marked with the following information: (a) the statement "Mixture for use in tattoos or permanent make-up"; (b) a reference number to uniquely identify the batch; (c) the list of ingredients in accordance with the nomenclature established in the glossary of common ingredient names pursuant to Article 33 of Regulation (EC) No 1223/2009, or in the absence of a common ingredient name, t IUPAC name. In the absence of a common ingredient mame of upsubstance, use for tattooing purposes purities shall not be regarded as ingredient. If the name of a subs		4. By way of derogation, paragraph 1 shall not apply to the following substances until 4 January 2023:
 stance such that the substance then becomes caught by point (a), (b), (c) or (d) of paragraph 1 of this entry, or suc that it then falls within a different one of those points from the one within which it fell previously, and the date of plication of that new or revised classification. 6. If Annex II or Annex IV to Regulation of that new or revised classification. 6. If Annex II or Annex IV to Regulation (EC) No 1223/2009 is amended after 4 January 2021 to list or change the li of a substance such that the substance then becomes caught by point (e), (f) or (g) of paragraph 1 of this entry, or such that it then falls within a different one of those points from the one within which it fell previously, and the amendment takes effect after the date referred to in paragraph 1 or, as the case may be, paragraph 4 of this entry that amendment shall, for the purposes of applying this entry to that substance, be treated as taking effect from date falling 18 months after entry into force of the act by which that amendment was made. 7. Suppliers placing a mixture on the market for use for tattooing purposes shall ensure that, after 4 January 2022 mixture is marked with the following information: (a) the statement "Mixture for use in tattoos or permanent make-up"; (b) a reference number to uniquely identify the batch; (c) the list of ingredients in accordance with the nomenclature established in the glossary of common ingredient name, the IUPAC name. In the absence of a common ingredient name or IUPAC name, the CAS and EC number. Ingredients be listed in descending order by weight or volume of the ingredients at the time of formulation. "Ingredient" means y substance added during the process of formulation and present in the mixture for use for tattooing purpose purities shall not be regarded as ingredients. The name of a substance, used as ingredient within the meaning ot his entry, is already required to be stated on the label in accordance with this Regulatio		(b) Pigment Green 7 (CI 74260, EC No 215-524-7, CAS No 1328-53-6).
 6. If Annex II or Annex IV to Regulation (EC) No 1223/2009 is amended after 4 January 2021 to list or change the li of a substance such that the substance then becomes caught by point (e), (f) or (g) of paragraph 1 of this entry, or such that it then falls within a different one of those points from the one within which it fell previously, and the amendment shall, for the purposes of applying this entry to that substance, be treated as taking effect from date falling 18 months after entry into force of the act by which that amendment was made. 7. Suppliers placing a mixture on the market for use for tattooing purposes shall ensure that, after 4 January 2022 mixture is marked with the following information: (a) the statement "Mixture for use in tattoos or permanent make-up"; (b) a reference number to uniquely identify the batch; (c) the list of ingredients in accordance with the nomenclature established in the glossary of common ingredient names pursuant to Article 33 of Regulation (EC) No 1223/2009, or in the absence of a common ingredient name, t IUPAC name, In the absence of a common ingredient name or IUPAC name, the CAS and EC number. Ingredients be listed in descending order by weight or volume of the ingredients at the time of formulation. "Ingredient" mea any substance added during the process of formulation and present in the mixture for use for tattooing purpose; purities shall not be marked in accordance with the Regulation; (d) the additional statement "PH regulator" for substance, substance, used as ingredient within the meaning of this entry, is already required to be stated on the label in accordance with Regulation (EC) No 1272/2008, that ing under point (d)(i) of paragraph 1; (e) the statement "Contains nickel. Can cause allergic reactions." if the mixture contains nickel below the concentration limit specified in Appendix 13; (f) the statement "Contains nickel. Can cause allergic reactions." if the mixture contains chromium (VI) be		stance such that the substance then becomes caught by point (a), (b), (c) or (d) of paragraph 1 of this entry, or suc that it then falls within a different one of those points from the one within which it fell previously, and the date of plication of that new or revised classification is after the date referred to in paragraph 1 or, as the case may be, pa graph 4 of this entry, that amendment shall, for the purposes of applying this entry to that substance, be treated
 7. Suppliers placing a mixture on the market for use for tattooing purposes shall ensure that, after 4 January 2022 mixture is marked with the following information: (a) the statement "Mixture for use in tattoos or permanent make-up"; (b) a reference number to uniquely identify the batch; (c) the list of ingredients in accordance with the nomenclature established in the glossary of common ingredient names pursuant to Article 33 of Regulation (EC) No 1223/2009, or in the absence of a common ingredient name, t IUPAC name. In the absence of a common ingredient name or IUPAC name, the CAS and EC number. Ingredients be listed in descending order by weight or volume of the ingredients at the time of formulation. "Ingredient" mea any substance added during the process of formulation and present in the mixture for use for tattooing purposes purities shall not be regarded as ingredients. If the name of a substance, used as ingredient within the meaning of this entry, is already required to be stated on the label in accordance with Regulation (EC) No 1272/2008, that ing ent does not need to be marked in accordance with this Regulation; (d) the additional statement "pH regulator" for substances falling under point (d)(i) of paragraph 1; (e) the statement "Contains nickel. Can cause allergic reactions." if the mixture contains nickel below the concentration limit specified in Appendix 13; (d) safety instructions for use insofar as they are not already required to be stated on the label by Regulation (EC) 1272/2008. The information shall be clearly visible, easily legible and marked in a way that is indelible. The information shall be written in the official language(s) of the Member State(s) where the mixture is placed on market, unless the Member State(s) concerned provide(s) otherwise. Where necessary because of the size of the package, the information listed in the first subparagraph, except for p (a), shall be included instead in		6. If Annex II or Annex IV to Regulation (EC) No 1223/2009 is amended after 4 January 2021 to list or change the list of a substance such that the substance then becomes caught by point (e), (f) or (g) of paragraph 1 of this entry, or such that it then falls within a different one of those points from the one within which it fell previously, and the amendment takes effect after the date referred to in paragraph 1 or, as the case may be, paragraph 4 of this entry that amendment shall, for the purposes of applying this entry to that substance, be treated as taking effect from the substance is a substance.
 (b) a reference number to uniquely identify the batch; (c) the list of ingredients in accordance with the nomenclature established in the glossary of common ingredient names pursuant to Article 33 of Regulation (EC) No 1223/2009, or in the absence of a common ingredient name, t IUPAC name. In the absence of a common ingredient name or IUPAC name, the CAS and EC number. Ingredients be listed in descending order by weight or volume of the ingredients at the time of formulation. "Ingredient" mea any substance added during the process of formulation and present in the mixture for use for tattooing purposes purities shall not be regarded as ingredients. If the name of a substance, used as ingredient within the meaning of this entry, is already required to be stated on the label in accordance with Regulation (EC) No 1272/2008, that ingredient "gent does not need to be marked in accordance with this Regulation; (d) the additional statement "PH regulator" for substances falling under point (d)(i) of paragraph 1; (e) the statement "Contains nickel. Can cause allergic reactions." if the mixture contains nickel below the concentriation limit specified in Appendix 13; (f) the statement "Contains chromium (VI). Can cause allergic reactions." if the mixture contains chromium (VI) be the concentration limit specified in Appendix 13; (g) safety instructions for use insofar as they are not already required to be stated on the label by Regulation (EC) 1272/2008. The information shall be clearly visible, easily legible and marked in a way that is indelible. The information shall be written in the official language(s) of the Member State(s) where the mixture is placed on market, unless the Member State(s) concerned provide(s) otherwise. Where necessary because of the size of the package, the information listed in the first subparagraph, except for procedure with the information marked on the package or included in the instructions for use pursuant to		7. Suppliers placing a mixture on the market for use for tattooing purposes shall ensure that, after 4 January 2022 mixture is marked with the following information:
names pursuant to Article 33 of Regulation (EC) No 1223/2009, or in the absence of a common ingredient name, the IUPAC name. In the absence of a common ingredient name or IUPAC name, the CAS and EC number. Ingredients be listed in descending order by weight or volume of the ingredients at the time of formulation. "Ingredient" maany substance added during the process of formulation and present in the mixture for use for tattooing purposes purities shall not be regarded as ingredients. If the name of a substance, used as ingredient within the meaning of this entry, is already required to be stated on the label in accordance with Regulation (EC) No 1272/2008, that ingrent does not need to be marked in accordance with this Regulation; (d) the additional statement "pH regulator" for substances falling under point (d)(i) of paragraph 1; (e) the statement "Contains nickel. Can cause allergic reactions." if the mixture contains nickel below the concentr tion limit specified in Appendix 13; (f) the statement "Contains chromium (VI). Can cause allergic reactions." if the mixture contains chromium (VI) be the concentration limit specified in Appendix 13; (g) safety instructions for use insofar as they are not already required to be stated on the label by Regulation (EC) 1272/2008. The information shall be clearly visible, easily legible and marked in a way that is indelible. The information shall be created step or provide(s) of the Member State(s) where the mixture is placed on market, unless the Member State(s) concerned provide(s) of the Member State(s) where the person undergoing (a), shall be included in stead in the instructions for use. Before using a mixture for tattooing purposes, the person using the mixture shall provide the person undergoing procedure with the information marked on the package or included in the instructions for use pursuant to this pace of the grady as the person undergoing procedure with the information marked on the package or included in the instructions for use pursuant to this pace or in		(b) a reference number to uniquely identify the batch;
 ent does not need to be marked in accordance with this Regulation; (d) the additional statement "PH regulator" for substances falling under point (d)(i) of paragraph 1; (e) the statement "Contains nickel. Can cause allergic reactions." if the mixture contains nickel below the concentr tion limit specified in Appendix 13; (f) the statement "Contains chromium (VI). Can cause allergic reactions." if the mixture contains chromium (VI) be the concentration limit specified in Appendix 13; (g) safety instructions for use insofar as they are not already required to be stated on the label by Regulation (EC) 1272/2008. The information shall be clearly visible, easily legible and marked in a way that is indelible. The information shall be written in the official language(s) of the Member State(s) where the mixture is placed on market, unless the Member State(s) concerned provide(s) otherwise. Where necessary because of the size of the package, the information listed in the first subparagraph, except for place, shall be included instead in the instructions for use. Before using a mixture for tattooing purposes, the person using the mixture shall provide the person undergoing procedure with the information marked on the package or included in the instructions for use pursuant to this paragraph. 		names pursuant to Article 33 of Regulation (EC) No 1223/2009, or in the absence of a common ingredient name, t IUPAC name. In the absence of a common ingredient name or IUPAC name, the CAS and EC number. Ingredients be listed in descending order by weight or volume of the ingredients at the time of formulation. "Ingredient" mea any substance added during the process of formulation and present in the mixture for use for tattooing purposes purities shall not be regarded as ingredients. If the name of a substance, used as ingredient within the meaning c
tion limit specified in Appendix 1.3; (f) the statement "Contains chromium (VI). Can cause allergic reactions." if the mixture contains chromium (VI) be the concentration limit specified in Appendix 13; (g) safety instructions for use insofar as they are not already required to be stated on the label by Regulation (EC) 1272/2008. The information shall be clearly visible, easily legible and marked in a way that is indelible. The information shall be written in the official language(s) of the Member State(s) where the mixture is placed on market, unless the Member State(s) concerned provide(s) otherwise. Where necessary because of the size of the package, the information listed in the first subparagraph, except for p (a), shall be included instead in the instructions for use. Before using a mixture for tattooing purposes, the person using the mixture shall provide the person undergoing procedure with the information marked on the package or included in the instructions for use pursuant to this pa- graph.		ent does not need to be marked in accordance with this Regulation; (d) the additional statement "pH regulator" for substances falling under point (d)(i) of paragraph 1; (e) the statement "Contains nickel. Can cause allergic reactions." if the mixture contains nickel below the concentr
The information shall be clearly visible, easily legible and marked in a way that is indelible. The information shall be written in the official language(s) of the Member State(s) where the mixture is placed on market, unless the Member State(s) concerned provide(s) otherwise. Where necessary because of the size of the package, the information listed in the first subparagraph, except for p (a), shall be included instead in the instructions for use. Before using a mixture for tattooing purposes, the person using the mixture shall provide the person undergoing procedure with the information marked on the package or included in the instructions for use pursuant to this pa graph.		(f) the statement "Contains chromium (VI). Can cause allergic reactions." if the mixture contains chromium (VI) bel the concentration limit specified in Appendix 13; (g) safety instructions for use insofar as they are not already required to be stated on the label by Regulation (EC)
market, unless the Member State(s) concerned provide(s) otherwise. Where necessary because of the size of the package, the information listed in the first subparagraph, except for p (a), shall be included instead in the instructions for use. Before using a mixture for tattooing purposes, the person using the mixture shall provide the person undergoing procedure with the information marked on the package or included in the instructions for use pursuant to this pa graph.		The information shall be clearly visible, easily legible and marked in a way that is indelible. The information shall be written in the official language(s) of the Member State(s) where the mixture is placed on
graph.		Where necessary because of the size of the package, the information listed in the first subparagraph, except for p (a), shall be included instead in the instructions for use.
		graph.

8. Mixtures that do not contain the statement "Mixture for use in tattoos or permanent make-up" shall not be used for tattooing purposes.

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Legend

9. This entry does not apply to substances that are gases at temperature of 20 °C and pressure of 101,3 kPa, or gener-ate a vapour pressure of more than 300 kPa at temperature of 50 °C, with the exception of formaldehyde (CAS No 50-00-0, EC No 200-001-8).

10. This entry does not apply to the placing on the market of a mixture for use for tattooing purposes, or to the use of a mixture for tattooing purposes, when placed on the market of a mixture low use for tattooing purposes, of to the use of a mixture for tattooing purposes, of to the use of a mixture for tattooing purposes, of to the use of a medical device or an accessory to a medical device, within the meaning of Regulation (EU) 2017/745, or when used exclusively as a medical device or an accessory to a medical device, within the same meaning. Where the placing on the market or use may not be exclusively as a medical device or an accessory to a medical device, the requirements of Regulation (EU) 2017/745 and of this previous the tattoo of the use of the us Regulation shall apply cumulatively.

List of substances subject to authorisation (REACH, Annex XIV)/SVHC - candidate list

Substance of Very High Concern (SVHC)							
Name acc. to invent- ory	CAS No	Listed in	Remarks	Latest ap- plication date	Sunset date	Date of in- clusion	
dodecamethylcyclo- hexasiloxane	540-97- 6	Candidate list	PBT A57d vPvB A57e			27.06.2018	
decamethylcyclopentasilox- ane	541-02- 6	Candidate list	PBT A57d vPvB A57e			27.06.2018	
octamethylcyclotetrasilox- ane	556-67- 2	Candidate list	PBT A57d vPvB A57e			27.06.2018	

Legend

Candidate list Substances meeting the criteria referred to in Article 57 and for eventual inclusion in Annex XIV PBT A57d Persistent, Bioaccumulative and Toxic (article 57d) vPvB A57e Very Persistent and very Bioaccumulative (article 57e)

Seveso Directive

2012/18/EU (Seveso III)							
Νο	Dangerous substance/hazard categories	Qualifying quantity (tonnes) for the ap- plication of lower and upper-tier re- quirements	Notes				
	not assigned						

Deco-Paint Directive

VOC content	0 %
VOC content	0 ^g / _l

Industrial Emissions Directive (IED)

VOC content	0 %
VOC content	0 ^g / _l

Directive on the restriction of the use of certain hazardous substances in electrical and electronic equipment (RoHS)

not listed

Regulation concerning the establishment of a European Pollutant Release and Transfer **Register (PRTR)**

not listed

according to Regulation (EC) No. 1907/2006 (REACH)



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ater Framework Directive (WFD) .ist of pollutants (WFD)					
Name of substance	Name acc. to inventory	CAS No	Listed in	Remarks	
Octamethylcyclotetrasiloxane	Substances and preparations, or the breakdown products of such, which have been proved to pos- sess carcinogenic or mutagenic properties or properties which may affect steroidogenic, thyroid, reproduction or other endocrine- related functions in or via the aquatic environment		a)		

Legend

a) Indicative list of the main pollutants

Regulation on the marketing and use of explosives precursors

not listed

Regulation on drug precursors

not listed

Regulation on substances that deplete the ozone layer (ODS)

not listed

Regulation concerning the export and import of hazardous chemicals (PIC)

not listed

Regulation on persistent organic pollutants (POP)

not listed

Other information

Directive 94/33/EC on the protection of young people at work. Observe employment restrictions under the Maternity Protection Directive (92/85/EEC) for expectant or nursing mothers.

National inventories

Country	Inventory	Status
AU	AIIC	all ingredients are listed
CA	DSL	all ingredients are listed
CN	IECSC	all ingredients are listed
EU	ECSI	all ingredients are listed
EU	REACH Reg.	all ingredients are listed
JP	CSCL-ENCS	all ingredients are listed
KR	KECI	all ingredients are listed
MX	INSQ	not all ingredients are listed
NZ	NZIoC	all ingredients are listed
PH	PICCS	all ingredients are listed
TR	CICR	not all ingredients are listed
TW	TCSI	all ingredients are listed

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Country	Inventory Status			
US	TSCA	all ingredients are listed (ACTIVE)		
VN	NCI	all ingredients are listed		
Legend AIIC CICR CSCL-ENCS DSL ECSI IECSC INSQ KECI NCI NZIOC PICCS REACH Reg. TCSI	Australian Inventory of Industrial Chemicals Chemical Inventory and Control Regulation List of Existing and New Chemical Substances (CSCL-ENCS) Domestic Substances List (DSL) EC Substance Inventory (EINECS, ELINCS, NLP) Inventory of Existing Chemical Substances Produced or Imported in China National Inventory of Chemical Substances Korea Existing Chemicals Inventory National Chemical Inventory National Chemical Inventory New Zealand Inventory of Chemicals Philippine Inventory of Chemicals and Chemical Substances (PICCS) REACH registered substances			

TSCA Toxic Substance Control Act

15.2 Chemical safety assessment

No Chemical Safety Assessment has been carried out for this substance. According to REACH, Article 14 (1) a chemical safety assessment has been carried out for this substance or components of this mixture when the substance has been registered in quantities of 10 tonnes or more per year per registrant.

SECTION 16: Other information

Indication of changes (revised safety data sheet)

Section	Former entry (text/value)	Actual entry (text/value)	Safety- relev- ant
2.3	Results of PBT and vPvB assessment: Containing a PBT-/vPvB-substance in a concen- tration of $\ge 0,1\%$.	Results of PBT and vPvB assessment: The substance was identified as a PBT (persist- ent, bioaccumulative and toxic). The substance was identified as a vPvB (very persistent and very bioaccumulative). Non-classified PBT sub- stance. Non-classified vPvB substance.	yes
14.8	Transport of dangerous goods by road, rail and inland waterway (ADR/RID/ADN) - Additional in- formation: Not subject to ADR, RID and ADN.		yes
15.1		Substance of Very High Concern (SVHC): change in the listing (table)	yes
15.1	VOC content: 7 % , 63,7 ^g / _l	VOC content: 0 %	yes
15.1		VOC content: 0 ^g / _l	yes
15.1	VOC content: 7 %	VOC content: 0 %	yes
15.1	VOC content: 63,7 ^g / _l	VOC content: 0 ^g / _l	yes
15.1		National inventories: change in the listing (table)	yes

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Section	Former entry (text/value)	Actual entry (text/value)	Safety- relev- ant
15.2	Chemical Safety Assessment: No Chemical Safety Assessment has been car- ried out for this substance.	Chemical safety assessment: No Chemical Safety Assessment has been car- ried out for this substance. According to REACH, Article 14 (1) a chemical safety assessment has been carried out for this substance or compon- ents of this mixture when the substance has been registered in quantities of 10 tonnes or more per year per registrant.	yes

Abbreviations and acronyms

Abbr.	Descriptions of used abbreviations	
ADR	Accord relatif au transport international des marchandises dangereuses par route (Agreement concern- ing the International Carriage of Dangerous Goods by Road)	
BCF	Bioconcentration factor	
BOD	D Biochemical Oxygen Demand	
CAS	Chemical Abstracts Service (service that maintains the most comprehensive list of chemical substances)	
CLP	Regulation (EC) No 1272/2008 on classification, labelling and packaging of substances and mixtures	
COD	Chemical oxygen demand	
DGR	Dangerous Goods Regulations (see IATA/DGR)	
DNEL	Derived No-Effect Level	
EC50	Effective Concentration 50 %. The EC50 corresponds to the concentration of a tested substance causing 50 % changes in response (e.g. on growth) during a specified time interval	
EC No	The EC Inventory (EINECS, ELINCS and the NLP-list) is the source for the seven-digit EC number, an identi- fier of substances commercially available within the EU (European Union)	
EINECS	European Inventory of Existing Commercial Chemical Substances	
ELINCS	European List of Notified Chemical Substances	
ErC50	= EC50: in this method, that concentration of test substance which results in a 50 % reduction in either growth (EbC50) or growth rate (ErC50) relative to the control	
GHS	"Globally Harmonized System of Classification and Labelling of Chemicals" developed by the United Na- tions	
ΙΑΤΑ	International Air Transport Association	
IATA/DGR	Dangerous Goods Regulations (DGR) for the air transport (IATA)	
ICAO	International Civil Aviation Organization	
IMDG	International Maritime Dangerous Goods Code	
index No	The Index number is the identification code given to the substance in Part 3 of Annex VI to Regulation (EC) No 1272/2008	
LC50	Lethal Concentration 50%: the LC50 corresponds to the concentration of a tested substance causing 50 % lethality during a specified time interval	
LD50	Lethal Dose 50 %: the LD50 corresponds to the dose of a tested substance causing 50 % lethality during specified time interval	
log KOW	n-Octanol/water	
NLP	No-Longer Polymer	
PBT	Persistent, Bioaccumulative and Toxic	

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Abbr.	Descriptions of used abbreviations
PNEC	Predicted No-Effect Concentration
REACH	Registration, Evaluation, Authorisation and Restriction of Chemicals
SVHC	Substance of Very High Concern
VOC	Volatile Organic Compounds
vPvB	Very Persistent and very Bioaccumulative

Key literature references and sources for data

Regulation (EC) No 1272/2008 on classification, labelling and packaging of substances and mixtures. Regulation (EC) No. 1907/2006 (REACH), amended by 2020/878/EU.

Agreement concerning the International Carriage of Dangerous Goods by Road (ADR). International Maritime Dangerous Goods Code (IMDG). Dangerous Goods Regulations (DGR) for the air transport (IATA).

Disclaimer

This information is based upon the present state of our knowledge. This SDS has been compiled and is solely intended for this product.