

## Safety Data Sheet

SDS EU format according to COMMISSION REGULATION (EU) 2020/878
Issue date: 7/2/2015 Revision date: 1/2/2023 Supersedes version of: 6/1/2017 Version: 4.00

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

#### 1.1. Product identifier

Product form : Mixture
Name : Acrylic filler
Trade name : PROTECT 320

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

#### 1.2.1. Relevant identified uses

Use of the substance/mixture : The product is intended for professional use

#### 1.2.2. Uses advised against

No additional information available

#### 1.3. Details of the supplier of the safety data sheet

NOVOL Sp. z o.o. Żabikowska 7/9 62-052 KOMORNIKI

Poland

T 0048618109800 - F 0048618109809

www.novol.com

E-mail address of competent person responsible for the SDS: dokumentacja@novol.com

#### 1.4. Emergency telephone number

Emergency number : 112

#### **SECTION 2: Hazards identification**

#### 2.1. Classification of the substance or mixture

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

Flammable liquids, Category 3 H226 Skin corrosion/irritation, Category 2 H315

Full text of H- and EUH-statements: see section 16

#### Adverse physicochemical, human health and environmental effects

No additional information available

## 2.2. Label elements

#### Labelling according to Regulation (EC) No. 1272/2008 [CLP]

Hazard pictograms (CLP) :





GHS02

GHS07

Signal word (CLP) : Warning
Contains : xylene

Hazard statements (CLP) : H226 - Flammable liquid and vapour.

H315 - Causes skin irritation.

Precautionary statements (CLP) : P210 - Keep away from heat, hot surfaces, sparks, open flames and other ignition sources.

No smoking.

P261 - Avoid breathing vapours, spray.

P271 - Use only outdoors or in a well-ventilated area.

P280 - Wear protective gloves, protective clothing, eye protection, face protection.

P312 - Call doctor if you feel unwell.

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**EUH-statements** 

: EUH211 - Warning! Hazardous respirable droplets may be formed when sprayed. Do not breathe spray or mist.

#### 2.3. Other hazards

Contains no PBT/vPvB substances ≥ 0.1% assessed in accordance with REACH Annex XIII

The mixture does not contain substance(s) included in the list established in accordance with Article 59(1) of REACH for having endocrine disrupting properties, or is not identified as having endocrine disrupting properties in accordance with the criteria set out in Commission Delegated Regulation (EU) 2017/2100 or Commission Regulation (EU) 2018/605 at a concentration equal to or greater than 0,1 %

## **SECTION 3: Composition/information on ingredients**

#### 3.1. Substances

Not applicable

#### 3.2. Mixtures

Name	Product identifier	%	Classification according to Regulation (EC) No. 1272/2008 [CLP]
xylene substance with national workplace exposure limit(s) (GB); substance with a Community workplace exposure limit (Note C)	CAS-No.: 1330-20-7 EC-No.: 215-535-7 EC Index-No.: 601-022-00-9 REACH-no: 01-2119488216- 32	13 – 18	Flam. Liq. 3, H226 Acute Tox. 4 (Dermal), H312 Acute Tox. 4 (Inhalation), H332 Skin Irrit. 2, H315
n-butyl acetate substance with national workplace exposure limit(s) (GB); substance with a Community workplace exposure limit	CAS-No.: 123-86-4 EC-No.: 204-658-1 EC Index-No.: 607-025-00-1 REACH-no: 01-2119485493-	10 – 15	Flam. Liq. 3, H226 STOT SE 3, H336
titanium dioxide; [in powder form containing 1 % or more of particles with aerodynamic diameter ≤ 10 µm] substance with national workplace exposure limit(s) (GB) (Note V)(Note W)(Note 10)	CAS-No.: 13463-67-7 EC-No.: 236-675-5 EC Index-No.: 022-006-00-2 REACH-no: 01-2119489379- 17	< 10	Carc. 2, H351
ethyl acetate substance with national workplace exposure limit(s) (GB); substance with a Community workplace exposure limit	CAS-No.: 141-78-6 EC-No.: 205-500-4 EC Index-No.: 607-022-00-5	1 – 2	Flam. Liq. 2, H225 Eye Irrit. 2, H319 STOT SE 3, H336

Note 10 : 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  $\mu$ m.

Note C: Some organic substances may be marketed either in a specific isomeric form or as a mixture of several isomers. In this case the supplier must state on the label whether the substance is a specific isomer or a mixture of isomers.

Note V : If the substance is to be placed on the market as fibres (with diameter <  $3 \mu m$ , length >  $5 \mu m$  and aspect ratio  $\geq 3:1$ ) or particles of the substance fulfilling the WHO fibre criteria or as particles with modified surface chemistry, their hazardous properties must be evaluated in accordance with Title II of this Regulation, to assess whether a higher category (Carc. 1B or 1A) and/or additional routes of exposure (oral or dermal) should be applied.

Note W: It has been observed that the carcinogenic hazard of this substance arises when respirable dust is inhaled in quantities leading to significant impairment of particle clearance mechanisms in the lung. This note aims to describe the particular toxicity of the substance; it does not constitute a criterion for classification according to this Regulation.

Full text of H- and EUH-statements: see section 16

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#### **SECTION 4: First aid measures**

#### 4.1. Description of first aid measures

First-aid measures general : General information. Refer to section 11.

First-aid measures after inhalation : If breathing is difficult, remove victim to fresh air and keep at rest in a position comfortable

for breathing

First-aid measures after skin contact : After contact with skin, take off immediately all contaminated clothing, and wash

immediately with plenty of water and soap. Rinse skin with water/shower. If skin irritation or rash occurs: Get medical advice/attention. If skin irritation continues, consult a doctor.

First-aid measures after eye contact : Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy

to do. Continue rinsing. Call a physician immediately. In case of contact with eyes, rinse

immediately with plenty of water and seek medical advice.

First-aid measures after ingestion : If swallowed: rinse mouth. Do NOT induce vomiting. Call a physician immediately.

#### 4.2. Most important symptoms and effects, both acute and delayed

Symptoms/effects after inhalation : Vapours may cause drowsiness and dizziness.

Symptoms/effects after skin contact : Prolonged or repeated contact may cause skin to become dry.

Symptoms/effects after eye contact : May cause eye irritation.

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

Treat symptomatically.

#### **SECTION 5: Firefighting measures**

#### 5.1. Extinguishing media

Suitable extinguishing media : Dry chemical, CO2, alcohol-resistant foam or waterspray.

Unsuitable extinguishing media : Do not use a heavy water stream.

#### 5.2. Special hazards arising from the substance or mixture

Hazardous decomposition products in case of fire : Carbon monoxide. Other toxic gases.

#### 5.3. Advice for firefighters

Protection during firefighting : Do not attempt to take action without suitable protective equipment. Self-contained

breathing apparatus. Complete protective clothing.

#### **SECTION 6: Accidental release measures**

#### 6.1. Personal precautions, protective equipment and emergency procedures

## 6.1.1. For non-emergency personnel

Protective equipment : Remove ignition sources. Ensure that there is a suitable ventilation system. Avoid any direct

or indirect contact with ingredients released. Avoid contact with skin and eyes. Use personal protective equipment as required. See Section 8.

protective equipment as required. See Section of

Protective equipment : Do not attempt to take action without suitable protective equipment. See Section 8.

#### 6.2. Environmental precautions

6.1.2. For emergency responders

Avoid release to the environment. Do not allow to enter into surface water or drains. Do not allow product to reach ground water, water bodies or sewage system, even in small quantities.

#### 6.3. Methods and material for containment and cleaning up

For containment : Cover spill with non combustible material, e.g.: sand, earth, vermiculite. Mechanically

recover the product.

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#### 6.4. Reference to other sections

Disposal considerations. See Section 13.

## **SECTION 7: Handling and storage**

#### 7.1. Precautions for safe handling

Precautions for safe handling : Ensure good ventilation of the work station. Keep away from heat, hot surfaces, sparks,

open flames and other ignition sources. No smoking. Use only outdoors or in a well-

ventilated area. Wear personal protective equipment.

Hygiene measures : Wash contaminated clothing before reuse. Contaminated work clothing should not be

allowed out of the workplace. Do not eat, drink or smoke when using this product. Always

wash hands after handling the product.

#### 7.2. Conditions for safe storage, including any incompatibilities

Technical measures : Ground/bond container and receiving equipment.

Storage conditions : Store in a well-ventilated place. Keep cool. Keep container tightly closed.

#### 7.3. Specific end use(s)

No additional information available

### **SECTION 8: Exposure controls/personal protection**

#### 8.1. Control parameters

#### 8.1.1 National occupational exposure and biological limit values

xylene (1330-20-7)		
EU - Indicative Occupational Exposure Limit (IOEL)		
Local name	Xylene, mixed isomers, pure	
IOEL TWA [ppm]	50 ppm	
IOEL STEL	442 mg/m³	
IOEL STEL [ppm]	100 ppm	
Remark	Skin	
Regulatory reference	COMMISSION DIRECTIVE 2000/39/EC	
United Kingdom - Occupational Exposure Limits		
Local name	Xylene	
WEL TWA (OEL TWA) [1]	220 mg/m³ o-,m-,p- or mixed isomers	
WEL TWA (OEL TWA) [2]	50 ppm o-,m-,p- or mixed isomers	
WEL STEL (OEL STEL)	441 mg/m³ o-,m-,p- or mixed isomers	
WEL STEL (OEL STEL) [ppm]	100 ppm o-,m-,p- or mixed isomers	
Remark	Sk (Can be absorbed through the skin. The assigned substances are those for which there are concerns that dermal absorption will lead to systemic toxicity)	
Regulatory reference	EH40/2005 (Fourth edition, 2020). HSE	
United Kingdom - Biological limit values		
Local name	Xylene, o-, m-, p- or mixed isomers	
BMGV	650 mmol/mol Creatinine Parameter: methyl hippuric acid - Medium: urine - Sampling time: Post shift	
Regulatory reference	EH40/2005 (Fourth edition, 2020). HSE	

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n-butyl acetate (123-86-4)	
EU - Indicative Occupational Exposure Lin	nit (IOEL)
Local name	n-Butyl acetate
IOEL TWA [ppm]	50 ppm
IOEL STEL	723 mg/m³
IOEL STEL [ppm]	150 ppm
Regulatory reference	COMMISSION DIRECTIVE (EU) 2019/1831
United Kingdom - Occupational Exposure	Limits
Local name	Butyl acetate
WEL TWA (OEL TWA) [1]	724 mg/m³
WEL TWA (OEL TWA) [2]	150 ppm
WEL STEL (OEL STEL)	966 mg/m³
WEL STEL (OEL STEL) [ppm]	200 ppm
Regulatory reference	EH40/2005 (Fourth edition, 2020). HSE
ethyl acetate (141-78-6)	
EU - Indicative Occupational Exposure Lin	nit (IOEL)
Local name	Ethyl acetate
IOEL TWA [ppm]	200 ppm
IOEL STEL	1468 mg/m³
IOEL STEL [ppm]	400 ppm
Regulatory reference	COMMISSION DIRECTIVE (EU) 2017/164
United Kingdom - Occupational Exposure	Limits
Local name	Ethyl acetate
WEL TWA (OEL TWA) [1]	734 mg/m³
WEL TWA (OEL TWA) [2]	200 ppm
WEL STEL (OEL STEL)	1468 mg/m³
WEL STEL (OEL STEL) [ppm]	400 ppm
Regulatory reference	EH40/2005 (Fourth edition, 2020). HSE
titanium dioxide; [in powder form containing 1 % or more of particles with aerodynamic diameter ≤ 10 μm] (13463-67-7)	
United Kingdom - Occupational Exposure	Limits
Local name	Titanium dioxide
WEL TWA (OEL TWA) [1]	4 mg/m³ respirable 10 mg/m³ total inhalable
Regulatory reference	EH40/2005 (Fourth edition, 2020). HSE

## 8.1.2. Recommended monitoring procedures

Monitoring methods	
9	EN 482. Workplace exposure - General requirements for the performance of procedures for the measurement of chemical agents.

#### 8.1.3. Air contaminants formed

No additional information available

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#### 8.1.4. DNEL and PNEC

8.1.4. DNEL and PNEC		
xylene (1330-20-7)		
DNEL/DMEL (Workers)	T	
Acute - systemic effects, inhalation	289 mg/m³	
Acute - local effects, inhalation	289 mg/m³	
Long-term - systemic effects, dermal	180 mg/kg bodyweight/day	
Long-term - systemic effects, inhalation	77 mg/m³	
DNEL/DMEL (General population)		
Acute - systemic effects, inhalation	174 mg/m³	
Acute - local effects, inhalation	174 mg/m³	
Long-term - systemic effects,oral	1.6 mg/kg bodyweight/day	
Long-term - systemic effects, inhalation	14.8 mg/m³	
Long-term - systemic effects, dermal	108 mg/kg bodyweight/day	
PNEC (Water)		
PNEC aqua (freshwater)	0.327 mg/l	
PNEC aqua (marine water)	0.327 mg/l	
PNEC aqua (intermittent, freshwater)	0.327 mg/l	
PNEC (Sediment)		
PNEC sediment (freshwater)	12.46 mg/kg dwt	
PNEC sediment (marine water)	12.46 mg/kg dwt	
PNEC (Soil)		
PNEC soil	2.31 mg/kg dwt	
PNEC (STP)		
PNEC sewage treatment plant	6.58 mg/l	
n-butyl acetate (123-86-4)		
PNEC (Water)		
PNEC aqua (freshwater)	0.18 mg/l	
PNEC aqua (marine water)	0.018 mg/l	
PNEC aqua (intermittent, freshwater)	0.36 mg/l	
PNEC (Sediment)		
PNEC sediment (freshwater)	0.981 mg/kg dwt	
PNEC sediment (marine water)	0.0981 mg/kg dwt	
PNEC (Soil)		
PNEC soil	0.0903 mg/kg dwt	
PNEC (STP)		
PNEC sewage treatment plant	35.6 mg/l	
ethyl acetate (141-78-6)		
DNEL/DMEL (Workers)		
Acute - systemic effects, inhalation	1468 mg/m³	
Acute - local effects, inhalation	1468 mg/m³	

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ethyl acetate (141-78-6)		
Long-term - systemic effects, dermal	63 mg/kg bodyweight/day	
Long-term - systemic effects, inhalation	734 mg/m³	
Long-term - local effects, inhalation	734 mg/m³	
DNEL/DMEL (General population)		
Acute - systemic effects, inhalation	734 mg/m³	
Acute - local effects, inhalation	734 mg/m³	
Long-term - systemic effects,oral	4.5 mg/kg bodyweight/day	
Long-term - systemic effects, inhalation	367 mg/m³	
Long-term - systemic effects, dermal	37 mg/kg bodyweight/day	
Long-term - local effects, inhalation	367 mg/m³	
PNEC (Water)		
PNEC aqua (freshwater)	0.24 mg/l	
PNEC aqua (marine water)	0.024 mg/l	
PNEC aqua (intermittent, freshwater)	1.65 mg/l	
PNEC (Sediment)		
PNEC sediment (freshwater)	1.15 mg/kg dwt	
PNEC sediment (marine water)	0.115 mg/kg dwt	
PNEC (Soil)		
PNEC soil	0.148 mg/kg dwt	
PNEC (Oral)		
PNEC oral (secondary poisoning)	0.2 g/kg food	
PNEC (STP)		
PNEC sewage treatment plant	650 mg/l	

## 8.1.5. Control banding

No additional information available

## 8.2. Exposure controls

#### 8.2.1. Appropriate engineering controls

#### Appropriate engineering controls:

Ensure good ventilation of the work station.

#### 8.2.2. Personal protection equipment

## Personal protective equipment symbol(s):







#### 8.2.2.1. Eye and face protection

## Eye protection:

Safety glasses

#### 8.2.2.2. Skin protection

## Skin and body protection:

Wear suitable protective clothing

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#### Hand protection:

Protective gloves

Hand protection					
Туре	Material	Permeation	Thickness (mm)	Penetration	Standard
Disposable gloves	Viton® II	6 (> 480 minutes)	0,7 mm		EN 374-3
Disposable gloves	Nitrile rubber (NBR)	2 (> 30 minutes)	0,4 mm		EN 374-3

#### 8.2.2.3. Respiratory protection

#### Respiratory protection:

In case of insufficient ventilation, wear suitable respiratory equipment

Respiratory protection			
Device	Filter type	Condition	Standard
Gas mask with filter type	Filter A1/B1		EN 14387

#### 8.2.2.4. Thermal hazards

No additional information available

#### 8.2.3. Environmental exposure controls

#### **Environmental exposure controls:**

Avoid release to the environment.

## **SECTION 9: Physical and chemical properties**

## 9.1. Information on basic physical and chemical properties

Physical state : Liquid Colour : Various colours. Odour : characteristic. Odour threshold : 0.9 – 9 mg/m³ Xylene Melting point : Not applicable Freezing point : Not available Boiling point : 126 - 145 °C Flammability : Not applicable Explosive properties : No data available. Explosive limits : Not available Lower explosion limit : 1.1 vol % Xylene Upper explosion limit : 8 vol % Xylene Flash point : 24 °C : 270 - 300 °C Auto-ignition temperature Decomposition temperature : Not available рΗ : Not available Viscosity, kinematic : 5000 mm<sup>2</sup>/s Solubility : Slightly soluble.

Partition coefficient n-octanol/water (Log Kow) : Not available Vapour pressure : Not available Vapour pressure at 50°C : Not available Density : 1.5 g/cm³ Relative density : Not available Relative vapour density at 20°C : Not available Particle characteristics : Not applicable

### 9.2. Other information

#### 9.2.1. Information with regard to physical hazard classes

No additional information available

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#### 9.2.2. Other safety characteristics

No additional information available

## **SECTION 10: Stability and reactivity**

#### 10.1. Reactivity

The product is non-reactive under normal conditions of use, storage and transport.

#### 10.2. Chemical stability

Stable under normal conditions of use.

#### 10.3. Possibility of hazardous reactions

No dangerous reactions known under normal conditions of use.

#### 10.4. Conditions to avoid

Keep away from sources of ignition. Prevent build-up of electrostatic charges (e.g, by grounding). Protect from sunlight. Avoid high temperatures.

#### 10.5. Incompatible materials

No contact with: strong acids, strong bases and strong oxidants.

#### 10.6. Hazardous decomposition products

Under normal conditions of storage and use, hazardous decomposition products should not be produced. Thermal decomposition may produce: Carbon monoxide. Other toxic gases.

#### **SECTION 11: Toxicological information**

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

Acute toxicity (oral) : Not classified (Based on available data, the classification criteria are not met)
Acute toxicity (dermal) : Not classified (Based on available data, the classification criteria are not met)
Acute toxicity (inhalation) : Not classified (Based on available data, the classification criteria are not met)

xylene (1330-20-7)		
LD50 oral rat	3523 mg/kg rat	
LD50 dermal rabbit	12126 mg/kg bodyweight Animal: rabbit, Animal sex: male	
LC50 Inhalation - Rat	27124 mg/l	
n-butyl acetate (123-86-4)		
LD50 oral rat	12.2 ml/kg Source: ECHA	
LC50 Inhalation - Rat (Vapours)	> 4.9 mg/l Source: ECHA	
ethyl acetate (141-78-6)		
LD50 oral rat	11.3 ml/kg Source: ECHA	
LD50 oral	4934 mg/kg bodyweight Animal: rabbit, Guideline: OECD Guideline 401 (Acute Oral Toxicity)	
LD50 dermal rabbit	> 20000 mg/kg bodyweight Animal: rabbit, Animal sex: male	
titanium dioxide; [in powder form containing 1 % or more of particles with aerodynamic diameter ≤ 10 μm] (13463-67-7)		
LC50 Inhalation - Rat (Dust/Mist)	> 6.82 mg/l Source: ECHA	
Skin corrosion/irritation	: Causes skin irritation.	

n-butyl acetate (123-86-4)	
рН	6.2 Temp.: 20 °C Concentration: 5,3 g/L

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Serious eye damage/irritation : Not classified (Based on available data, the classification criteria are not met)  n-butyl acetate (123-86-4)  ph	titanium dioxide; [in powder form containing	1 % or more of particles with aerodynamic diameter ≤ 10 μm] (13463-67-7)
n-butyl acetate (123-86-4) pH   6.2 Temp.: 20 °C Concentration: 5.3 g/L  titanium dioxide; [in powder form containing 1 % or more of particles with aerodynamic diameter ≤ 10 µm] (13463-67-7) pH   7 Source: ECHA  Respiratory or skin sensitisation   Not classified (Based on available data, the classification criteria are not met)  Particles with aerodynamic diameter ≤ 10 µm] (13463-67-7) pH   7 Source: ECHA  Respiratory or skin sensitisation   Not classified (Based on available data, the classification criteria are not met)  Particles with aerodynamic diameter ≤ 10 µm] (13463-67-7)  PH   Not classified (Based on available data, the classification criteria are not met)  Particles with aerodynamic diameter ≤ 10 µm] (13463-67-7)  PH   Not classified (Based on available data, the classification criteria are not met)  Particles with aerodynamic diameter ≤ 10 µm] (13463-67-7)  PH   Not classified (Based on available data, the classification criteria are not met)  Particles with aerodynamic diameter ≤ 10 µm] (13463-67-7)  PH   Not classified (Based on available data, the classification criteria are not met)  PROTESTIGE exposure   May cause drowsiness or dizziness.  PROTESTIGE exposure   May cause drowsiness or dizziness.  PROTESTIGE exposure   Not classified (Based on available data, the classification criteria are not met)  PROTESTIGE exposure   Not classified (Based on available data, the classification criteria are not met)  PROTESTIGE exposure   Not classified (Based on available data, the classification criteria are not met)  PROTESTIGE exposure   Not classified (Based on available data, the classification criteria are not met)  PROTESTIGE exposure   Not classified (Based on available data, the classification criteria are not met)  PROTESTIGE exposure   Not classified (Based on available data, the classification criteria are not met)  PROTESTIGE exposure   Not classified (Based on available data, the classification criteria are not met)  PROTECT 320  Viscosity, kinematic   Not classified (Based on available data,	рН	7 Source: ECHA
titanium dioxide; [in powder form containing 1 % or more of particles with aerodynamic diameter ≤ 10 μm] (13463-67-7) pH 7 Source; ECHA Respiratory or skin sensitisation : Not classified (Based on available data, the classification criteria are not met) Aeronicogenicity : Not classified (Based on available data, the classification criteria are not met) Aeronicogenicity : Not classified (Based on available data, the classification criteria are not met) Aeronicogenicity : Not classified (Based on available data, the classification criteria are not met) Aeronicogenicity : Not classified (Based on available data, the classification criteria are not met) Aeronicover toxicity : Not classified (Based on available data, the classification criteria are not met) Aeronicover : Not classified (Based on available data, the classification criteria are not met) Aeronicover : Not classified (Based on available data, the classification criteria are not met) Aeronicover : Not classified (Based on available data, the classification criteria are not met)  **ToT-single exposure	Serious eye damage/irritation :	Not classified (Based on available data, the classification criteria are not met)
titianium dioxide; [in powder form containing 1 % or more of particles with aerodynamic diameter ≤ 10 μm] (13463-67-7) pH 7 Source; ECHA Respiratory or skin sensitisation 7 Source; ECHA Respiratory or skin sensitisation 8 Not classified (Based on available data, the classification criteria are not met) Per de imutagenicity 8 Not classified (Based on available data, the classification criteria are not met) Per de imutagenicity 9 Source; ECHA Respiratory or skin sensitisation 9 Source; ECHA Respiratory or skin sensitisation 1 Source; ECHA Respiratory or skin sensitisation criteria are not met) Respiratory or skin sensitisation criteria are	n-butyl acetate (123-86-4)	
Prince	рН	6.2 Temp.: 20 °C Concentration: 5,3 g/L
Respiratory or skin sensitisation  In Not classified (Based on available data, the classification criteria are not met)  Sarm cell mutagenicity  Not classified (Based on available data, the classification criteria are not met)  Accreiospenicity  Not classified (Based on available data, the classification criteria are not met)  Itianium dioxide; [in powder form containing 1 % or more of particles with aerodynamic diameter £ 10 µm] (13463-67-7)  IARG group  Be Possibly carcinogenic to humans  Reproductive toxicity  Not classified (Based on available data, the classification criteria are not met)  In-butyl acetate (123-86-4)  STOT-single exposure  May cause drowsiness or dizziness.  Why cause drowsiness or dizziness.  Why cause drowsiness or dizziness.  Why cause drowsiness or dizziness.  In StOT-single exposure  May cause drowsiness or dizziness.  STOT-single exposure  Not classified (Based on available data, the classification criteria are not met)  STOT-single exposure  May cause drowsiness or dizziness.  STOT-single exposure  Not classified (Based on available data, the classification criteria are not met)  STOT-single exposure  Not classified (Based on available data, the classification criteria are not met)  STOT-single exposure  Not classified (Based on available data, the classification criteria are not met)  STOT-single exposure  STOT-single exposure  Not classified (Based on available data, the classification criteria are not met)  STOT-single exposure  STOT-single exposure  Not classified (Based on available data, the classification criteria are not met)  STOT-single exposure  STOT-single exposure  Not classified (Based on available data, the classification criteria are not met)  STOT-single exposure  STOT-single exposure  Not classified (Based on available data, the classification criteria are not met)  STOT-single exposure  STOT-single exposure  STOT-single exposure  STOT-single exposure  Not classified (Based on available data, the classification criteria are not met)  STOT-single exposure  STOT-sing	titanium dioxide; [in powder form containing	1 % or more of particles with aerodynamic diameter ≤ 10 µm] (13463-67-7)
aerm cell mutagenicity : Not classified (Based on available data, the classification criteria are not met) carcinogenicity : Not classified. (Based on available data, the classification criteria are not met) titanium dioxide; [in powder form containing 1 % or more of particles with aerodynamic diameter ≤ 10 μm] (13463-67-7)  IARC group 2B - Possibly carcinogenic to humans  Reproductive toxicity : Not classified (Based on available data, the classification criteria are not met) to classified (Based on available data, the classification criteria are not met)  STOT-single exposure   May cause drowsiness or dizziness.  ethyl acetate (123-86-4)  STOT-single exposure   May cause drowsiness or dizziness.  ethyl acetate (141-78-6)  STOT-single exposure   May cause drowsiness or dizziness.  ethyl acetate (1330-20-7)  LOAEL (oral, rat, 90 days)   150 mg/kg bodyweight Animal: rat, Animal sex: male, Guideline: CPA OPP 82-1 (90-Day Ora Toxicity)  n-butyl acetate (123-86-4)  LOAEL (oral, rat, 90 days)   500 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 798.2650 (90-Day Oral Toxicity in Rodents)  NOAEL (oral, rat, 90 days)   125 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 798.2650 (90-Day Oral Toxicity in Rodents)  NOAEL (oral, rat, 90 days)   3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 798.2650 (90-Day Oral Toxicity in Rodents)  NOAEL (oral, rat, 90 days)   3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 798.2650 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)   3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)   3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)   3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)   3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)	рН	7 Source: ECHA
titanium dioxide; [in powder form containing 1 % or more of particles with aerodynamic diameter ≤ 10 µm] (13463-67-7)  IARC group  28 - Possibly carcinogenic to humans  Reproductive toxicity: Not classified (Based on available data, the classification criteria are not met)  STOT-single exposure: Not classified (Based on available data, the classification criteria are not met)  stoTo-single exposure  May cause drowsiness or dizziness.  ethyl acetate (141-78-6)  STOT-single exposure: May cause drowsiness or dizziness.  ethyl acetate (93-86-9)  STOT-single exposure: Not classified (Based on available data, the classification criteria are not met)  stylene (1330-20-7)  LOAEL (oral, rat, 90 days)  150 mg/kg bodyweight Animal: rat, Animal sex: male, Guideline: OECD Guideline 408 (Repeated Dose 90-Day Oral Toxicity in Rodents), Guideline: EPA OPP 82-1 (90-Day Oral Toxicity)  n-butyl acetate (123-86-4)  LOAEL (oral, rat, 90 days)  500 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 798.2650 (90-Day Oral Toxicity in Rodents)  NOAEL (oral, rat, 90 days)  125 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 798.2650 (90-Day Oral Toxicity in Rodents)  NOAEL (oral, rat, 90 days)  3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 798.2650 (90-Day Oral Toxicity in Rodents)  NOAEL (oral, rat, 90 days)  3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat	Respiratory or skin sensitisation :	Not classified (Based on available data, the classification criteria are not met)
titanium dioxide; [in powder form containing 1 % or more of particles with aerodynamic diameter ≤ 10 µm] (13463-67-7)  IARC group  2B - Possibly carcinogenic to humans  Reproductive toxicity : Not classified (Based on available data, the classification criteria are not met) : Not classified (Based on available data, the classification criteria are not met)  Thoutyl acetate (123-86-4)  STOT-single exposure  May cause drowsiness or dizziness.  STOT-repated exposure  May cause drowsiness or dizziness.  STOT-repated exposure  Not classified (Based on available data, the classification criteria are not met)  xylene (1330-20-7)  LOAEL (oral, rat, 90 days)  150 mg/kg bodyweight Animal: rat, Animal sex: male, Guideline: DECD Guideline 408 (Repeated Dose 90-Day Oral Toxicity in Rodents), Guideline: EPA OPP 82-1 (90-Day Oral Toxicity)  n-butyl acetate (123-86-4)  LOAEL (oral, rat, 90 days)  125 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 798.2650 (90-Day Oral Toxicity in Rodents)  NOAEL (oral, rat, 90 days)  125 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 798.2650 (90-Day Oral Toxicity in Rodents)  Rodents)  Phyl acetate (141-78-6)  LOAEL (oral, rat, 90 days)  3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)	Germ cell mutagenicity :	Not classified (Based on available data, the classification criteria are not met)
ARC group   2B - Possibly carcinogenic to humans	Carcinogenicity :	Not classified. (Based on available data, the classification criteria are not met)
Reproductive toxicity : Not classified (Based on available data, the classification criteria are not met) STOT-single exposure : Not classified (Based on available data, the classification criteria are not met)  Thutyl acetate (123-86-4) STOT-single exposure   May cause drowsiness or dizziness.  May cause drowsiness or dizziness.  May cause drowsiness or dizziness.  STOT-single exposure   May cause drowsiness or dizziness.  STOT-repeated exposure   May cause drowsiness or dizziness.  STOT-repeated exposure   STOT-repeated exposure   Not classified (Based on available data, the classification criteria are not met)  xylene (1330-20-7)  LOAEL (oral, rat, 90 days)   150 mg/kg bodyweight Animal: rat, Animal sex: male, Guideline: DECD Guideline 408 (Repeated Dose 90-Day Oral Toxicity in Rodents), Guideline: EPA OPP 82-1 (90-Day Oral Toxicity)  In-butyl acetate (123-86-4)  LOAEL (oral, rat, 90 days)   500 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 798.2650 (90-Day Oral Toxicity in Rodents)  NOAEL (oral, rat, 90 days)   125 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 798.2650 (90-Day Oral Toxicity in Rodents)  LOAEL (oral, rat, 90 days)   125 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)   900 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)   900 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)   900 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)   900 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)   900 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)   900 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)   900 mg/kg bodyweig	titanium dioxide; [in powder form containing	1 % or more of particles with aerodynamic diameter ≤ 10 μm] (13463-67-7)
STOT-single exposure : Not classified (Based on available data, the classification criteria are not met)  n-butyl acetate (123-86-4)  STOT-single exposure May cause drowsiness or dizziness.  May cause drowsiness or dizziness.  STOT-single exposure May cause drowsiness or dizziness.  STOT-single exposure : Not classified (Based on available data, the classification criteria are not met)  xylene (1330-20-7)  LOAEL (oral, rat, 90 days) 150 mg/kg bodyweight Animal: rat, Animal sex: male, Guideline: OECD Guideline 408 (Repeated Dose 90-Day Oral Toxicity in Rodents), Guideline: EPA OPP 82-1 (90-Day Oral Toxicity)  n-butyl acetate (123-86-4)  LOAEL (oral, rat, 90 days) 500 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 798.2650 (90-Day Oral Toxicity in Rodents)  NOAEL (oral, rat, 90 days) 125 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 798.2650 (90-Day Oral Toxicity in Rodents)  LOAEL (oral, rat, 90 days) 3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days) 3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days) 3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days) 3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days) 5000 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days) 5000 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days) 5000 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days) 5000 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days) 5000 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days) 5000 mg/kg bodyweigh	IARC group	2B - Possibly carcinogenic to humans
n-butyl acetate (123-86-4)  STOT-single exposure  thyl acetate (141-78-6)  STOT-single exposure  May cause drowsiness or dizziness.  May cause drowsiness or dizziness.  STOT-repeated exposure  Not classified (Based on available data, the classification criteria are not met)  xylene (1330-20-7)  LOAEL (oral, rat, 90 days)  150 mg/kg bodyweight Animal: rat, Animal sex: male, Guideline: OECD Guideline 408 (Repeated Dose 90-Day Oral Toxicity in Rodents), Guideline: EPA OPP 82-1 (90-Day Oral Toxicity)  n-butyl acetate (123-86-4)  LOAEL (oral, rat, 90 days)  500 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 798.2650 (90-Day Oral Toxicity in Rodents)  NOAEL (oral, rat, 90 days)  125 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 798.2650 (90-Day Oral Toxicity in Rodents)  LOAEL (oral, rat, 90 days)  3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  5000 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  5000 mm²/s  NOO mm²/s	Reproductive toxicity :	
STOT-single exposure  thyl acetate (141-78-6)  STOT-single exposure  May cause drowsiness or dizziness.  May cause drowsiness or dizziness.  STOT-repeated exposure  Not classified (Based on available data, the classification criteria are not met)  xylene (1330-20-7)  LOAEL (oral, rat, 90 days)  150 mg/kg bodyweight Animal: rat, Animal sex: male, Guideline: OECD Guideline 408 (Repeated Dose 90-Day Oral Toxicity in Rodents), Guideline: EPA OPP 82-1 (90-Day Ora Toxicity)  n-butyl acetate (123-86-4)  LOAEL (oral, rat, 90 days)  500 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 798.2650 (90-Day Oral Toxicity in Rodents)  NOAEL (oral, rat, 90 days)  125 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 798.2650 (90-Day Oral Toxicity in Rodents)  LOAEL (oral, rat, 90 days)  3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  900 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  900 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  900 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  5000 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  5000 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  5000 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  5000 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  5000 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)	STOT-single exposure :	Not classified (Based on available data, the classification criteria are not met)
ethyl acetate (141-78-6)  STOT-single exposure	n-butyl acetate (123-86-4)	
May cause drowsiness or dizziness.  STOT-repeated exposure : Not classified (Based on available data, the classification criteria are not met)  xylene (1330-20-7)  LOAEL (oral, rat, 90 days)   150 mg/kg bodyweight Animal: rat, Animal sex: male, Guideline: OECD Guideline 408 (Repeated Dose 90-Day Oral Toxicity in Rodents), Guideline: EPA OPP 82-1 (90-Day Oral Toxicity)  n-butyl acetate (123-86-4)  LOAEL (oral, rat, 90 days)   500 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 798.2650 (90-Day Oral Toxicity in Rodents)  NOAEL (oral, rat, 90 days)   125 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 798.2650 (90-Day Oral Toxicity in Rodents)  ethyl acetate (141-78-6)  LOAEL (oral, rat, 90 days)   3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)   900 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)   900 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)   900 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)   900 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)   900 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)   900 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)   900 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)	STOT-single exposure	May cause drowsiness or dizziness.
STOT-repeated exposure : Not classified (Based on available data, the classification criteria are not met)  xylene (1330-20-7)  LOAEL (oral, rat, 90 days)	ethyl acetate (141-78-6)	
LOAEL (oral, rat, 90 days)  150 mg/kg bodyweight Animal: rat, Animal sex: male, Guideline: OECD Guideline 408 (Repeated Dose 90-Day Oral Toxicity in Rodents), Guideline: EPA OPP 82-1 (90-Day Ora Toxicity)  n-butyl acetate (123-86-4)  LOAEL (oral, rat, 90 days)  500 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 798.2650 (90-Day Oral Toxicity in Rodents)  NOAEL (oral, rat, 90 days)  125 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 798.2650 (90-Day Oral Toxicity in Rodents)  ethyl acetate (141-78-6)  LOAEL (oral, rat, 90 days)  3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  900 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  Aspiration hazard  : Not classified (Based on available data, the classification criteria are not met)  PROTECT 320  Viscosity, kinematic  5000 mm²/s  5000 mm²/s	STOT-single exposure	May cause drowsiness or dizziness.
LOAEL (oral, rat, 90 days)  150 mg/kg bodyweight Animal: rat, Animal sex: male, Guideline: OECD Guideline 408 (Repeated Dose 90-Day Oral Toxicity in Rodents), Guideline: EPA OPP 82-1 (90-Day Oral Toxicity)  n-butyl acetate (123-86-4)  LOAEL (oral, rat, 90 days)  500 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 798.2650 (90-Day Oral Toxicity in Rodents)  NOAEL (oral, rat, 90 days)  125 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 798.2650 (90-Day Oral Toxicity in Rodents)  ethyl acetate (141-78-6)  LOAEL (oral, rat, 90 days)  3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  900 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  Aspiration hazard  PROTECT 320  Viscosity, kinematic  5000 mm²/s  n-butyl acetate (123-86-4)	STOT-repeated exposure :	Not classified (Based on available data, the classification criteria are not met)
(Repeated Dose 90-Day Oral Toxicity in Rodents), Guideline: EPA OPP 82-1 (90-Day Oral Toxicity)  n-butyl acetate (123-86-4)  LOAEL (oral, rat, 90 days)  NOAEL (oral, rat, 90 days)  125 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 798.2650 (90-Day Oral Toxicity in Rodents)  NOAEL (oral, rat, 90 days)  125 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 798.2650 (90-Day Oral Toxicity in Rodents)  ethyl acetate (141-78-6)  LOAEL (oral, rat, 90 days)  3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  3900 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  3000 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  Not classified (Based on available data, the classification criteria are not met)  PROTECT 320  Viscosity, kinematic  5000 mm²/s  n-butyl acetate (123-86-4)	xylene (1330-20-7)	
LOAEL (oral, rat, 90 days)  500 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 798.2650 (90-Day Oral Toxicity in Rodents)  NOAEL (oral, rat, 90 days)  125 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 798.2650 (90-Day Oral Toxicity in Rodents)  ethyl acetate (141-78-6)  LOAEL (oral, rat, 90 days)  3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  900 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  Aspiration hazard  Not classified (Based on available data, the classification criteria are not met)  PROTECT 320  Viscosity, kinematic  5000 mm²/s  n-butyl acetate (123-86-4)	LOAEL (oral, rat, 90 days)	(Repeated Dose 90-Day Oral Toxicity in Rodents), Guideline: EPA OPP 82-1 (90-Day Oral
Rodents)  NOAEL (oral, rat, 90 days)  125 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 798.2650 (90-Day Oral Toxicity in Rodents)  ethyl acetate (141-78-6)  LOAEL (oral, rat, 90 days)  3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  900 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  Aspiration hazard  : Not classified (Based on available data, the classification criteria are not met)  PROTECT 320  Viscosity, kinematic  5000 mm²/s  n-butyl acetate (123-86-4)	n-butyl acetate (123-86-4)	
ethyl acetate (141-78-6)  LOAEL (oral, rat, 90 days)  NOAEL (oral, rat, 90 days)  Sepiration hazard  PROTECT 320  Viscosity, kinematic  Rodents)  Aspiration Loae (123-86-4)  Rodents)  Rodents)  Aspiration Loae (141-78-6)  Rodents)  Aspiration Mayor (141-78-6)  Sepiration Mayor	LOAEL (oral, rat, 90 days)	500 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 798.2650 (90-Day Oral Toxicity in Rodents)
LOAEL (oral, rat, 90 days)  3600 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  NOAEL (oral, rat, 90 days)  900 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  Aspiration hazard  : Not classified (Based on available data, the classification criteria are not met)  PROTECT 320  Viscosity, kinematic  5000 mm²/s  n-butyl acetate (123-86-4)	NOAEL (oral, rat, 90 days)	125 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 798.2650 (90-Day Oral Toxicity in Rodents)
Toxicity Test)  NOAEL (oral, rat, 90 days)  900 mg/kg bodyweight Animal: rat, Guideline: EPA OTS 795.2600 (Subchronic Oral Toxicity Test)  Aspiration hazard  : Not classified (Based on available data, the classification criteria are not met)  PROTECT 320  Viscosity, kinematic  5000 mm²/s  n-butyl acetate (123-86-4)	ethyl acetate (141-78-6)	
Toxicity Test) Aspiration hazard : Not classified (Based on available data, the classification criteria are not met)  PROTECT 320  Viscosity, kinematic 5000 mm²/s  n-butyl acetate (123-86-4)	LOAEL (oral, rat, 90 days)	,
PROTECT 320 Viscosity, kinematic 5000 mm²/s n-butyl acetate (123-86-4)	NOAEL (oral, rat, 90 days)	
Viscosity, kinematic 5000 mm²/s n-butyl acetate (123-86-4)	Aspiration hazard :	Not classified (Based on available data, the classification criteria are not met)
n-butyl acetate (123-86-4)	PROTECT 320	
	Viscosity, kinematic	5000 mm²/s
Viscosity, kinematic 0.83 mm²/s Temp.: '20°C' Parameter: 'kinematic viscosity (in mm²/s)'	n-butyl acetate (123-86-4)	
	Viscosity, kinematic	0.83 mm²/s Temp.: '20°C' Parameter: 'kinematic viscosity (in mm²/s)'

## 11.2. Information on other hazards

No additional information available

1/2/2023 (Revision date) GB - en 10/15

## Safety Data Sheet

SDS EU format according to COMMISSION REGULATION (EU) 2020/878

## **SECTION 12: Ecological information**

## 12.1. Toxicity

Hazardous to the aquatic environment, short-term

(acute)

(chronic)

Hazardous to the aquatic environment, long-term

: Not classified (Based on available data, the classification criteria are not met)

: Not classified (Based on available data, the classification criteria are not met)

Not rapidly degradable

xylene (1330-20-7)		
LC50 - Fish [1]	2.6 mg/l Test organisms (species): Oncorhynchus mykiss (previous name: Salmo gairdneri)	
EC50 - Crustacea [1]	> 3.4 mg/l Test organisms (species): Ceriodaphnia dubia	
NOEC chronic fish	> 1.3 mg/l Test organisms (species): Oncorhynchus mykiss (previous name: Salmo gairdneri) Duration: '56 d'	
n-butyl acetate (123-86-4)		
LC50 - Fish [1]	18 mg/l Source: ECHA	
EC50 - Crustacea [1]	44 mg/l Source: ECHA	
EC50 - Other aquatic organisms [1]	32 mg/l Test organisms (species): Artemia salina	
EC50 72h - Algae [1]	674.7 mg/l Test organisms (species): Desmodesmus subspicatus (previous name: Scenedesmus subspicatus)	
EC50 72h - Algae [2]	246 mg/l Test organisms (species): Pseudokirchneriella subcapitata (previous names: Raphidocelis subcapitata, Selenastrum capricornutum)	
LOEC (chronic)	47.6 mg/l Test organisms (species): Daphnia magna Duration: '21 d'	
NOEC (chronic)	23.2 mg/l Test organisms (species): Daphnia magna Duration: '21 d'	
ethyl acetate (141-78-6)		
LC50 - Fish [1]	230 mg/l Source: ECHA	
NOEC (chronic)	2.4 mg/l Test organisms (species): Daphnia magna Duration: '21 d'	
titanium dioxide; [in powder form con	taining 1 % or more of particles with aerodynamic diameter ≤ 10 μm] (13463-67-7)	
LC50 - Fish [1]	> 100 mg/l	
EC50 72h - Algae [1]	> 50 mg/l Source: ECHA	

#### 12.2. Persistence and degradability

No additional information available

## 12.3. Bioaccumulative potential

n-butyl acetate (123-86-4)		
Partition coefficient n-octanol/water (Log Pow)	1.78 Source: HSDB	
ethyl acetate (141-78-6)		
Partition coefficient n-octanol/water (Log Pow)	0.73 Source: ICSC	

#### 12.4. Mobility in soil

No additional information available

#### 12.5. Results of PBT and vPvB assessment

No additional information available

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#### 12.6. Endocrine disrupting properties

No additional information available

#### 12.7. Other adverse effects

No additional information available

### **SECTION 13: Disposal considerations**

#### 13.1. Waste treatment methods

Regional legislation (waste)

Waste treatment methods

Sewage disposal recommendations

Product/Packaging disposal recommendations

Additional information

European List of Waste (LoW) code

- : Disposal must be done according to official regulations.
- : Dispose of contents/container in accordance with licensed collector's sorting instructions.
- : Do not discharge into drains.
- : This material and its container must be disposed of as hazardous waste. Do not dispose of with domestic waste. After cleaning, recycle or dispose of at an authorised site.
- : Flammable vapours may accumulate in the container.
- : 08 01 11\* waste paint and varnish containing organic solvents or other dangerous

substances

15 01 10\* - packaging containing residues of or contaminated by dangerous substances

#### **SECTION 14: Transport information**

In accordance with ADR / IMDG / IATA

ADR	IMDG	IATA
14.1. UN number or ID number		
UN 1263	UN 1263	UN 1263
14.2. UN proper shipping name		
PAINT	PAINT	Paint
Transport document description		
UN 1263 PAINT, 3, III, (D/E)	UN 1263 PAINT, 3, III (24°C c.c.)	UN 1263 Paint, 3, III
14.3. Transport hazard class(es)		
3	3	3
3	3	3
14.4. Packing group		
III	III	III
14.5. Environmental hazards		•
Dangerous for the environment: No	Dangerous for the environment: No Marine pollutant: No	Dangerous for the environment: No
No supplementary information available		,

#### 14.6. Special precautions for user

## Overland transport

Classification code (ADR) : F1
Limited quantities (ADR) : 5I
Special packing provisions (ADR) : PP1
Mixed packing provisions (ADR) : MP19
Transport category (ADR) : 3

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Special provisions for carriage - Packages (ADR) : V12

Tunnel restriction code (ADR) : D/E EAC code : •3Y

Transport by sea

Special provisions (IMDG) : 163, 223, 367, 955

Limited quantities (IMDG) : 5 L
Special packing provisions (IMDG) : PP1
EmS-No. (Fire) : F-E
EmS-No. (Spillage) : S-E
Stowage category (IMDG) : A

#### Air transport

No data available

#### 14.7. Maritime transport in bulk according to IMO instruments

Not applicable

#### **SECTION 15: Regulatory information**

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

#### 15.1.1. EU-Regulations

#### **REACH Annex XVII (Restriction List)**

Contains no substance(s) listed on REACH Annex XVII (Restriction Conditions)

#### REACH Annex XIV (Authorisation List)

Contains no substance(s) listed on REACH Annex XIV (Authorisation List)

#### **REACH Candidate List (SVHC)**

Contains no substance(s) listed on the REACH Candidate List

#### PIC Regulation (Prior Informed Consent)

Contains no substance(s) listed on the PIC list (Regulation EU 649/2012 concerning the export and import of hazardous chemicals)

#### **POP Regulation (Persistent Organic Pollutants)**

Contains no substance(s) listed on the POP list (Regulation EU 2019/1021 on persistent organic pollutants)

#### Ozone Regulation (1005/2009)

Contains no substance(s) listed on the Ozone Depletion list (Regulation EU 1005/2009 on substances that deplete the ozone layer)

#### **Explosives Precursors Regulation (2019/1148)**

Contains no substance(s) listed on the Explosives Precursors list (Regulation EU 2019/1148 on the marketing and use of explosives precursors)

#### **Drug Precursors Regulation (273/2004)**

Contains no substance(s) listed on the Drug Precursors list (Regulation EC 273/2004 on the manufacture and the placing on market of certain substances used in the illicit manufacture of narcotic drugs and psychotropic substances)

#### 15.1.2. National regulations

No additional information available

#### 15.2. Chemical safety assessment

No chemical safety assessment has been carried out

#### **SECTION 16: Other information**

#### Indication of changes:

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Abbreviations and acronyms:		
ADN	European Agreement concerning the International Carriage of Dangerous Goods by Inland Waterways	
ADR	European Agreement concerning the International Carriage of Dangerous Goods by Road	
ATE	Acute Toxicity Estimate	
BCF	Bioconcentration factor	
BLV	Biological limit value	
BOD	Biochemical oxygen demand (BOD)	
COD	Chemical oxygen demand (COD)	
DMEL	Derived Minimal Effect level	
DNEL	Derived-No Effect Level	
EC-No.	European Community number	
EC50	Median effective concentration	
EN	European Standard	
IARC	International Agency for Research on Cancer	
IATA	International Air Transport Association	
IMDG	International Maritime Dangerous Goods	
LC50	Median lethal concentration	
LD50	Median lethal dose	
LOAEL	Lowest Observed Adverse Effect Level	
NOAEC	No-Observed Adverse Effect Concentration	
NOAEL	No-Observed Adverse Effect Level	
NOEC	No-Observed Effect Concentration	
OECD	Organisation for Economic Co-operation and Development	
OEL	Occupational Exposure Limit	
PBT	Persistent Bioaccumulative Toxic	
PNEC	Predicted No-Effect Concentration	
RID	Regulations concerning the International Carriage of Dangerous Goods by Rail	
SDS	Safety Data Sheet	
STP	Sewage treatment plant	
ThOD	Theoretical oxygen demand (ThOD)	
TLM	Median Tolerance Limit	
VOC	Volatile Organic Compounds	
CAS-No.	Chemical Abstract Service number	
N.O.S.	Not Otherwise Specified	
vPvB	Very Persistent and Very Bioaccumulative	
ED	Endocrine disrupting properties	

Data sources : ECHA (European Chemicals Agency).

Training advice : Handle in accordance with good industrial hygiene and safety procedures.

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Full text of H- and EUH-statements:		
Acute Tox. 4 (Dermal)	Acute toxicity (dermal), Category 4	
Acute Tox. 4 (Inhalation)	Acute toxicity (inhal.), Category 4	
Carc. 2	Carcinogenicity, Category 2	
EUH211	Warning! Hazardous respirable droplets may be formed when sprayed. Do not breathe spray or mist.	
Eye Irrit. 2	Serious eye damage/eye irritation, Category 2	
Flam. Liq. 2	Flammable liquids, Category 2	
Flam. Liq. 3	Flammable liquids, Category 3	
H225	Highly flammable liquid and vapour.	
H226	Flammable liquid and vapour.	
H312	Harmful in contact with skin.	
H315	Causes skin irritation.	
H319	Causes serious eye irritation.	
H332	Harmful if inhaled.	
H336	May cause drowsiness or dizziness.	
H351	Suspected of causing cancer.	
Skin Irrit. 2	Skin corrosion/irritation, Category 2	
STOT SE 3	Specific target organ toxicity – Single exposure, Category 3, Narcosis	

Classification and procedure used to derive the classification for mixtures according to Regulation (EC) 1272/2008 [CLP]:				
Flam. Liq. 3	H226	On basis of test data		
Skin Irrit. 2	H315	Calculation method		

Safety Data Sheet (SDS), EU

This information is based on our current knowledge and is intended to describe the product for the purposes of health, safety and environmental requirements only. It should not therefore be construed as guaranteeing any specific property of the product.