



CONSEIL DE L'EUROPI

Bezafibrate

Safety Data Sheet

according to the REACH Regulation (EC) 1907/2006 amended by Regulation (EU) 2020/878 Issue date: 03/04/2023 Version: 1.0

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

1.1. Product identifier Product form : Substance Substance name : Bezafibrate : 201600105 Product code : Bezafibrate, salts, hydrates, isomers and impurities where applicable Synonyms 1.2. Relevant identified uses of the substance or mixture and uses advised against 1.2.1. Relevant identified uses : The product is intended for research, analysis and scientific education. Main use category Use of the substance/mixture : For professional use only Function or use category : Laboratory chemicals 1.2.2. Uses advised against Restrictions on use : Do not use : Ingestion, Inhalation, Dermal 1.3. Details of the supplier of the safety data sheet European Directorate for the Quality of Medicines & Healthcare EDQM, Council of Europe 7, Allée Kastner, CS30026 F- 67081 Strasbourg France T +33(0)388412035 - F +33(0)388412771 sds@edqm.eu - www.edqm.eu 1.4. Emergency telephone number Emergency number : +33(0)390215608 **SECTION 2: Hazards identification** 2.1. Classification of the substance or mixture Classification according to Regulation (EC) No. 1272/2008 [CLP] H302 Acute toxicity (oral), Category 4 Full text of H- and EUH-statements: see section 16 Adverse physicochemical, human health and environmental effects Expert judgement and weight of evidence determination. 2.2. Label elements Labelling according to Regulation (EC) No. 1272/2008 [CLP] Hazard pictograms (CLP) GHS07 Signal word (CLP) : Warning Hazard statements (CLP) : H302 - Harmful if swallowed. Precautionary statements (CLP) : P301+P312 - IF SWALLOWED: Call a doctor if you feel unwell. Labelling according to: exemption for inner packaging where the contents do not exceed 10ml No labelling required

2.3. Other hazards

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SECTION 3: Composition/information on ingredients					
3.1. Substances					
Name	Product identifier	%	Classification according to Regulation (EC) No. 1272/2008 [CLP]		
Bezafibrate	-	≤ 100	Acute Tox. 4 (Oral), H302 (ATE=300 mg/kg bodyweight)		
Full text of H- and EUH-statements: see section 16		-			
3.2. Mixtures					
Not applicable					
SECTION 4: First aid measures					
4.1. Description of first aid measures					
First-aid measures after skin contact First-aid measures after eye contact					
4.2. Most important symptoms and effects	both acute and delayed				
No additional information available					
4.3. Indication of any immediate medical attention and special treatment needed					
No additional information available					
SECTION 5: Firefighting measures					
5.1. Extinguishing media					
Suitable extinguishing media	: Extinguishing blanket.				
5.2. Special hazards arising from the subst	ance or mixture				
Fire hazard	: See Heading 2.2.				
5.3. Advice for firefighters					
Firefighting instructions	: Use extinguishing media appro	opriate for surrou	unding fire.		
SECTION 6: Accidental release measures					
6.1. Personal precautions, protective equipment and emergency procedures					

6.1.1. For non-emergency personnel

: Avoid all unnecessary exposure.

Emergency procedures

6.1.2. For emergency responders

No additional information available

6.2. Environmental precautions

No additional information available

6.3. Methods and material for containment and cleaning up

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6.4. Reference to other sections

No additional information available

SECTION 7: Handling and storage

7.1. Precautions for safe handling

No additional information available

7.2. Conditions for safe storage, including any incompatibilities

No additional information available

7.3. Specific end use(s)

See Heading 1.

SECTION 8: Exposure controls/personal protection

8.1. Control parameters

8.1.1 National occupational exposure and biological limit values

No additional information available

8.1.2. Recommended monitoring procedures

No additional information available

8.1.3. Air contaminants formed

No additional information available

8.1.4. DNEL and PNEC

No additional information available

8.1.5. Control banding

No additional information available

8.2. Exposure controls

8.2.1. Appropriate engineering controls

Appropriate engineering controls:

Keep in a well-ventilated room. Both local exhaust and general room ventilation are usually required.

8.2.2. Personal protection equipment

8.2.2.1. Eye and face protection

No additional information available

8.2.2.2. Skin protection

Skin and body protection: Lab coat

Hand protection:

Protective gloves

8.2.2.3. Respiratory protection

No additional information available

8.2.2.4. Thermal hazards

No additional information available

8.2.3. Environmental exposure controls

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SECTION 9: Physical and chemical properties			
9.1. Information on basic physical and o	chemical properties		
Physical state Colour Appearance Odour Odour threshold	: Solid : Not available : Powder. : Not available : Not available		
Melting point Freezing point Boiling point Flammability Explosive limits	: Not available : Not available : Not available : Not available : Not applicable		
Lower explosion limit Upper explosion limit Flash point Auto-ignition temperature	Not applicableNot applicableNot applicableNot applicableNot applicable		
Decomposition temperature pH pH solution Viscosity, kinematic Solubility	 Not available Not available Not available Not applicable Water: < 0,1 g/l Not available 		
Partition coefficient n-octanol/water (Log Kow) Vapour pressure Vapour pressure at 50°C Density Relative density Relative vapour density at 20°C Particle size	 Not available Not available Not available Not available Not available Not available Not applicable Not available 		

9.2. Other information

9.2.1. Information with regard to physical hazard classes

No additional information available

9.2.2. Other safety characteristics No additional information available

SECTION 10: Stability and reactivity

10.1. Reactivity

Stable under normal conditions.

10.2. Chemical stability

No additional information available

10.3. Possibility of hazardous reactions

None known.

10.4. Conditions to avoid

No additional information available

10.5. Incompatible materials

None under normal use. See Section 7.

10.6. Hazardous decomposition products

When heated to decomposition, emits dangerous fumes.

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11.1. Information on hazard classes as defined in Regulation (EC) No 1272/2008			
Acute toxicity (oral)	: Harmful if swallowed.		
Acute toxicity (dermal)	: Not classified		
Acute toxicity (inhalation)	: Not classified		
Bezafibrate			
LD50 oral rat	300 – 2000 mg/kg		
Skin corrosion/irritation	: Not classified (Lack of data)		
Serious eye damage/irritation	: Not classified (Lack of data)		
Respiratory or skin sensitisation	: Not classified (Based on available data, the classification criteria are not met)		
Germ cell mutagenicity	: Not classified (Lack of data)		
Carcinogenicity	: Not classified (Lack of data)		
Reproductive toxicity	: Not classified (Further details: Acute toxicity)		
STOT-single exposure	: Not classified (Further details: Acute toxicity)		
STOT-repeated exposure	: Not classified (Further details: Acute toxicity)		
Aspiration hazard	: Not classified (Based on available data, the classification criteria are not met)		

No additional information available

SECTION 12: Ecological information 12.1. Toxicity : Presents no specific risk for the environment. Ecology - general Hazardous to the aquatic environment, short-term : Not classified (acute) : Not classified Hazardous to the aquatic environment, long-term (chronic) 12.2. Persistence and degradability No additional information available 12.3. Bioaccumulative potential No additional information available 12.4. Mobility in soil No additional information available 12.5. Results of PBT and vPvB assessment No additional information available 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

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SECTION 14: Transport information In accordance with ADR / IMDG / IATA ADR IMDG ΙΑΤΑ 14.1. UN number or ID number Not applicable Not applicable Not applicable 14.2. UN proper shipping name Not applicable Not applicable Not applicable 14.3. Transport hazard class(es) Not applicable Not applicable Not applicable 14.4. Packing group Not applicable Not applicable Not applicable 14.5. Environmental hazards Not applicable Not applicable Not applicable No supplementary information available

14.6. Special precautions for user

Overland transport

Not applicable

Transport by sea

Not applicable

Air transport

Not applicable

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)

Not listed on REACH Annex XVII

REACH Annex XIV (Authorisation List)

Not listed on REACH Annex XIV (Authorisation List)

REACH Candidate List (SVHC)

Not listed on the REACH Candidate List

PIC Regulation (Prior Informed Consent)

Not applicable.

POP Regulation (Persistent Organic Pollutants)

Not applicable.

Ozone Regulation (1005/2009)

Not listed on the Ozone Depletion list (Regulation EU 1005/2009)

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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 additional information available

SECTION 16: Other information

Full text of H- and EUH-statements:		
Acute Tox. 4 (Oral)	Acute toxicity (oral), Category 4	
H302	Harmful if swallowed.	

Safety Data Sheet (SDS), EU

DISCLAIMER OF LIABILITY The EDQM has created this SDS as a downstream user for regulatory compliance to rules applicable to chemicals only. This article is intended only for small-volume laboratory analysis and other routine testing prescribed in the pharmacopoeia or EDQM study protocol, under controlled conditions and by professionals only. Any other use of this article or the SDS information is the sole responsibility of the user. This substance is present in articles in quantities totalling under 10 kg per year. There is no human or environmental exposure under intended and foreseeable conditions of use. The user has the responsibility for handling, storage, use conditions and disposal of this article and for any use of the information in this SDS. The information has been obtained from suppliers and is without any warranty, express or implied, regarding its correctness. For this and other reasons, we do not assume responsibility and expressly disclaim liability for loss, damage or expense arising out of or in any way connected with the handling, storage, use or disposal of the article.