

Fetal Bovine Serum

According to Regulation (EC) No. 1907/2006 (REACH)Date of compilation04.10.2017Revision Date01.10.2024Version007/10.2024Replaces version of28.08.2023 (Version 006/08.2023)

Section 1 Identification of the substance/mixture and of the company/undertaking

Product identifiers

Product name:

Fetal Bovine Serum

Catalogue No. & Product name:

~	110.011000000	
	10-FBS-11F	– Fetal Bovine Serum Advanced (FBS Minis), Collected in South America
	FBS-11A	– Fetal Bovine Serum Advanced (FBS Advanced), Collected in South America
	FBS-11B	– Fetal Bovine Serum Advanced (FBS Advanced), Collected in South America
	FBS-HI-11A	- Fetal Bovine Serum Advanced (FBS Advanced), Heat Inactivated (HI), Collected in South America
	FBS-HI-11B	- Fetal Bovine Serum Advanced (FBS Advanced), Heat Inactivated (HI), Collected in South America
	10-FBS-12F	– Fetal Bovine Serum (FBS Minis), Collected in South America
	FBS-12A	– Fetal Bovine Serum (FBS), Collected in South America
	FBS-12B	– Fetal Bovine Serum (FBS), Collected in South America
	FBS-CS-12A	– Fetal Bovine Serum (FBS), Charcoal Stripped, Collected in South America
	FBS-CS-12B	– Fetal Bovine Serum (FBS), Charcoal Stripped, Collected in South America
	FBS-DIA-12A	- Fetal Bovine Serum (FBS), Dialyzed, Collected in South America
	FBS-DIA-12B	– Fetal Bovine Serum (FBS), Dialyzed, Collected in South America
	FBS-ED-12B	– Fetal Bovine Serum (FBS), Exosome Depleted, Collected in South America
	FBS-ED-12F	– Fetal Bovine Serum (FBS), Exosome Depleted, Collected in South America
	FBS-ED-12G	- Fetal Bovine Serum (FBS), Exosome Depleted, Collected in South America
	FBS-ES-12A	- Fetal Bovine Serum, Embryonic Stem Cell Pre-tested (ES Cell FBS), Collected in South America
	FBS-ES-12B	- Fetal Bovine Serum, Embryonic Stem Cell Pre-tested (ES Cell FBS), Collected in South America
	FBS-GI-12A	– Fetal Bovine Serum (FBS), Gamma Irradiated, Collected in South America
	FBS-GI-12B	– Fetal Bovine Serum (FBS), Gamma Irradiated, Collected in South America
	FBS-HI-12A	 Fetal Bovine Serum (FBS), Heat Inactivated, Collected in South America
	FBS-HI-12B	 Fetal Bovine Serum (FBS), Heat Inactivated, Collected in South America
	FBS-IG-12A	 Fetal Bovine Serum (FBS), IgG Depleted, Collected in South America
	FBS-IG-12B	– Fetal Bovine Serum (FBS), IgG Depleted, Collected in South America
	FBS-LE-12A	– Fetal Bovine Serum (FBS), Low Endotoxin, Collected in South America
	FBS-LE-12B	 Fetal Bovine Serum (FBS), Low Endotoxin, Collected in South America
	FBS-TET-12A	 Fetal Bovine Serum (FBS), Tetracycline Negative, Collected in South America
	FBS-TET-12B	 Fetal Bovine Serum (FBS), Tetracycline Negative, Collected in South America
	FBS-22A	– Fetal Bovine Serum (FBS), Origin USA
	FBS-22B	– Fetal Bovine Serum (FBS) Origin USA
	FBS-HI-22A	– Fetal Bovine Serum (FBS) Heat Inactivated, Origin USA
	FBS-HI-22B	– Fetal Bovine Serum (FBS) Heat Inactivated, Origin USA
	FBS-24A	 Fetal Bovine Serum (FBS) BioProcess, 9CFR Tested, Origin USA
	FBS-24B	 Fetal Bovine Serum (FBS) BioProcess, 9CFR Tested, Origin USA

REACH No.:

A registration number is not available for this substance as the substance or its uses are exempted from registration, the annual tonnage does not require a registration, or the registration is envisaged for a later registration deadline.



Fetal Bovine Serum

According to Regulation (EC) No. 1907/2006 (REACH)Date of compilation04.10.2017Revision Date01.10.2024Version007/10.2024Replaces version of28.08.2023 (Version 006/08.2023)

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

Identified uses: For research use only. Not intended for human or animal diagnostic or therapeutic uses.

Details of the supplier of the safety data sheet

Company:	Capricorn Scientific GmbH
	Auf der Lette 13 A
	35085 Ebsdorfergrund
	Germany
Phone:	+49 6424 94464 0
E-Mail:	info@capricorn-scientific.com
Website:	www.capricorn-scientific.com

Emergency telephone number

+49 6424 94464 0 (on workdays, Mon. to Fri. from 8:00 – 16:00 h CET/CEST)

Classification of the substance or mixture

Not a hazardous substance or mixture according to Regulation (EC) No. 1272/2008.

Label elements

Emergency Phone:

No hazard pictogram, no signal word, no hazard statement(s), no precautionary statement(s) required according to Regulation (EC) No 1272/2008.

Other hazards

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

Ecological information:

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

Toxicological information:

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



Fetal Bovine Serum

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

 Date of compilation
 04.10.2017

 Revision Date
 01.10.2024

 Version
 007/10.2024

 Replaces version of
 28.08.2023 (Version 006/08.2023)

Section 3	Composition/information on ingredients	
Mixture		
Synonyms:	FBS	
No components need to be disclosed according to the applicable regulations.		

Section 4	First aid measures	
Description of first aid measures		
General notes	Show this material safety data sheet to the doctor in attendance.	
Following skin contact	Wash off with soap and plenty of water. In case of modifications on the skin, seek medical advice with this safety data sheet.	
Following eye contact	Remove contact lenses. Rinse immediately carefully and thoroughly with eyebath or water. If eye irritation persists: Get medical advice/attention.	
Following ingestion	Make victim drink water (two glasses at most). Consult doctor if feeling unwell. Never give anything by mouth to an unconscious person. Do not induce vomiting without medical advice.	
Following inhalation	If breathed in, move person into fresh air. If not breathing, give artificial respiration.	
Following swallowing	Make victim drink water (two glasses at most). Consult doctor if feeling unwell. Never give anything by mouth to an unconscious person.	
Self-protection of the first aider	Consider personal protective equipment as indicated in section 8.	
Most important symptoms and eff The most important known sympto	ects, both acute and delayed oms and effects are described in the labelling (see section 2) and/or in section 11.	
Indication of any immediate medical attention and special treatment needed		

Indication of any immediate medical attention and special treatment needed No data available.

Section 5	Firefighting measures
Extinguishing media Suitable extinguishing media Unsuitable extinguishing media	Foam Carbon dioxide (CO2) Dry powder For this substance/mixture no limitations of extinguishing agents are given.



Fetal Bovine Serum

According to Regulation (EC) No. 1907/2006 (REACH)Date of compilation04.10.2017Revision Date01.10.2024Version007/10.2024Replaces version of28.08.2023 (Version 006/08.2023)

Special hazards arising from the substance or mixture

Hazardous combustion products:	Nature of decomposition products not known. Combustible. Development of hazardous combustion gases possible in the event of fire.
Advice for firefighters	In the event of fire, wear self-contained breathing apparatus. Suppress (knock down) gases/vapors/mists with a water spray jet. Prevent fire extinguishing water from contaminating surface water or the ground water system.

Section 6 Accidental release measures

Personal precautions, protective equipment and emergency procedures

Advice for non-emergency personnel: Do not breathe vapors, aerosols. Evacuate the danger area, observe emergency procedures, consult an expert. For personal protection see section 8.

Environmental precautions Do not let product enter drains.

Methods and material for containment and cleaning up

Cover drains. Collect, bind, and pump off spills. Observe possible material restrictions (see sections 7 and 10). Take up with liquid-absorbent material. Dispose of properly. Clean up affected area.

Reference to other sections For disposal see section 13.

Section 7	Handling and storage		
Precautions for safe handling	For precautions see section	0.2.	
Conditions for safe storage, including any incompatibilities			
Storage	<u>Conditions</u> : Tightly closed. <u>Storage class</u> (TRGS 510):	<u>Stability</u> : Recommended storage temperature: ≤ -20 °C. 10 - Combustible liquids	
Specific end use(s)	Apart from the uses mentic	oned in section 1 no other specific uses are stipulated.	



Fetal Bovine Serum

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

 Date of compilation
 04.10.2017

 Revision Date
 01.10.2024

 Version
 007/10.2024

 Replaces version of
 28.08.2023 (Version 006/08.2023)

Section 8	Exposure controls/personal protection	
Control parameters	Components with workplace control parameters.	
Exposure controls		
Personal protective equipment		
Respiratory protection	Not required; except in case of aerosol formation.	
Eye protection	Safety glasses tested and approved under appropriate government standards such as NIOSH (US) or EN166 (EU).	
Environmental exposure controls Do not let product enter drains.		

Section 9

Physical and chemical properties

Information on basic physical and chemical properties

Physical state	dark amber liquid
Colour	No data available
Odour (Treshold)	No data available
Melting/freezing point	No data available
Initial boiling point and range	No data available
Flammability (solid, gas)	No data available
Upper/lower explosion limit	No data available
Flash point	No data available
Autoignition temperature	No data available
Decomposition temperature	No data available
рН	6,8 - 8,2
Viscosity - Kinematic:	No data available
- Dynamic:	No data available
Water solubility	No data available
Partition coefficient [n-octanol/water]	No data available
Vapour pressure	No data available
Density/Relative density	No data available
Relative vapour density	No data available
Particle characteristics	No data available
Oxidizing properties	none

Other information

No data available



Fetal Bovine Serum

According to Regulation (EC) No. 1907/2006 (REACH)Date of compilation04.10.2017Revision Date01.10.2024Version007/10.2024Replaces version of28.08.2023 (Version 006/08.2023)

Section 10	ability and Reactivity	
Reactivity	No data available	
Chemical stability	The product is chemically stable under recommended conditions.	
Possibility of hazardous reactions	No data available	
Conditions to avoid	Heat	
Incompatible materials	Strong oxidizing agents	
Hazardous decomposition produc	In the event of fire: see section 5	

Section 11	Toxicological information
------------	---------------------------

Information on toxicological effects

Acute toxicity	Oral: No data available Inhalation: No data available Dermal: No data available
Skin corrosion/irritation	No data available
Serious eye damage/eye irritation	No data available
Respiratory or skin sensitization	No data available
Germ cell mutagenicity	No data available
Carcinogenicity	No data available
Reproductive toxicity	No data available
Specific target organ toxicity	
- single exposure	No data available
- repeated exposure	No data available
Aspiration hazard	No data available

Additional Information

Endocrine disrupting properties

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

To the best of our knowledge, the chemical, physical, and toxicological properties have not been thoroughly investigated.



Fetal Bovine Serum

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

 Date of compilation
 04.10.2017

 Revision Date
 01.10.2024

 Version
 007/10.2024

 Replaces version of
 28.08.2023 (Version 006/08.2023)

Section 12	Ecolog	gical information	
Toxicity	No data	No data available	
Persistence and degradability	No data available		
Bioaccumulative potential	No data available		
Mobility in soil	No data available		
Results of PBT and vPvB assessn	This substance/mixture contains no components considered to be either persistent, bioaccumulative and toxic (PBT), or very persistent and very bioaccumulative (vPvB) at levels of 0.1% or higher.		
Endocrine disrupting properties			
Product:			
	CH Article	e does not contain components considered to have endocrine disrupting e 57(f) or Commission Delegated regulation (EU) 2017/2100 or Commission 0.1% or higher.	
Other adverse effects		No data available	
Section 13	Disposal considerations		
Waste treatment methods			
Product:	Discour	age sewage disposal. Dispose of waste according to applicable legislation.	
Contaminated packaging	Dispose of as unused product. Dispose of waste according to applicable legislation.		



Fetal Bovine Serum

According to Regulation (EC) No. 1907/2006 (REACH)Date of compilation04.10.2017Revision Date01.10.2024Version007/10.2024Replaces version of28.08.2023 (Version 006/08.2023)

Section 14

Transport Information

In accordance with ADR / RID / IMDG / IATA

ADR/RID	IMDG	IATA		
UN number				
-	-	-		
UN proper shipping name				
Not dangerous goods	Not dangerous goods	Not dangerous goods		
Transport hazard classes				
-	-	-		
Packaging group				
-	-	-		
Environmental hazards				
no	Marine pollutant: no	no		

Special precautions for user

No data available

Maritime transport in bulk according to IMO instrumentsNo data available

Section 15

Regulatory information

Safety, health and environmental regulations/legislation specific for the substance or mixture This safety datasheet complies with the requirements of Regulation (EC) No. 1907/2006.

Chemical Safety Assessment

For this product a chemical safety assessment was not carried out.



Fetal Bovine Serum

According to Regulation (EC) No. 1907/2006 (REACH)Date of compilation04.10.2017Revision Date01.10.2024Version007/10.2024Replaces version of28.08.2023 (Version 006/08.2023)

Section 16 Other information

The above information is believed to be correct but does not purport to be all inclusive and shall be used only as a guide. The information in this document is based on the present state of our knowledge and is applicable to the product regarding ap-propriate safety precautions. It does not represent any guarantee of the properties of the product.

Full text of other abbreviations

ADN - European Agreement concerning the International Carriage of Dangerous Goods by Inland Waterways;

ADR - Agreement concerning the International Carriage of Dangerous Goods by Road;

AllC - Australian Inventory of Industrial Chemicals; ASTM - American Society for the Testing of Materials; bw - Body weight;

CMR - Carcinogen, Mutagen or Reproductive Toxicant; DIN - Standard of the German Institute for Standardisation;

DSL - Domestic Substances List (Canada); ECx – Concentration associated with x% response; ELx - Loading rate associated with x% response; EmS - Emergency Schedule; ENCS - Existing & New Chemical Substances (Japan);

ErCx - Concentration associated with x% growth rate response; GHS - Globally Harmonized System; GLP - Good Laboratory Practice;

IARC - International Agency for Research on Cancer; IATA - International Air Transport Association;

IBC - International Code for the Construction and Equipment of Ships carrying Dangerous Chemicals in Bulk;

IC50 - Half maximal inhibitory concentration; ICAO - International Civil Aviation Organization;

IECSC - Inventory of Existing Chemical Substances in China; IMDG - International Maritime Dangerous Goods;

IMO - International Maritime Organization; ISHL - Industrial Safety and Health Law (Japan);

ISO - International Organisation for Standardization; KECI - Korea Existing Chemicals Inventory;

LC50 - Lethal Concentration to 50 % of a test population; LD50 - Lethal Dose to 50% of a test population (Median Lethal Dose);

MARPOL - International Convention for the Prevention of Pollution from Ships; n.o.s. - Not Otherwise Specified;

NO(A)EC - No Observed (Adverse) Effect Concentration; NO(A)EL - No Observed (Adverse) Effect Level;

NOELR - No Observable Effect Loading Rate; NZIoC - New Zealand Inventory of Chemicals;

OECD - Organization for Economic Co-operation and Development; OPPTS - Office of Chemical Safety and Pollution Prevention;

PBT - Persistent, Bioaccumulative and Toxic substance; PICCS - Philippines Inventory of Chemicals and Chemical Substances; (Q)SAR - (Quantitative) Structure Activity Relationship; REACH - Regulation (EC) No 1907/2006 of the European Parliament and of the Council concerning the Registration, Evaluation, Authorisation and Restriction of Chemicals;

RID - Regulations concerning the International Carriage of Dangerous Goods by Rail; SADT - Self-Accelerating Decomposition Temperature; SDS - Safety Data Sheet; TCSI - Taiwan Chemical Substance Inventory; TECI - Thailand Existing Chemicals Inventory; TSCA - Toxic Substances Control Act (United States); UN - United Nations;

UNRTDG - United Nations Recommendations on the Transport of Dangerous Goods; vPvB - Very Persistent and Very Bioaccumulative