

According to REACH Regulation 1907/2006 and CLP Regulation 1272/2008.

SECTION 1: IDENTIFICATION OF THE SUBSTANCE/MIXTURE AND OF THE COMPANY/UNDERTAKING

1.1 Product identifier

Product Name Foetal Bovine Serum

Product code S-001#
CAS No. Not applicable.
EC No. Not applicable.
REACH Registration No. Not known.

1.2 Relevant identified uses of the substance or mixture and uses advised against Identified Use(s) Foetal Bovine Serum, filtered to 0.1 micron.

For Research Use Only.

Uses Advised Against Not for drug, household or other uses.

1.3 Details of the supplier of the safety data sheet

Manufacturer

Company Identification Life Science Group Ltd Address of Manufacturer Unit 14, Crowhill Farm,

Wilden

Bedfordshire Postal code MK44 2QE

Telephone: +44 (0) 1234 889180

Fax Not known.

E-mail sales@lifesciencegroup.co.uk

Office hours 08:00 - 18:00

Supplier

Company Identification Life Science Group Ltd Address of Supplier Unit 14, Crowhill Farm,

Wilden

Bedfordshire MK44 2QE

Postal code MK44 2QE Telephone: +44 (0) 1234 889180

Fax Not known.

E-mail sales@lifesciencegroup.co.uk

Office hours 08:00 - 18:00

1.4 Emergency telephone number

Emergency Phone No. +44 (0) 1234 889180 Contact Jenny Murray

SECTION 2: HAZARDS IDENTIFICATION

2.1 Classification of the substance or mixture

Not a hazardous substance or mixture according to Regulation (EC) No 1272/2008. This substance is not classified as dangerous according to Directive 67/548/EEC.

Although classed as non-hazardous contains animal source material. Handle as though capable of transmitting infectious agents. This material has been viral tested

and should be handled at Biohazard Safety Level 2.

2.2 Label elements

Product Name Foetal Bovine Serum

Hazard Pictogram(s) None.

Signal Word(s) None.

Hazard Statement(s) None.

Precautionary Statement(s) None.

2.3 Other hazards

None known.

2.4 Additional Information

None.

SECTION 3: COMPOSITION/INFORMATION ON INGREDIENTS

3.1 Substances

	CAS No.	EC No. / Registration	%W/W		Hazard
INGREDIENT(S)		number(s)			Pictogram(s)
Foetal Bovine Serum	n/a	n/a	>99	n/a	n/a

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The product contains no substances which at their given concentration, are considered to be hazardous to health. Although the serum used to manufacture this product has been tested and shown to be negative for certain infectious agents, no test is 100% accurate. All materials derived from blood should be handled as if capable of transmitting infection.

3.2 Mixtures

Not applicable.

For full text of H/P Statements see section 16

SECTION 4: FIRST AID MEASURES

4.1 Description of first aid measures

Inhalation If breathing is difficult, remove victim to fresh air and keep at rest in a position

comfortable for breathing.

Skin Contact Wash skin with soap and plenty of water.

Eye Contact Flush eyes with water as a precaution for at least 15 minutes.

Ingestion Never give anything by mouth to an unconscious person. Rinse mouth with

water. Call medical assistance immediately.

4.2 Most important symptoms and effects, both acute and delayed

None anticipated. Treat symptomatically.

To the best of our knowledge, the chemical, physical, and toxicological properties

have not been thoroughly investigated.

4.3 Indication of any immediate medical attention and special treatment needed

No information available.

SECTION 5: FIREFIGHTING MEASURES

5.1 Extinguishing media

Suitable Extinguishing media

As appropriate for surrounding fire.

Unsuitable extinguishing media None.

5.2 Special hazards arising from the substance or mixture

None anticipated.

5.3 Advice for firefighters

Wear self-contained breathing apparatus for firefighting if necessary.

SECTION 6: ACCIDENTAL RELEASE MEASURES

6.1 Personal precautions, protective equipment and emergency procedures

Provide adequate ventilation. Avoid breathing vapours, mist or gas. Wear suitable

gloves if prolonged skin contact is likely.

6.2 Environmental precautions

Contain the spill immediately and decontaminate using a string solution of sodium hypochlorite (bleach). Volume of bleach should be not less than 10% of the spill volume. Allow at least 10 minutes for neutralisation. Absorb spill using suitable

absorbent materials.

Dispose as clinical waste.

6.3 Methods and material for containment and cleaning up

Dispose as clinical waste.

6.4 Reference to other sections

See Also Section 8, 13.

SECTION 7: HANDLING AND STORAGE

7.1 Precautions for safe handling

Not known.

7.2 Conditions for safe storage, including any incompatibilities

Keep container tightly closed in a dry and well-ventilated place. Containers which

are opened must be carefully resealed and kept upright to prevent leakage.

Storage temperature -20°C

Storage life Stable under normal conditions.

Incompatible materials None known

7.3 Specific end use(s)

No information available.

SECTION 8: EXPOSURE CONTROLS/PERSONAL PROTECTION

8.1 Control parameters

8.1.1 Occupational Exposure Limits Contains no substances with occupational exposure limit values.

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8.2 Exposure controls

8.2.1. Appropriate engineering controls 8.2.2. Personal protection equipment

General industrial hygiene practice.

Eye Protection Use equipment for eye protection tested and approved under appropriate

government standards such as NIOSH (US) or EN 166(EU).



Handle with gloves. Gloves must be inspected prior to use. Use proper glove Skin protection

removal technique (without touching glove's outer surface) to avoid skin contact with this product. Dispose of contaminated gloves after use in accordance with

applicable laws and good laboratory practices.

Wash and dry hands.

The selected protective gloves have to satisfy the specifications of EU Directive

89/686/EEC and the standard EN 374 derived from it.

Respiratory protection

Respiratory protection not required. For nuisance exposures use type OV/AG (US) or type ABEK (EU EN 14387) respirator cartridges. Use respirators and components tested and approved under appropriate government standards such as NIOSH (US)

or CEN (EU).



Thermal hazards None known.

Environmental Exposure Controls Do not release large quantities into the surface water or into drains.

SECTION 9: PHYSICAL AND CHEMICAL PROPERTIES

9.1 Information on basic physical and chemical properties

Appearance Liquid.

Colour: Amber Odour Not known. Odour threshold Not known. Not known. Melting point/freezing point Not known. Initial boiling point and boiling range Not known. Flash Point Not known. Not known. Evaporation rate Flammability (solid, gas) Not known. Upper/lower flammability or explosive Not known.

limits

Vapour pressure Not known. Vapour density Not known Density (g/ml) Not known. Relative density Not known. Solubility(ies) Not known. Partition coefficient: n-octanol/water Not known. Auto-ignition temperature Not known. Decomposition Temperature (°C) Not known. Not known. Viscosity Explosive properties Not known. Oxidising properties Not known. 9.2 Other information

None.

SECTION 10: STABILITY AND REACTIVITY

10.1 Reactivity

No information available.

10.2 Chemical Stability

No information available.

10.3 Possibility of hazardous reactions

No information available.

10.4 Conditions to avoid

No information available.

10.5 Incompatible materials

Strong oxidising agents.

10.6 Hazardous decomposition products

No information available.

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SECTION 11: TOXICOLOGICAL INFORMATION

11.1 Information on toxicological effects

Acute toxicity - Ingestion
Acute toxicity - Skin Contact
Acute toxicity - Inhalation
Skin corrosion/irritation
Serious eye damage/irritation
Skin sensitization data
Respiratory sensitization data
Germ cell mutagenicity

No information available.

Carcinogenicity No component of this product present at levels greater than or equal to 0.1% is

identified as probable, possible or confirmed human carcinogen by IARC.

Reproductive toxicity
Lactation
STOT - single exposure
STOT - repeated exposure
Aspiration hazard
No information available.
No information available.
No information available.
No information available.

11.2 Other information

To the best of our knowledge, the chemical, physical, and toxicological properties

have not been thoroughly investigated.

SECTION 12: ECOLOGICAL INFORMATION

12.1 Toxicity

Toxicity - Aquatic invertebrates
Toxicity - Fish
Toxicity - Algae
Toxicity - Sediment Compartment
Toxicity - Terrestrial Compartment
12.2 Persistence and degradability

No information available.
No information available.
No information available.
No information available.

12.3 Bioaccumulative potential

No information available.

12.4 Mobility in soil

No information available.

12.5 Results of PBT and vPvB assessment

No information available.

12.6 Other adverse effects

No information available.

SECTION 13: DISPOSAL CONSIDERATIONS

13.1 Waste treatment methods

Observe all local and government environmental regulations.

Product: Offer surplus and non-recyclable solutions to a licensed disposal company.

Packaging: Dispose of as unused product.

13.2 Additional Information

No information available.

SECTION 14: TRANSPORT INFORMATION

IATA AIR TRANSPORTATION

Proper shipping name: Biological substance, Category B

Hazard Class: Class 6.2

14.1 UN number

UN3373

14.2 UN proper shipping name

Biological substance, Category B ADR/RID: Not dangerous goods. IMDG: Not dangerous goods. IATA: Not dangerous goods.

14.3 Transport hazard class(es)

Hazard Class: Class 6.2

14.4 Packing group

No information available



14.5 Environmental hazards

ADR/RID: No

IMDG Marine pollutant: No

IATA: No

14.6 Special precautions for user

Not known

14.7 Transport in bulk according to Annex II of Marpol and the IBC Code

Harmonised tariff code: 3002 1098

SECTION 15: REGULATORY INFORMATION

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

United Kingdom Regulations - Authorisations and/or Restrictions On Use

UK REACH Candidate List of Substances Not listed of Very High Concern for Authorisation UK REACH Authorisation List (Annex Not listed

XIV) list of substances subject to

authorisation

UK REACH Restrictions List (Annex XIV) Calcium chloride (CaCl2), dihydrate (10035-04-8)

Not listed

Not listed

Not listed

Restrictions on the manufacture, placing on the market and use of certain dangerous substances, mixtures and

articles

The Persistent Organic Pollutants

Regulations 2007 (SI 2007/3106) as

amended

The Ozone-Depleting Substances and

Fluorinated Greenhouse Gases (Amendment etc.) (EU Exit) Regulations

2019 (SI 2019/583)

The Prior Informed Consent (PIC)

Regulations concerning the export and

import of hazardous chemicals SI2008/2108 as amended

European Regulations - Authorisations and/or Restrictions On Use

Community Rolling Action Plan (CoRAP) Not listed

15.2 Chemical Safety Assessment

A REACH chemical safety assessment has not been carried out. United Kingdom

SECTION 16: OTHER INFORMATION

The following sections contain revisions or new statements:

LEGEND

Hazard Pictogram(s) None.

Hazard classification None.

Hazard Statement(s) None.

Precautionary Statement(s)

Acronyms

ATE: Acute Toxicity Estimate CAS : Chemical Abstracts Service

DNEL: Derived No Effect Level EC: European Community

EINECS: European Inventory of Existing Commercial Chemical Substances

LTEL: Long term exposure limit

PBT : Persistent, Bioaccumulative and Toxic PNEC: Predicted No Effect Concentration

REACH: Registration, Evaluation, Authorisation and Restriction of Chemicals

STEL: Short term exposure limit STOT: Specific Target Organ Toxicity

vPvB: very Persistent and very Bioaccumulative

Key literature references and sources for GB CLP Regulation, UK SI 2019/720 and UK SI 2020/1567

data used to compile the SDS

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