



# MATERIAL SAFETY DATA SHEET

according to Regulation (EC) No. 1907/2006 (REACH), Annex II,  
as amended by Regulation (EU) No. 453/2010

## LABiTec -Control Plasma 1-

### SECTION 1: Identification of the Substance/ Mixture and of the Company/ Undertaking

#### 1.1 Product identifier

Product name: **LABiTec Control Plasma 1**  
Product number: **210-12-000-00**

#### 1.2 Relevant identified uses of the substances or mixtures and uses advised against

LABiTec Control Plasma 1 may be used as a normal control when measuring PT, APTT and Fibrinogen. The LABiTec Control Plasma 1 is used in the same manner as a fresh plasma pool. The control plasma should be used to gauge internal factors in each laboratory's system. Source material for the plasma product has been tested and found negative for the presence of HIV and HCV antibodies as well as Hepatitis B Surface Antigen by approved test methods. However, no known test method can offer assurance that products derived from human blood are free of infectious agents. Therefore, handle this material observing the same safety precautions employed when handling any potentially infectious material.

#### 1.3. Details of the supplier of the Safety Data Sheet:

Contract Manufacturer: LABiTec® LAbor BioMedical Technologies GmbH  
Address: An der Strusbek 6  
Place: D-22926 Ahrensburg, Germany  
Telephone: +49(0) 4102/4795-0  
Fax: +49(0) 4102/4795-35  
E-mail: info@labitec.de

#### 1.4. Emergency contact

Telephone Number: +49(0) 4102/4795-0  
Opening hours ( ): 8:30am – 3:30pm



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### SECTION 2: Hazards Identification

#### 2.1 Classification of the substance/ mixture:

**2.1.1 Classification according to Regulation (EC) No 1272/2008 [CLP]**  
Non-hazardous

**2.1.2 Additional information:**  
Routine PPE is recommended for handling of biological material. For the full text of R-phrases mentioned, see SECTION 16.

#### 2.2 Label elements:

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

Hazard pictogram:  
None required since product non-hazardous

Signal word: None required

Hazard statement(s): None required

Precautionary statements: None required

Supplement Hazard Information (EU): Not applicable

#### 2.3 Other hazards:

None

### SECTION 3: Composition/ Information on Ingredients

#### 3.1 Mixtures:

Description of the mixture: LABiTec Control Plasma 1 is prepared from a frozen pool of citrated plasma and comprises of buffer and lyoprotectant to ensure stability of all plasma constituents.

##### Hazardous Ingredients:

Chemical concentrations in LABiTec Control Plasma 1 are not classifiable as hazardous according to (EC) No 1272/2008 [CLP].

### SECTION 4: First Aid Measures

#### 4.1 Description of first aid measures:

**General advice:** Consult a physician. Show this SDS to the doctor in attendance.

**After Inhalation:** - If inhaled, remove person to fresh air. If breathing becomes difficult obtain immediate medical attention, give artificial respiration. Consult a physician.



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**After Skin Contact:** - Immediately wash skin with soap and copious amounts of water. Consult a physician.

**After Eye Contact:** - Irrigate with copious amounts of clean fresh water for at least 15 minutes. If irritation persists seek medical attention.

**After Ingestion:** - Wash out mouth with water provided person is conscious. Urgently seek immediate medical attention. Consult a doctor/ physician or poison centre.

**Self-protection for first aider:** Personal protective equipment for first aid responder is recommended.

### **SECTION 5: Fire-fighting Measures**

#### **5.1 Extinguishing media:**

Suitable extinguishing media: Water, alcohol resistant foam, CO<sub>2</sub>, Dry Chemical Extinguisher.

#### **5.2 Special hazards arising from the substance or mixture:**

No hazardous decomposition materials known.

#### **5.3 Advice for firefighters**

Wear suitable protective clothing/ equipment. Wear self-contained breathing apparatus for firefighting if necessary.

### **SECTION 6: Accidental Release Measures**

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

##### **6.1.1 For non-emergency personnel**

Protective Equipment: Exercise appropriate precautions to minimise direct contact with skin or eyes and prevent inhalation of vapours – Wear personal protective equipment (PPE) such as gloves, safety glasses.

Emergency Procedures: Wear respiratory protection, avoid dust formation and avoid breathing vapours, dust, mist or gas. Ensure adequate ventilation and evacuate personnel to safe areas.

##### **6.1.2 For emergency responders**

Personal protective equipment for first aid responder/s is/ are recommended.

#### **6.2 Environmental precautions:**

Prevent further leakage or spillage if safe to do so and do not let product enter drains.



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

- 6.3.1 **For containment:** Spillages should be contained using absorbent material to prevent contamination of drains and watercourses.
- 6.3.2 **For cleaning up:** Pick up and arrange disposal without additional leaking of material. Use adsorbent materials and wash spill site for decontamination after material pickup. Waste material from spillages should be disposed of in accordance with local regulations.
- 6.3.3 **Other information:** Spilled material may cause surfaces to become slippery, clear spills immediately.

### 6.4 Reference to other sections

Refer to section 8 for exposure controls and personal protection and sections 13 for disposal considerations.

## SECTION 7: Handling and Storage

### 7.1 Precautions for safe handling

- 7.1.1 **Recommendations:** Wear appropriate PPE. Avoid contact with eyes, skin and clothing. For IVD Use Only. Not for medicinal use. DO NOT INGEST. Reduce the release of substance to the environment by avoiding spillages. For precautions see section 2.2.
- 7.1.2 **General occupational hygiene:** Do not eat, drink or smoke in work areas. Wash hands after use. Remove protective equipment before entering eating areas.

### 7.2 Conditions for safe storage including any incompatibilities:

Store securely in the original labelled container. Storage conditions according to product label.

### 7.3 Specific end use:

Use in accordance with product Instructions For Use provided in Kit. For In Vitro Diagnostic use only.

## SECTION 8: Exposure Controls/ Personal Protection

### 8.1 Control Parameters

- 8.1.1 **Components with workplace control parameters:** Contains no substances with occupational exposure limit values.



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### 8.2 Exposure controls

#### 8.2.1 Appropriate engineering controls:

Follow good clinical hygiene practices adopting suitable individual protective measures. Avoid contact with skin, eyes and clothing. Do not eat or smoke while handling the product. Wash hands before breaks and immediately after handling the product.

#### 8.2.2 Personal Protection Equipment:

##### 8.2.2.1 Eye/ face Protection:

Wear safety glasses or goggles. Use equipment for eye protection tested and approved under appropriate government standards such as EN 166(EU).

##### 8.2.2.2 Skin Protection:

Handle with gloves. Gloves must be inspected prior to use to ensure good condition. Use proper glove removal technique (without touching gloves outer surface) to avoid skin contact with this product. Dispose of contaminated gloves after use in accordance with good laboratory practice and applicable laws. Wash hands and dry.

##### 8.2.2.3 Body Protection:

Wear laboratory coat/ work coat or apron in line with protective equipment at the specific work place.

##### 8.2.2.4 Respiratory Protection

Wear simple mask to prevent inhalation of dust / vapour.

#### 8.2.3 Environmental Exposure Controls

Prevent further leakage or spillage if safe to do so as described in section 6.3. Do not let product enter drains.

## SECTION 9: Physical and Chemical Properties

### 9.1 Information on basic physical and chemical properties

- |    |   |  |
|----|---|--|
| a) | <b>Appearance:</b>                              | - Lyophilised: Buff compact cake at base of vial, sealed with stopper and screw cap. Once reconstituted the plasma is a pale straw colour. |
| b) | <b>Odour:</b>                                   | Odourless  |
| c) | <b>Odour threshold:</b>                         | No data available  |
| d) | <b>pH:</b>                                      | Neutral  |
| e) | <b>Melting point/ freezing point:</b>           | Melting point/ range: No data available  |
| f) | <b>Initial boiling point and boiling range:</b> | No data available  |
| g) | <b>Flash Point:</b>                             | No data available  |



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h)	Evaporation Weight	No data available
i)	Flammability (solid, gas):	No data available
j)	Upper/ lower flammability or explosive limits:	No data available
k)	Vapour Pressure:	No data available
l)	Vapour Density:	No data available
m)	Relative Density	No data available
n)	Water Solubility	No data available
o)	Partition Coefficient: n-octanol/ water	No data available
p)	Auto-ignition temperature	No data available
q)	Decomposition temperature	No data available
r)	Viscosity	No data available
s)	Explosive properties	No data available
t)	Oxidising Properties	No data available

### 9.2 Other safety information

Water soluble

### SECTION 10: Stability and Reactivity

**10.1 Reactivity:** No data available

**10.2 Chemical stability:** This product is stable in normal conditions of use and storage.

**10.3 Possibility of hazardous reactions:** No data available

**10.4 Conditions to avoid:** No data available

**10.5 Incompatible materials:** Bases, oxidising agents, iron and irons salts, copper.

**10.6 Hazardous decomposition products:** No known hazardous decomposition products

### SECTION 11: Toxicological Information

**11.1 Information on toxicological effects:**

**Acute Toxicity:** No data available

**Skin corrosion/ irritation:** No data available

**Serious Eye damage/ irritation:** No data available

**Respiratory or skin sensitisation:** No data available

**Germ cell mutagenicity:** No data available

**Carcinogenicity: IARC:** No data available

**Reproductive Toxicity:** No data available

**Summary of evaluation of the CMR properties:** No data available

**STOT-single exposure:** No data available

**STOT-repeated exposure:** No data available



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**Aspiration hazard:** No data available  
**Additional Information:** No data available

### SECTION 12: Ecological Information

**12.1 Toxicity:** No data available

**12.2 Persistence and degradability:** No data available

**12.3 Bio-accumulative potential:** No data available

**12.4 Mobility in soil:** No data available

**12.5 Results of PBT and vPvB assessment:** No data available

**12.6 Other adverse effects:** No data available

Use according to good clinical hygiene practices; avoid dispersion of the product in the environment.

### SECTION 13: Disposal Considerations

#### **13.1 Waste treatment methods**

**Product:** Disposal of waste must always comply with existing EEC, national and local regulations. Registered waste carriers and licensed disposal sites must be used.

**Contaminated packaging:** Dispose of as unused product.

### SECTION 14: Transport Information

**14.1 UN number:** No number available/ not required for mixture

**14.2 UN Proper Shipping name:** Not required

**14.3 Transport Hazard Classes:** Not required

**14.4 Packaging Group:** Not required

**14.5 Environmental Hazards:** Not required

**14.6 Special precautions for user:** Not required

**14.7 Transport in bulk according to Annex II MARPOL73/78 and the IBC Code:** It is not intended that this product is transported as bulk and therefore the shipping/ transport regulations for bulk hazardous material do not apply.



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### SECTION 15: Regulatory Information

**15.1 Safety, health and environmental regulations/ legislation specific for the substance or mixture:** None required

**15.2 Chemical Safety Assessment:** No chemical safety assessment has been carried out for this substance/ mixture by the supplier.

### SECTION 16: Other Information

**Full text of H-Statements referred to under sections 2 and 3**

Not applicable

**Full text of R-phrases referred to under sections 2 and 3**

Not applicable

Classification and procedure used to derive the classification for mixtures according to Regulation (EC) 1272/2008 [CLP]:

- LABiTec Control Plasma 1 is a non-hazardous product

**Note for the user:** The data contained in this sheet is the result of the most up to date information available in our company. The user must ensure that the information is suitable and complete in relation to the specific use of the product.