

SAFETY DATA SHEET

STREPTOCOCCUS Latex Test Kit

1. PRODUCT AND COMPANY INFORMATION

Company name: **Lab21 Healthcare Ltd,**
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Web site: www.Lab21.com

Emergency contact: As above, during office hours.

Product name: **Streptococcus Latex Test Kit**

Product code: **Test Kits:** STE010; 2/763
Strep Latex; Groups:
A (STE/100A and versions thereof), **B** (STE/100B and versions thereof),
C (STE/100C and versions thereof), **D** (STE/100D and versions thereof),
F (STE/100F and versions thereof) and **G** (STE/100G and versions thereof)
Extraction enzyme (STE/100EE)

2. HAZARD IDENTIFICATION

Main hazards: These products are for in vitro diagnostic use only.
Specimen material may contain pathogenic organisms. Handle with the appropriate precautions, according to good laboratory practices.
Product contains no hazardous constituents or the concentrations of all chemical constituents are below the regulatory threshold described in Article 31 Requirements for Safety Data Sheets.
All reagents contain less than 0.1% w/w sodium azide (NaN₃) as preservative.

3. COMPOSITION / INFORMATION ON INGREDIENTS

Composition: Test Latex: Sodium Azide <0.1%
Extraction Enzyme: Merthiolate (Thiomersal) <0.01% in working strength

Hazardous Components: SODIUM AZIDE: <1% w/w EINECS: 247-852-1 CAS: 26628-22-8
[T+] R28; [-] R32; [N] R50/53

THIOMERSAL (Merthiolate): <0.01% CAS:54-64-8 [T+] R26/27/28; [Xn-]
R33; [N] R50/53

4. FIRST AID MEASURES

Skin contact: May cause mild irritation at the site of contact. Wash skin with soap and water.
Eye contact: May cause irritation and redness. Flush with water.
Ingestion: Avoid hand to mouth contact.

5. FIRE-FIGHTING MEASURES

Extinguishing media: Not combustible. Suitable extinguishing media for the surrounding fire should be used.
Exposure hazards: None in small quantities

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6. ACCIDENTAL RELEASE MEASURES

- Personal precautions:** Wear appropriate protective clothing. Refer to section 8 of MSDS for personal protection details.
- Environmental precautions:** Properly disinfect any spills. Do not discharge into drains or rivers. Contain large spillages using bunding.
- Clean-up procedures:** Absorb into dry earth or sand. Transfer to a closable, labelled salvage container for disposal by an appropriate method.

7. HANDLING AND STORAGE

- Handling requirements:** For in vitro diagnostic use only. Read the instructions for use. Avoid the formation of aerosols. Avoid direct contact with the substance
- Storage conditions:** Store in cool (2° to 8°C), well-ventilated area. Keep container tightly closed.

8. EXPOSURE CONTROLS / PERSONAL PROTECTION

Hazardous ingredients:

SODIUM AZIDE: WEL (8 hr TWA): 0.1 mg/m³ WEL (15 min STEL): 0.3 mg/m³

Respiratory protection: Respiratory protection not required.

Hand protection: Protective disposable gloves.

Eye protection: Safety glasses. Ensure eye bath is to hand.

Skin protection: Protective clothing.

THIOMERSAL: WEL (8 hr TWA): 0.05 mg/m³ WEL (15 min STEL): 0.15 mg/m³

Respiratory protection: Respiratory protection must be available.

Hand protection: Impermeable gloves.

Eye protection: Safety glasses. Ensure eye bath is to hand.

Skin protection: Impermeable protective clothing.

9. PHYSICAL AND CHEMICAL PROPERTIES

- State:** Liquid
- Colour:** Latex: White/Off white
Positive and Negative Controls: colourless/ straw coloured
- Odour:** Odourless
- pH:** pH6.8 – pH 8.4
- Solubility:** Soluble
- Flammability:** Not combustable

10. STABILITY AND REACTIVITY

- Stability:** Stable under normal storage and handling conditions. Do not use after expiry date.
- Materials to avoid:** Avoid contact of the products with lead and copper (plumbing metals), mercury, acids, and oxidising agents.

Hazardous decomposition products: In combustion emits toxic fumes.

11. TOXICOLOGICAL INFORMATION

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Hazardous ingredients: **SODIUM AZIDE:**

ORL MUS LD50 27 mg/kg

ORL RAT LD50 27 mg/kg

SKN RAT LD50 50 mg/kg

THIOMERSAL:

IPR MUS LD50 54 mg/kg

IVN MUS LD50 45 mg/kg

ORL MUS LD50 91 mg/kg

ORL RAT LD50 75 mg/kg

SCU MUS LD50 66 mg/kg

SCU RAT LD50 98 mg/kg

UNR RAT LD50 40 mg/kg

Routes of exposure: Refer to section 4 of SDS for routes of exposure and corresponding symptoms.

12. ECOLOGICAL INFORMATION

Mobility: Readily absorbed into soil.

Persistence and degradability: No data available.

Bio-accumulative potential: No data available.

Other adverse effects: No data available.

13. DISPOSAL CONSIDERATIONS

Waste code number: 18 01 07 Hazardous waste.

NB: The user's attention is drawn to the possible existence of regional or national regulations regarding disposal.

14. TRANSPORT INFORMATION

SODIUM AZIDE:

ADR / RID

UN no: - Not applicable.

Shipping name: Not classified as dangerous in the meaning of transport regulations.

IMDG / IMO

UN no: - **Marine pollutant:** NO

IATA / ICAO

UN no: - Not applicable.

THIOMERSAL:

ADR / RID

ADR Class 6.1

UN no:2024

Shipping name: MERCURY COMPOUND, LIQUID N.O.S

Labelling: 6.1 Classification Code: T4 Hazard ID: 66

IMDG / IMO

Class 6.1

UN no:2024

Marine Pollutant:PP

Shipping name: MERCURY COMPOUND, LIQUID N.O.S

Labelling: 6.1 EmS: F-A,S-A

IATA/ICAO

Class 6.1

UN no:2024

Packing instructions: 610(P&CA); 605(CAO)

Labelling: 6.1

15. REGULATORY INFORMATION

Date of first issue: 30/07/2008

Version: 02 01 November 2012

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SODIUM AZIDE:

Hazard symbols:

Harmful.

Risk phrases:

R22: Harmful if swallowed.

R32: Contact with acids liberates very toxic gas.

EC classification:

Xn- Harmful

Safety phrases:

S29/35: Do not empty into drains; dispose of this material and its container in a safe way.

S36/37/39: Wear suitable protective clothing, gloves and eye / face protection.

S46: If swallowed, seek medical advice immediately and show this container or label.

THIOMERSAL:

Hazard symbols:

Harmful

Risk phrases:

R20/21/22: Harmful by inhalation, in contact with skin and if swallowed.

R52/53: Harmful to aquatic organisms, may cause long-term adverse effects in the aquatic environment.

EC classification:

Xn- Harmful

Safety phrases:

S24/25: Avoid contact with skin and eyes.

S26: In case of contact with skin, wash immediately with plenty of water.

Note: The regulatory information given above only indicates the principal regulations specifically applicable to the product described in the safety data sheet. The user's attention is drawn to the possible existence of additional provisions that complete these regulations. Refer to all applicable national, international and local regulations or provisions.

16. OTHER INFORMATION

Warning: Because no test method can offer complete assurance that HIV, HCV, HbsAg or other infectious agents are absent, the components of this kit should be handled accordingly.

Legal disclaimer: The above information is believed to be correct but does not purport to be all inclusive and shall be used only as a guide. This company shall not be held liable for any damage resulting from handling or from contact with the above product.

Sources of information used in this data sheet:

Regulation (EC) No. 1907/2006 of the European Parliament and of the Council concerning the registration, evaluation, authorisation and restriction of chemicals (REACH): Article 31:- Requirements for safety data sheets, and Annex II:- Guide to the Compilation of Safety Data Sheets, OJ L 136/35 – L 136/36 and L 136/84 – L 136/92, 29.5.2007.

The Chemicals (Hazard Information & Packaging for Supply) Regulations [CHIP], United Kingdom Statutory Instrument 2002 No. 1689, based on the European Directives on Dangerous Substances (67/548/EEC) and Dangerous Preparations (1999/45/EC).

Approved Supply List (8th edition), information approved for the classification and labelling of substances and preparations dangerous for supply, United Kingdom Health & Safety Commission, 2005, based on Annex I to 67/548/EEC.

Commission Decision 2000/532/EC establishing a list of wastes pursuant to Article 1(a) of Directive 75/442/EEC on waste and Article 1(4) of Directive 91/689/EEC on hazardous waste.
CONSLEG: 2000D0532 – 01/01/2002, Office for Official Publications of the European Communities.
Directive 98/79/EC of the European Parliament and of the Council on *in vitro* diagnostic medical devices, Annex I, Essential Requirements, OJ L 331/20, 7.12.98.

List of approved workplace exposure limits, Table 1 of EH40/2005, United Kingdom Health & Safety Commission.