



Works:

Plot Nos. M 46 - 47, Phase III B, Verna Industrial Estate,
Verna, Goa - 403 722, INDIA.

Tel.: (0832) 6624502 / 6682000, Fax: (0832) 6680156.

E-mail: zephyr@tulipgroup.com

MATERIAL SAFETY DATA SHEET

1. IDENTIFICATION

1.1. Identification of the product

Product name: **electra HSV 1,2 IgM**

Chemiluminescence Assay for Qualitative Detection of HSV 1,2 IgM antibody in Human serum.

Product classification: In-vitro diagnostics

1.2. Manufacturer identification

Company Name: ZEPHYR BIOMEDICALS-A Div. of Tulip Diagnostics [P] Ltd.

Address: Plot Nos. M 46/47, Phase III-B, Verna Industrial Estate,
Verna, Goa, 403 722, India

Phone: 91 832 6682000

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E-mail: zephyr@tulipgroup.com

2. HAZARDS IDENTIFICATION

2.1. Potential biohazard:

The calibrator, negative and positive controls are formulated with a buffer base, animal serum, and human serum. The human serums are tested by a licensed method and found to be non-reactive for HIV-1, HIV-2, Hepatitis B surface antigen and HCV. Because no test method can offer absolute assurance that these agents are absent, reagents should be handled at the Biosafety Level 2, as recommended for any potentially infectious human blood product

2.2. Chemical hazard:

Chemiluminescent substrate A with enhanced Luminol solution:

Eye: Causes eye irritation.

Skin: Causes skin irritation.

Ingestion: May cause gastrointestinal irritation with nausea, vomiting and diarrhea.

Inhalation: Causes respiratory tract irritation.

Chemiluminescent substrate B with stabilized peroxide solution:

May cause fire or explosion; strong oxidizer

Causes severe skin burns and eye damage

Harmful if swallowed or if inhaled

3. COMPOSITION OF THE KIT COMPONENTS

Coated Microwells: Purified HSV 1,2 antigen coated wells

Sample Diluent

Negative Control





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Positive Control

Calibrator

Enzyme Conjugate

Wash Buffer Concentrate (20X)

Substrate A: Chemiluminescent substrate containing enhanced luminol solution.

Substrate B: Chemiluminescent substrate containing stabilized peroxide solution.

4. FIRST-AID MEASURES

4.1. Skin contact: Remove contaminated clothing and wash the exposed skin area thoroughly with soap and water.

4.2. Eye contact: Flush with copious amounts of water for at least 15 minutes. Seek medical attention.

4.3. Inhalation: Remove to fresh air, give oxygen if breathing becomes difficult and seek medical attention.

4.4. Ingestion: Flush mouth with copious amounts of water, provided that the person is conscious and seek medical attention.

5. FIRE-FIGHTING MEASURES

5.1. Suitable extinguishing media: Chemical or water fire extinguisher.

5.2. Extinguishing media not to be used: None known.

5.3. Special fire-fighting procedures: None known.

5.4. Unusual fire and explosion hazards: Not available

5.5. NFPA rating: Health: 2, Flammability: 0, Reactivity: 1

6. ACCIDENTAL RELEASE MEASURES

6.1. Body precautions: Wear rubber gloves, impermeable shoe covers and long laboratory coat.

6.2. Environmental precautions: Contain the spill to the smallest area possible.

6.3. Cleaning measures: Take care not to contaminate body. Absorb the material with disposable towels. Soak area with a 2% Sodium Hypochlorite solution and wipe up with disposable towels. Dispose of all contaminated trash in accordance with local regulations.

7. HANDLING AND STORAGE

7.1. Handling: Wear suitable personal protective equipment. Take care not to splash spill or splatter reagents. Do not eat, drink, smoke or apply cosmetics in laboratory areas. Do not pipette samples or reagents by mouth.

7.2. Storage: Store the test kits in 2 - 8° C refrigerators designated and labeled to contain human blood products.

8. EXPOSURE CONTROLS/PERSONAL PROTECTION

8.1. Body protection: Wear long laboratory coat.

8.2. Respiratory protection: In case of fire, wear self-contained breathing apparatus.





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8.3. Hand protection: Wear non-permeable rubber, neoprene, latex or nitrile disposable gloves.
Change gloves when they become contaminated.

8.4. Eye protection: Wear safety glasses or goggles when a splash hazard exists.

9. PHYSICAL AND CHEMICAL PROPERTIES

9.1. Physical state (at 20°C): Microtiter plate is in solid form. All other reagents are in liquid form.

9.2. Odor: Odorless

9.3. pH value: All reagents 6.8 ~ 8.0

9.4. Melting point/freezing Point: Not determined

9.5. Boiling point: Not determined

9.6. Evaporation rate: Not determined

9.7. Flammability (solid, gas): Not flammable

9.8. Upper/lower flammability or explosive limits: Not applicable

9.9. Vapor pressure: Not determined

9.10. Vapor density: Not determined

9.11. Relative density: Not determined

9.12. Solubility: Water soluble

9.13. Auto-ignition temperature: Not applicable

9.14. Decomposition temperature: Not determined

9.15. Viscosity: Not determined

9.16. Explosive properties: None

9.17. Oxidizing properties: Not determined

10. STABILITY AND REACTIVITY

10.1. Stability

Stable under recommended storage conditions.

10.2. Reactivity

No known reactivity hazards associated with product.

10.3. Possibility of hazardous reactions

No hazardous polymerization

10.4. Conditions to avoid

Excessive heat and light

10.5. Incompatible materials

Acids





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11. TOXICOLOGICAL INFORMATION

- 11.1. **Acute toxicity:** Not determined
- 11.2. **Skin corrosion/irritation:** In case of contact, flush skin with plenty of water. Remove contaminated clothing and shoes. Get medical aid if irritation develops and persists. Wash clothing before reuse.
- 11.3. **Serious eye damage/irritation:** Not determined
- 11.4. **Respiratory or skin sensitization:** Not determined
- 11.5. **Germ cell mutagenicity:** Not determined
- 11.6. **Carcinogenicity:** None known
- 11.7. **Reproductive toxicity:** Not determined
- 11.8. **STOT-single exposure:** Not determined
- 11.9. **STOT-repeated exposure:** Not determined
- 11.10. **Aspiration hazard:** Not determined

12. ECOLOGICAL INFORMATION None known.

13. DISPOSAL CONSIDERATIONS

- 13.1. **Disposal of reagents:** Dispose according to current regional and national regulations.
- 13.2. **Contaminated containers:** Dispose following current regional and national regulations.

14. TRANSPORT INFORMATION

- 14.1. **UN number:** Not available
- 14.2. **UN proper shipping name:** Not available
- 14.3. **Transport hazard class(es):** Not available
- 14.4. **Packing group:** Not available
- 14.5. **Environmental hazards:**
 - Overland transport (ADR/RID): None
 - Water transport (ADN/IMDG): None
 - Air transport (ICAO/IATA): None
- 14.6. **Special precautions for user:** None

15. REGULATORY INFORMATION

- 15.1. **Classification:** This preparation is classified and labeled in accordance with applicable national and international regulations.
- 15.2. **Symbols of danger:** N/A
- 15.3. **"R" phrases indicating specific risks:** Irritant R36/37/38, Irritating to the eyes, respiratory system and skin.
- 15.4. **"S" phrases indicating caution:** S26, In case of contact with any reagents rinse with copious amounts of water for at least 15 minutes. If irritation persists, seek medical attention. S36, Wear suitable protective clothing.





TULIP DIAGNOSTICS (P) LTD.

a revvity company

CIN: U85195GA1988PTC005717

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16. OTHER INFORMATION

Recommended Use: For In Vitro Diagnostic Use only.

Not for use on or in human and animals. Information contained in this MSDS relies upon our best knowledge of the product upon issue of this document to describe the use, storage, transport and disposal under safe and secure measures. Such information is not a guarantee of the characteristics and specific quality of the product. This MSDS concerns the stated preparation and cannot apply to the product if used with other materials or in a process other than those specifically stated.

MSDS established : May 2022

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MSDS electra HSV 1,2 IgM, Page 5 of 5



Zephyr Biomedicals

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