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Material Safety Data Sheet RhoGAM UF	Page: 1 Rev. Date 09/04/97
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1. CHEMICAL PRODUCT AND COMPANY IDENTIFICATION

Ortho Diagnostic Systems Inc.
U.S. Route 202
Raritan, NJ 08869

TELEPHONE NUMBER: (800)322-ODSI

PRODUCT NAME: RhoGAM UF - RhoGAM, MicRhoGAM
PRODUCT CODE: 780705,780725,780795,7807971,780805,780825

SYNONYMS: MICRhoGAM-5%
RhoGAM Rho(D)

2. COMPOSITION/INFORMATION ON INGREDIENTS

This product contains no hazardous components as defined in OSHA 29 CFR 1910.1200

This product contains Thimerosal as a preservative at concentrations less than 0.1%

3. HAZARDS IDENTIFICATION

POTENTIAL HEALTH EFFECTS

EYES
None known

SKIN
None known

INGESTION
None known

INHALATION
None known

4. FIRST AID MEASURES

EYES
In case of eye contact, flush eyes with plenty of water for at least 15 minutes. Seek medical attention if irritation develops.

SKIN
Promptly wash exposed areas thoroughly with soap and water. If irritation persists, seek medical attention.

INGESTION
Seek medical attention.

INHALATION
Remove individual to fresh air. Seek medical attention.

5. FIRE FIGHTING MEASURES

FLAMMABLE PROPERTIES
FLASH POINT: None
LOWER EXPLOSIVE LIMIT (%): N/A
UPPER EXPLOSIVE LIMIT (%): N/A

FIRE AND EXPLOSION HAZARDS
None known

EXTINGUISHING MEDIA
Use any extinguishing agent which is suitable for the surrounding fire.

5. FIRE FIGHTING MEASURES - Continued**FIRE FIGHTING INSTRUCTIONS**

None normally required.

6. ACCIDENTAL RELEASE MEASURES

Contain spill by placing a suitable absorbent material around the edges of the spill and work inward. Carefully scoop up into an appropriate waste container for disposal. Dispose of in accordance with local, state and federal requirements.

7. HANDLING AND STORAGE**HANDLING AND STORAGE PRECAUTIONS**

Do not use beyond expiration date. Store at 2° to 8°C. May be at room temperature (20°C to 30°C) while in use.

CAUTION: All blood products should be treated as potentially infectious. Source material from which this product was derived was found negative when tested in accordance with FDA required tests. No known test methods can offer assurance that products derived from human blood will not transmit infectious agents. Handle as if capable of transmitting infectious agents.

WARNING: For intramuscular injection only. Do not inject intravenously.

Parenteral drug products should be inspected visually for particulate matter and discoloration prior to administration, whenever solution and container permit.

Since there is no lot number and expiration date on the prefilled syringes, they should not be removed from the protective pouch until immediately before use.

WORK/HYGIENIC PRACTICES

Use good hygienic practices. Wash thoroughly after handling.

8. EXPOSURE CONTROLS/PERSONAL PROTECTION**ENGINEERING CONTROLS**

None normally required.

EYE/FACE PROTECTION

None normally required.

SKIN PROTECTION

Latex gloves are recommended.

RESPIRATORY PROTECTION

None normally required.

OTHER/GENERAL PROTECTION

None normally required.

9. PHYSICAL AND CHEMICAL PROPERTIES**APPEARANCE**

Clear liquid

ODOR

Odorless

BASIC PHYSICAL PROPERTIES

VAPOR PRESSURE: Not Determined

SPECIFIC GRAVITY: 1.02

SOLUBILITY (H₂O): Not Determined

pH: 6.3-6

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09/04/97**10. STABILITY AND REACTIVITY****STABILITY:** Stable**CONDITIONS TO AVOID (STABILITY)**

None known

INCOMPATIBLE MATERIALS

None known

HAZARDOUS DECOMPOSITION PRODUCTS

None known

HAZARDOUS POLYMERIZATION: Will Not Occur**11. TOXICOLOGICAL INFORMATION****MISCELLANEOUS TOXICOLOGICAL INFORMATION**

Carcinogenicity: NTP: No IARC: No OSHA: No

Systemic reactions associated with administration of MICRhoGAM are extremely rare. Discomfort at the site of injection has been reported and a small number of women have noted a slight elevation in temperature.

MEDICAL CONDITIONS AGGRAVATED BY EXPOSURE

Individuals known to have had an anaphylactic or severe systematic reaction to human globulin should not receive MICRhoGAM or any other Rho(D) Immune Globulin (Human).

12. ECOLOGICAL INFORMATION**OTHER ENVIRONMENTAL INFORMATION**

This product does NOT contain any ingredients which are regulated on the U.S. EPA List of Toxic Chemicals (40 CFR 372), and is therefore not subject to release reporting under Section 313 of EPCRA.

The health effects noted above do not meet the criteria for establishment of a SARA Hazard Classification for this product.

This product contains Thimerosal as a preservative at concentrations less than 0.1%.

13. DISPOSAL CONSIDERATIONS

Dispose of in accordance with local, state and federal regulations.

Certain states may govern disposal of this product under regulated medical waste requirements.

14. TRANSPORT INFORMATION**PROPER SHIPPING NAME:** None required**15. REGULATORY INFORMATION**

NO DATA GIVEN

16. OTHER INFORMATION**REFERENCE DOCUMENTATION**

Primary references used in the preparation of this document:

1. Ortho Diagnostics Product Circular
2. OSHA Z-Table, 1910.1000 (Revised Final Rule)
3. ACGIH Threshold Limit Values and Biological Exposure Indices (1996)

16. OTHER INFORMATION - Continued

REFERENCE DOCUMENTATION - Continued

N/A = Not Applicable

- = Approximately Equal To

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Ortho Diagnostic Systems Inc.