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FUJISAWA USA INC, FUJISAWA PHARMACEUTICAL CO DIV -- KANAMYCIN SULFATE INJECTION -- 6505-01-346-2061

============= Product Identification =================

Product ID:KANAMYCIN SULFATE INJECTION

MSDS Date:10/07/1992

FSC:6505

NIIN:01-346-2061

MSDS Number: BQXYY === Responsible Party ===

Company Name: FUJISAWA USA INC, FUJISAWA PHARMACEUTICAL CO DIV

Address: PARKWAY NORTH CENTER, 3 PARKWAY N

City:DEERFIELD

State:IL ZIP:6001 5-2548 Country:US

Info Phone Num:708-317-0800

Emergency Phone Num:708-317-0800, 800-888-7704

Preparer's Name: GREGORY KOLAR

CAGE:24832

=== Contractor Identification ===

Company Name: FUJISAWA USA INC (LYPHOMED INC)

Address: 3 PARKWAY NORTH

City:DEERFIELD

State:IL

ZIP:60015-2548

Country:US

Phone:800-727-7003/708-317-8800

CAGE:24832

Company Name: FUJISAWA USA, INC

Address:3 PKY N Box:City:DEERFIELD

State:IL

ZIP:60015-2548 Country:US

Phone:847-317-1256

CAGE:39832

Company Name: FUJISAWA USA, INC.

Address

:3 PARKWAY NORTH Box:City:DEERFIELD State:IL

ZIP:60015-2548 Country:US

Phone:708-317-0800 OR 800-888-7704

CAGE:0K706

======= Composition/Information on Ingredients ========

Ingred Name: KANAMYCIN SULFATE

CAS:25389-94-0 RTECS #:NZ3225000

Other REC Limits: NONE RECOMMENDED

Ingred Name: SODIUM METABISULFITE

CAS:7681-57-4

RTECS #:UX8225000

Other REC Limits: NONE RECOMMENDED

Ingred Name: SODIUM CITRATE

CAS:6132-04-3

Other REC Limits: NONE RECOMMENDED

Ingred Name: WATER FOR INJECTION (NOTE:

PH ADJUSTED WITH SULFURIC ACID)

CAS:7732-18-5

RTECS #:ZC0110000

Other REC Limits: NONE RECOMMENDED

========== Hazards Identification =================

LD50 LC50 Mixture:INTRAVENOUS RAT LD50: 225 MG/KG Routes of Entry: Inhalation:YES Skin:YES Ingestion:YES Reports of Carcinogenicity:NTP:NO IARC:NO OSHA:NO

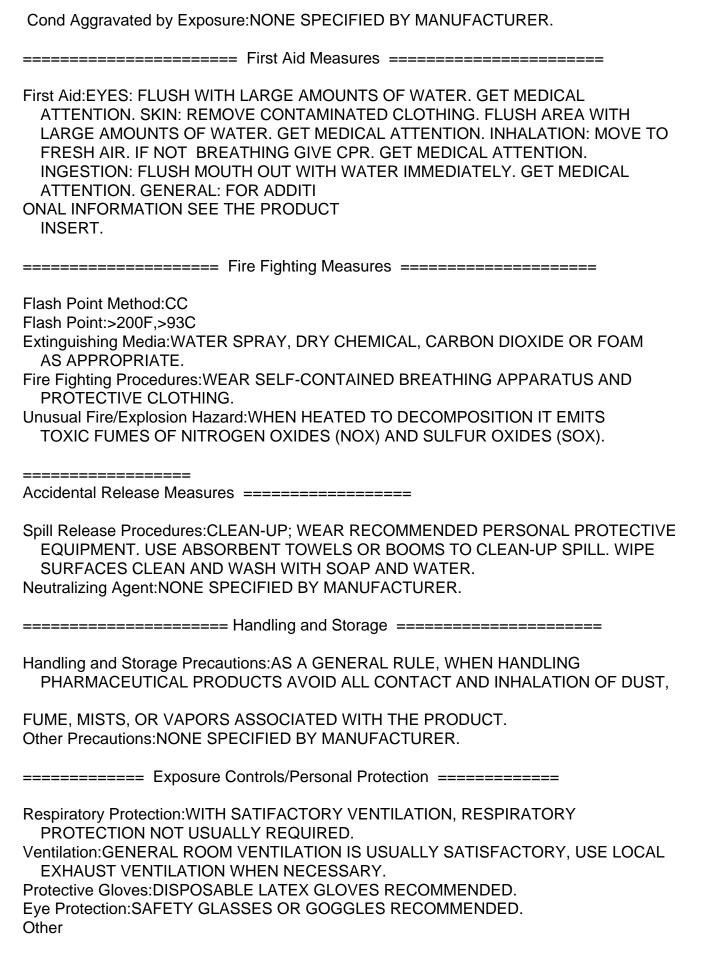
Health Hazards Acute and Chronic:EYE, SKIN, AND RESPIRATORY IRRITATION MAY OCCUR. THIS PRODUCT IS INTENDED FOR THERAPUTIC USE ONLY WHEN PRESCRIBED BY A PHYSICIAN.

POTENTIAL ADVERSE REACTIONS FROM

PRESCRIBED DOSES AND OVERDOSES ARE DE SCRIBED IN THE PACKAGE INSERT. OCCUPATIONAL EXPOSURE HAS NOT BEEN FULLY INVESTIGATED.

Effects of Overexposure:EYE, SKIN, AND RESPIRATORY IRRITATION MAY OCCUR. THIS PRODUCT IS INTENDED FOR THERAPUTIC USE ONLY WHEN PRESCRIBED BY A PHYSICIAN. POTENTIAL ADVERSE REACTIONS FROM PRESCRIBED DOSES AND OVERDOSES ARE DE SCRIBED IN THE PACKAGE INSERT. OCCUPATIONAL EXPOSURE HAS NOT BEEN FULLY INVESTIGATED.

Medical



Stability Indicator/Materials to Avoid:YES
NONE SPECIFIED BY MANUFACTURER.
Stability Condition to Avoid:PROTECT FROM FREEZING.
Hazardous Decomposition Products:WHEN HEATED TO DECOMPOSITION IT EMITS
TOXIC FUMES OF NITROGEN OXIDES (NOX) AND SULFUR OXIDES (SOX).
Conditions to Avoid Polymerization:WILL NOT OCCUR.

======= Disposal Considerations ===========

Waste Disposal Methods:DISPOSE OF WASTE IN ACCORDANCE WITH LOCAL, STATE AND FEDERAL REGULATIONS.

## Disc

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