

SAFETY DATA SHEET

SECTION 1 : IDENTIFICATION

Product Name:	Heparin Sodium in 5% Dextrose Injection
Synonyms:	Heparin Sodium in 5% Dextrose Injection (freeflex®)
Distributor Name:	Fresenius Kabi USA, LLC
Address:	Three Corporate Drive Lake Zurich, Illinois 60047
General Phone Number:	(847) 550-2300
Customer Service Phone Number:	(888) 386-1300
Health Issues Information:	(800) 551-7176
SDS Creation Date:	April 23, 2018
SDS Revision Date:	March 06, 2024

SECTION 2 : HAZARD(S) IDENTIFICATION

GHS Class:	Not classified
Emergency Overview:	This product is intended for therapeutic use only when prescribed by a physician. Potential adverse reactions from prescribed doses and overdoses are described in the package insert.
Route of Exposure:	Inhalation Ingestion Eye contact Skin Absorption. Injection.
Potential Health Effects:	
Eye:	Contact with eyes may cause irritation.
Skin:	May cause skin irritation.
Inhalation:	May cause irritation of respiratory tract.
Ingestion:	May cause irritation.
Signs/Symptoms:	Adverse reactions from prescribed doses include: hemorrhage, local irritation, erythema, mild pain, hematoma, ulceration, hypersensitivity reactions (chills, fever, urticaria), asthma, rhinitis, lacrimation, headache, nausea, vomiting, anaphylactic reactions including shock, itching, burning, and thrombocytopenia. The chief sign of heparin overdose is bleeding (nosebleeds, blood in urine, tarry stools, easy bruising, or petechial formations). When clinical circumstances require reversal of heparinization, protamine sulfate should be administered. Occupational exposure has not been fully investigated.
Aggravation of Pre-Existing Conditions:	Individuals with documented hypersensitivity to heparin sodium and individuals with thrombocytopenia or increased risk for hemorrhage.

SECTION 3 : COMPOSITION/INFORMATION ON INGREDIENTS

Chemical Name	CA S#	Ingredient Percent	EC Num.
Citric Acid, anhydrous, USP	77-92-9	0.51 mg/mL by weight	
Sodium Metabisulfite, NF	7681-57-4	0.2 mg/mL by weight	
Dextrose, USP (monohydrate)	14431-43-7	50 mg/mL by weight	
Heparin Sodium	9041-08-1	40 U/mL, 50 U/mL or 100 U/mL by weight	
Sodium Citrate, Dihydrate, USP	6132-04-3	3.34 mg/mL by weight	
Water for Injection	7732-18-5	QS to 1 mL by weight	

SECTION 4 : FIRST AID MEASURES

Eye Contact:

Immediately flush eyes with plenty of water for at least 15 to 20 minutes. Ensure adequate flushing of the eyes by separating the eyelids with fingers. Get immediate medical attention.

Skin Contact:

Immediately wash skin with plenty of soap and water for 15 to 20 minutes, while removing contaminated clothing and shoes. Get medical attention if irritation develops or persists.

Inhalation:	If inhaled, remove to fresh air. If not breathing, give artificial respiration or give oxygen by trained personnel. Seek immediate medical attention.
Ingestion:	If conscious, flush mouth out with water immediately. Call a physician or poison control center immediately. Do not induce vomiting unless directed to do so by medical personnel. Never give anything by mouth to an unconscious person.
Other First Aid:	For Adverse Event Information, please call (800) 551-7176.

SECTION 5 : FIRE FIGHTING MEASURES

Flash Point:	Not established.
Flash Point Method:	Not established.
Auto Ignition Temperature:	Not established.
Lower Flammable/Explosive Limit:	Not established.
Upper Flammable/Explosive Limit:	Not established.
Fire Fighting Instructions:	Evacuate area of unprotected personnel. Use cold water spray to cool fire exposed containers to minimize risk of rupture. Do not enter confined fire space without full protective gear. If possible, contain fire run-off water.
Extinguishing Media:	Use alcohol resistant foam, carbon dioxide, dry chemical, or water fog or spray when fighting fires involving this material. Use extinguishing measures that are appropriate to local circumstances and the surrounding environment.
Protective Equipment:	As in any fire, wear Self-Contained Breathing Apparatus (SCBA), MSHA/NIOSH (approved or equivalent) and full protective gear.
Hazardous Combustion Byproducts:	Thermal decomposition products may include smoke and toxic fumes. Oxides of carbon, oxides of nitrogen and other organic substances may be formed. Other undetermined low molecular weight hydrocarbon compounds may be released in small quantities depending upon specific conditions of combustion.

SECTION 6 : ACCIDENTAL RELEASE MEASURES

Personal Precautions:	Evacuate area and keep unnecessary and unprotected personnel from entering the spill area. Avoid personal contact and breathing vapors or mists. Use proper personal protective equipment as listed in Section 8.
Environmental Precautions:	Avoid runoff into storm sewers, ditches, and waterways.
Methods for containment:	Contain spills with an inert absorbent material such as soil, sand or oil dry.
Methods for cleanup:	Absorb spill with inert material (e,g., dry sand or earth), then place in a chemical waste container. After removal, flush spill area with soap and water to remove trace residue.

SECTION 7 : HANDLING and STORAGE

Handling:	When handling pharmaceutical products, avoid all contact and inhalation of vapor, mists and/or fumes. Use with adequate ventilation. Use only in accordance with directions.
Storage:	Store at 20 to 25 °C (68 to 77 °F) [See USP Controlled Room Temperature]. Avoid excessive heat. Do not freeze.
Work Practices:	Facilities storing or utilizing this material should be equipped with an eyewash facility and a safety shower.
Hygiene Practices:	Wash thoroughly after handling. Avoid contact with eyes and skin. Avoid inhaling vapor or mist.

SECTION 8: EXPOSURE CONTROLS, PERSONAL PROTECTION

Engineering Controls:	General ventilation is sufficient if this product is being used in a controlled medical setting (clinic, hospital, medical office) for its sole intended parenteral (injection) purpose. Otherwise, use appropriate engineering control such as process enclosures, local exhaust ventilation, or other engineering controls including use of a biosafety cabinet / fume hood to control airborne levels below recommended exposure limits.
Eye/Face Protection:	Chemical splash goggles. Wear a face shield also when splash hazard exist.
Skin Protection Description:	Protective laboratory coat, apron, or disposable garment recommended.
Hand Protection Description:	Wear appropriate protective gloves. Consult glove manufacturer's data for permeability data. Nitrile rubber or natural rubber gloves are recommended.
Respiratory Protection:	No personal respiratory protective equipment is normally required when this product is being used/administered by a licensed healthcare practitioner (i.e. an end-user such as a clinician / doctor / nurse) for its sole intended parenteral (injection) purpose in a controlled medical setting. The need for respiratory protection will vary according to the airborne concentrations and environmental conditions. A NIOSH approved air-purifying respirator with an organic vapor cartridge or canister may be permissible under certain circumstances. Consult the NIOSH web site (http://www.cdc.gov/niosh/npptl/topics/respirators/) for a list of respirator types and approved suppliers.
Other Protective:	Consult with local procedures for selection, training, inspection and maintenance of the personal protective equipment.

SECTION 9 : PHYSICAL and CHEMICAL PROPERTIES

Physical State:	Liquid solution.
Color:	Clear and colorless
Boiling Point:	Not established.
Melting Point:	Not established.
Solubility:	Soluble. in water.
Vapor Density:	Not established.
Vapor Pressure:	Not established.
Percent Volatile:	Not established.
pH:	5.0 - 7.5
Molecular Formula:	Mixture
Molecular Weight:	Variable.
Flash Point:	Not established.
Flash Point Method:	Not established.
Auto Ignition Temperature:	Not established.

SECTION 10 : STABILITY and REACTIVITY

Chemical Stability:	Stable under normal temperatures and pressures.
Hazardous Polymerization:	Not reported.
Conditions to Avoid:	Protect from freezing.

SECTION 11 : TOXICOLOGICAL INFORMATION

Acute Toxicity:	Eye, skin, and respiratory irritation may occur.
Heparin Sodium :	
Acute Toxicity:	LD50: IV Rat 2449 mg/kg LD50: IV Mouse 2800 mg/kg
Acute Effects:	Eye, skin, and respiratory irritation may occur.
Chronic Effects:	None known.
Citric Acid, anhydrous, USP :	
RTECS Number:	GE7350000
Skin:	Dermal - Rat LD50: >2000 mg/kg [Details of toxic effects not reported other than lethal dose value]
Ingestion:	Oral - Rat LD50: 5400 mg/kg [Details of toxic effects not reported other than lethal dose value]
Sodium Metabisulfite, NF :	
RTECS Number:	UX8225000
Skin:	Dermal - Rat LD50: >2000 mg/kg [Details of toxic effects not reported other than lethal dose value]
Ingestion:	Oral - Rat LD50: 1540 mg/kg [Details of toxic effects not reported other than lethal dose value]
Heparin Sodium :	
RTECS Number:	MI0850000
RTECS Number: Ingestion:	MI0850000 Oral - Rat LD50: >779000 iu/kg [Details of toxic effects not reported other than lethal dose value] Oral - Mouse LD50: >5 gm/kg [Details of toxic effects not reported other than lethal dose value]
	Oral - Rat LD50: >779000 iu/kg [Details of toxic effects not reported other than lethal dose value] Oral - Mouse LD50: >5 gm/kg [Details of toxic effects not reported other than lethal dose value] Intravenous Rat LD50: 391821 iu/kg [Details of toxic effects not reported other than lethal dose value] Intravenous Mouse LD50: 2800 mg/kg [Behavioral - convulsions or effect on seizure threshold] Intravenous Mouse LD50: 2800 units/kg [Blood - hemorrhage Blood - change in clotting factors] Intravenous Rat TDL0: 300 units/kg [Blood - hemorrhage Blood - change in clotting factors] Intravenous Guinea pig TDL0: 160 units/kg [Blood - change in clotting factors] Intravenous Rat TDL0: 84 ku/kg/28D (intermittent) [Musculoskeletal - other changes Nutritional and Gross Metabolic - changes in calcium Nutritional and Gross Metabolic - changes in phosphorus] Subcutaneous - Rat LD50: 46715 iu/kg [Details of toxic effects not reported other than lethal dose value] Subcutaneous - Mouse LD50: >2500 mg/kg [Details of toxic effects not reported other than lethal dose value] Subcutaneous - Human TDL0: 3600 units/kg/18D (intermittent) [Blood - hemorrhage Related to Chronic Data - death]
Ingestion:	Oral - Rat LD50: >779000 iu/kg [Details of toxic effects not reported other than lethal dose value] Oral - Mouse LD50: >5 gm/kg [Details of toxic effects not reported other than lethal dose value] Intravenous Rat LD50: 391821 iu/kg [Details of toxic effects not reported other than lethal dose value] Intravenous Mouse LD50: 2800 mg/kg [Behavioral - convulsions or effect on seizure threshold] Intravenous Mouse LD50: 2800 mg/kg [Behavioral - convulsions or effect on seizure threshold] Intravenous Rat TDLo: 300 units/kg [Blood - hemorrhage Blood - change in clotting factors] Intravenous Guinea pig TDLo: 160 units/kg [Blood - change in clotting factors] Intravenous Rat TDLo: 84 ku/kg/28D (intermittent) [Musculoskeletal - other changes Nutritional and Gross Metabolic - changes in calcium Nutritional and Gross Metabolic - changes in phosphorus] Subcutaneous - Rat LD50: 46715 iu/kg [Details of toxic effects not reported other than lethal dose value] Subcutaneous - Mouse LD50: >2500 mg/kg [Details of toxic effects not reported other than lethal dose value] Subcutaneous - Mouse LD50: >2500 mg/kg [Details of toxic effects not reported other than lethal dose value] Subcutaneous - Human TDLo: 3600 units/kg/18D (intermittent) [Blood - hemorrhage Related to
Ingestion: Other Toxicological Information:	Oral - Rat LD50: >779000 iu/kg [Details of toxic effects not reported other than lethal dose value] Oral - Mouse LD50: >5 gm/kg [Details of toxic effects not reported other than lethal dose value] Intravenous Rat LD50: 391821 iu/kg [Details of toxic effects not reported other than lethal dose value] Intravenous Mouse LD50: 2800 mg/kg [Behavioral - convulsions or effect on seizure threshold] Intravenous Mouse LD50: 2800 mg/kg [Behavioral - convulsions or effect on seizure threshold] Intravenous Rat TDLo: 300 units/kg [Blood - hemorrhage Blood - change in clotting factors] Intravenous Guinea pig TDLo: 160 units/kg [Blood - change in clotting factors] Intravenous Rat TDLo: 84 ku/kg/28D (intermittent) [Musculoskeletal - other changes Nutritional and Gross Metabolic - changes in calcium Nutritional and Gross Metabolic - changes in phosphorus] Subcutaneous - Rat LD50: 46715 iu/kg [Details of toxic effects not reported other than lethal dose value] Subcutaneous - Mouse LD50: >2500 mg/kg [Details of toxic effects not reported other than lethal dose value] Subcutaneous - Human TDLo: 3600 units/kg/18D (intermittent) [Blood - hemorrhage Related to Chronic Data - death] Intraperitoneal Mouse LD50: >2500 mg/kg [Details of toxic effects not reported other than lethal

Skin:	Administration onto the skin - Rabbit LD50: >10 gm/kg [Details of toxic effects not reported other than lethal dose value] Administration onto the skin - Rabbit Standard Draize test.: 50 mg/24H [mild] Administration onto the skin - Rabbit Standard Draize test.: 500 mg/24H [mild]
Inhalation:	Inhalation - Rat LC50: >42 gm/m3/1H [Details of toxic effects not reported other than lethal dose value]
Ingestion:	Oral - Mouse LD50: 4 gm/kg [Details of toxic effects not reported other than lethal dose value] Oral - Rat LD50: 3000 mg/kg [Details of toxic effects not reported other than lethal dose value]
Other Toxicological Information:	Intravenous Mouse LD50: 645 mg/kg [Details of toxic effects not reported other than lethal dose value] Intravenous Rabbit LDLo: 1100 mg/kg [Behavioral - convulsions or effect on seizure threshold Behavioral - muscle contraction or spasticity Cardiac - other changes] Intravenous Guinea pig LDLo: 300 mg/kg [Details of toxic effects not reported other than lethal dose value] Intravenous Mouse TDLo: 2.1 mg/kg [Vascular - other changes Blood - hemorrhage Skin and Appendages - dermatitis, irritative (after systemic exposure)] Intravenous Rabbit LDLo: 1.5 mg/kg [Vascular - other changes Blood - hemorrhage Skin and Appendages - dermatitis, irritative (after systemic exposure)] Subcutaneous Rabbit TDLo: 0.04 mg/kg [Vascular - other changes Blood - hemorrhage Skin and Appendages - dermatitis, irritative (after systemic exposure)] Subcutaneous - Rabbit TDLo: 3500 mg/kg [Behavioral - irritability] Subcutaneous - Rubuse LD50: 3 gm/kg [Details of toxic effects not reported other than lethal dose value] Subcutaneous - Guinea pig LDLo: 2160 mg/kg [Details of toxic effects not reported other than lethal dose value] Subcutaneous - Rubbit TDLo: 0.04 mg/kg [Vascular - other changes Skin and Appendages - dermatitis, irritative (after systemic exposure)] Subcutaneous - Rubbit TDLo: 0.04 mg/kg [Vascular - other changes Skin and Appendages - dermatitis, irritative (after systemic exposure)] Subcutaneous - Mouse TDLo: 1900 mg/kg [Reproductive - Effects on Embryo or Fetus - fetal death] Subcutaneous - Mouse TDLo: 2500 mg/kg [Reproductive - Specific Developmental Abnormalities - musculoskeletal system] Intraperitoneal Mouse LD50: 2600 mg/kg [Details of toxic effects not reported other than lethal dose value] Subcutaneous - Mouse TDLo: 13440 mg/kg [Reproductive - Fertility - abortion] Intraperitoneal Rat LD50: 2600 mg/kg [Details of toxic effects not reported other than lethal dose value] Intraperitoneal Rat LDLo: 3.72 gm/kg [Behavioral - tremor Behavioral - convulsions or effect on seizure threshold] Intraperit

Ecotoxicity:	No ecotoxicity data was found for the product.
Environmental Stability:	No environmental information found for this product.

SECTION 13 : DISPOSAL CONSIDERATIONS

Waste Disposal:

Dispose of in accordance with Local, State, Federal and Provincial regulations.

SECTION 14 : TRANSPORT INFORMATION

DOT Shipping Name:	Not Regulated.
DOT UN Number:	Not Regulated.
IATA Shipping Name:	Non regulated.
IATA UN Number:	Non regulated.
IMDG UN Number :	Non regulated.
IMDG Shipping Name :	Non regulated.

SECTION 15 : REGULATORY INFORMATION

Heparin Sodium :		
TSCA Inventory Status:	Listed	
Canada DSL:	Listed	
Sodium Citrate, Dihydrate, USP :		
TSCA Inventory Status:	Listed	

SECTION 16	: ADDITIONAL	INFORMATION
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HMIS Ratings: HMIS Health Hazard: 1 HMIS Fire Hazard: 0 HMIS Reactivity: 0 HMIS Personal Protection: Х SDS Creation Date: April 23, 2018 SDS Revision Date: March 06, 2024 SDS Revision Notes: Overall SDS review - no changes to formulation. Added product name synonym in Section 1. The information contained herein pertains to this material. It is the responsibility of each individual party to determine for themselves the proper means of handling and using these materials based on their purpose and intended use. Fresenius-Kabi assumes no liability resulting from the use of or reliance upon the information contained in this material safety data sheet. This material safety data sheet does not constitute the guaranty or specifications of the product. Disclaimer:

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