

SAFETY DATA SHEET

SECTION 1: IDENTIFICATION

Haloperidol Decanoate Injection Product Name:

Manufacturer Name: Fresenius Kabi USA, LLC Address: Three Corporate Drive Lake Zurich, Illinois 60047

General Phone Number: (847) 550-2300 Customer Service Phone (888) 386-1300

Number:

Health Issues Information: (800) 551-7176 SDS Creation Date: January 08, 2009 SDS Revision Date: June 10, 2015

(M)SDS Format:

SECTION 2: HAZARD(S) IDENTIFICATION

GHS Pictograms:



WARNING. Signal Word:

GHS Class:

Reproductive toxicity. Category 2.

Acute Oral Toxicity. Category 4.

Specific Target Organ Toxicity - STOT, Single Exposure SE. Category 3.

Reproductive toxicity. Effects on or via lactation.

Hazard Statements: Suspected of damaging fertility or the unborn child. Harmful if swallowed.

May cause respiratory irritation.
May cause harm to breast-fed children.

Precautionary Statements: Obtain special instructions before use.

Obtain special instructions before use.

Do not handle until all safety precautions have been read and understood.

Do not breathe dust/fume/gas/mist/vapours/spray.

Avoid breathing dust/fume/gas/mist/vapours/spray.

Avoid contact during pregnancy and while nursing.

Wash hands thoroughly after handling.

Do not eat, drink or smoke when using this product.

Use only outdoors or in a well-ventilated area.

Wear protective gloves/protective clothing/eye protection/face protection.

IF SWALLOWED: Call a POISON CENTER or doctor/physician if you feel unwell.

IF INHALED: Remove victim to fresh air and keep at rest in a position comfortable for breathing. IF exposed or concerned: Get medical advice/attention.

Call a POISON CENTER or doctor/physician if you feel unwell.

Rinse mouth.

Store in a well-ventilated place. Keep container tightly closed.

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

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. Emergency Overview:

Inhalation Ingestion Eye contact Skin Absorption. Injection.

Potential Health Effects:

Route of Exposure:

Contact with eyes may cause irritation. Eye:

Skin: May cause skin irritation.

Inhalation: May cause irritation of respiratory tract.

Inaestion: May cause irritation

Signs/Symptoms: Adverse effects from therapeutic doses include: extrapyramidal symptoms (Parkinson-like symptoms,

Adverse effects from therapeutic doses include: extrapyramidal symptoms (Parkinson-like symptoms akathisia, or dystonia), tardive dyskinesia (involuntary, dyskinetic movements), tardive dystonia, insomnia, restlessness, anxiety, euphoria, agitation, drowsiness, depression, lethargy, headache, confusion, vertigo, grand mal seizures, exacerbation of psychotic symptoms, neuroleptic malignant syndrome, tachycardia, hypotension, hypertension, mild and transient leukopenia and leukocytosis, impaired liver function, maculopapular and acneiform skin reactions, lactation, breast engorgement, anorexia, constipation, dry mouth, blurred vision, laryngospasm, broncospasm, cataracts, and visual disturbances. Occupational exposure has not been fully investigated.

Aggravation of Pre-Existing Conditions:

Individuals who are hypersensitive to this drug or have Parkinson's Disease.

SECTION 3: COMPOSITION/INFORMATION ON INGREDIENTS

Chemical Name CAS# **Ingredient Percent** EC Num.

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74050-97-8 Haloperidol Decanoate

Benzyl Alcohol 100-51-6 1.2%~%~1.2% by weight 1.2% by

Volume

Sesame Oil 526-07-8 Quantity Sufficient

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

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 Inaestion:

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: 200 °F

Flash Point Method: Cleveland closed cup. Auto Ignition Temperature: Not established. Lower Flammable/Explosive Limit: Not established. Upper Flammable/Explosive Limit: Not established.

Evacuate area of unprotected personnel. Use cold water spray to cool fire exposed containers to Fire Fighting Instructions:

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

Use extinguishing measures that are appropriate to local circumstances and the surrounding

As in any fire, wear Self-Contained Breathing Apparatus (SCBA), MSHA/NIOSH (approved or equivalent) Protective Equipment:

Hazardous Combustion

Byproducts:

Storage:

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

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 Personnel Precautions:

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.

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. Methods for cleanup:

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.

Store at controlled room temperature 20 to 25°C (68 to 77°F). [See USP Controlled Room Temperature]. Protect from light. Do not refrigerate or freeze. Retain in carton until time of use.

Work Practices: Facilities storing or utilizing this material should be equipped with an eyewash facility and a safety

shower

Wash thoroughly after handling. Avoid contact with eyes and skin. Avoid inhaling vapor or mist. Hygiene Practices:

SECTION 8: EXPOSURE CONTROLS, PERSONAL PROTECTION

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 Controls:**

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.

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Eve/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.

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. Consult with local procedures for selection, training, inspection and maintenance of the personal

EXPOSURE GUIDELINES

Other Protective:

SECTION 9: PHYSICAL and CHEMICAL PROPERTIES

Physical State: Liquid solution.

Color: Pale Yellow light amber

Boiling Point: Not established. Melting Point: Not established.

Solubility: Insoluble in water.(0.01 mg/ml). Soluble in most organic solvents.

Vapor Density: Not established. Vapor Pressure: Not established. Percent Volatile: Not established. 3.0 - 3.8 pH: Molecular Formula: Mixture

530.12 Molecular Weight: 200 °F Flash Point:

Flash Point Method: Cleveland closed cup. 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 light, heat, and freezing. Do not refrigerate.

Incompatible Materials: Avoid contact with oxidizing agents

SECTION 11: TOXICOLOGICAL INFORMATION

Acute Toxicity: Eye, skin, and respiratory irritation may occur. Moderately toxic by ingestion.

Haloperidol Decanoate:

Acute Toxicity: LD50 IP Rat: 27 mg/kg

LD50 IP Kat: 27 mg/kg LD50 IP Mouse: 30 mg/kg LD50 SC Mouse: 41 mg/kg LD50 IV Mouse: 15 mg/kg TDLO Multiple Man: 343 mcg/kg LD50 IV Dog: 18 mg/kg

Acute Effects: Eye, skin, and respiratory irritation may occur. Moderately toxic by ingestion.

Chronic Effects: Hypersensitivity reactions ranging from mild to severe may occur.

Haloperidol Decanoate:

RTECS Number: HD9350000

Ingestion:

Oral - Rat LD50: 1717 mg/kg [Sense Organs and Special Senses (Eye) - Lacrimation Sense Organs and Special Senses (Eye) - Ptosis Behavioral - Somnolence (general depressed activity)]
Oral - Mouse LD50: 739 mg/kg [Sense Organs and Special Senses (Eye) - Ptosis Behavioral - Somnolence (general depressed activity)]

Other Toxicological Information:

Subcutaneous - Rat LD50: 780 mg/kg [Sense Organs and Special Senses (Eye) - lacrimation Sense Organs and Special Senses (Eye) - ptosis Behavioral - somnolence (general depressed activity)]
Subcutaneous - Mouse LD50: 1990 mg/kg [Sense Organs and Special Senses (Eye) - ptosis Behavioral - somnolence (general depressed activity)]
Intraperitoneal. - Rat LD50: 328 mg/kg [Sense Organs and Special Senses (Eye) - lacrimation Sense Organs and Special Senses (Eye) - ptosis Behavioral - somnolence (general depressed activity)] Intraperitoneal. - Mouse LD50: 288 mg/kg [Sense Organs and Special Senses (Eye) - ptosis Behavioral - somnolence (general depressed activity) Gastrointestinal - hypermotility, diarrhea]

Benzyl Alcohol:

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DN3150000 RTECS Number:

Administration onto the skin - Rabbit LD50: 2000 mg/kg [Details of toxic effects not reported other Skin:

than lethal dose value]

Administration onto the skin - Rabbit Standard Draize test.: 100 mg/24H

Administration onto the skin - Rat LD50: 100 pph/90M [Details of toxic effects not reported other than

lethal dose value]

Inhalation: Inhalation - Mouse LC50: >500 mg/m3 [Behavioral - Somnolence (general depressed activity)

Behavioral - Ataxia Lungs, Thorax, or Respiration - Respiratory depression]
Inhalation - Rat LC50: >500 mg/m3 [Behavioral - Somnolence (general depressed activity) Behavioral

- Ataxia Lungs, Thorax, or Respiration - Respiratory depression]

Oral - Rat LD50: 1230 mg/kg [Behavioral - Somnolence (general depressed activity) Behavioral - Excitement Behavioral - Coma]
Oral - Mouse LD50: 1360 mg/kg [Details of toxic effects not reported other than lethal dose value] Ingestion:

Oral - Mouse LD50: 1360 mg/kg [Behavioral - Somnolence (general depressed activity) Behavioral - Ataxia Lungs, Thorax, or Respiration - Respiratory depression]

Oral - Rat LD50: 1660 mg/kg [Behavioral - Somnolence (general depressed activity) Behavioral - Ataxia Lungs, Thorax, or Respiration - Respiratory depression]

Oral - Rat LD50: 1.5 mL/kg [Details of toxic effects not reported other than lethal dose value]

Other Toxicological Information:

Intravenous. - Rat LD50: 53 mg/kg [Lungs, Thorax, or Respiration - dyspnea] Intravenous. - Mouse LD50: 324 mg/kg [Details of toxic effects not reported other than lethal dose

Subcutaneous - Rat LDLo: 1700 mg/kg [Sense Organs and Special Senses (Eye) - miosis (pupillary constriction) Behavioral - coma Kidney/Ureter/Bladder - other changes]

Intraperitoneal. - Rat LD50: 400 mg/kg [Details of toxic effects not reported other than lethal dose

value]

Intraperitoneal. - Mouse LD50: 650 mg/kg [Behavioral - altered sleep time (including change in righting reflex) Behavioral - somnolence (general depressed activity) Lungs, Thorax, or Respiration -

Intraperitoneal. - Rat LDLo: 650 mg/kg [Behavioral - somnolence (general depressed activity) Behavioral - ataxia Lungs, Thorax, or Respiration - respiratory depression] Intraperitoneal. - Rat TDLo: 514 mg/kg [Behavioral - ataxia]

SECTION 12: ECOLOGICAL INFORMATION

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.

SECTION 15: REGULATORY INFORMATION

Haloperidol Decanoate:

277-679-7 EINECS Number: Canada DSL: Listed

Benzyl Alcohol:

TSCA Inventory Status: Listed EINECS Number: 202-859-9 Canada DSL: Listed

Canada IDL: Identified under the Canadian Hazardous Products Act Ingredient Disclosure List: 0.1%.169(170)

SECTION 16: ADDITIONAL INFORMATION

HMIS Ratings:

HMIS Health Hazard: 1 HMIS Fire Hazard: 2 HMIS Reactivity: 1 HMIS Personal Protection:

SDS Creation Date: January 08, 2009 SDS Revision Date: June 10, 2015

SDS Format:

Disclaimer:

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