

ATROPINE SULFATE - atropine sulfate injection, solution
Sparhawk Laboratories, Inc.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

ATROPINE SULFATE LA 15 MG

INJECTABLE SOLUTION

FOR ANIMAL USE ONLY

KEEP OUT OF REACH OF CHILDREN

CAUTION: Federal law restricts this drug to use by or on the order of a licensed veterinarian.

Each mL contains:

Atropine Sulfate 15 mg
Sodium Chloride 9 mg
Benzyl Alcohol 1%
Water for Injection q.s.

INDICATIONS:

For use as an antidote in the treatment of organophosphate insecticide poisoning of cattle, horses and sheep.

WARNING:

Poisonous alkaloid. Keep out of reach of children. Antidotes: warmth, emetics, cholinergics

Store at controlled room temperature between 15°-30° C (59°-86°)

DOSAGE AND ADMINISTRATION; INITIAL DOSE:

Cattle: 20 mg per 100 lbs. of body weight
Horses: 6.5 mg per 100 lbs. of body weight
Sheep: 20 mg per 100 lbs. of body weight

The recommended average initial dose should be split, injecting one quarter (1/4) to one-third (1/3) slowly I.V. and the remainder I.M. or S.C. After symptoms appear to be under control, repeated maintenance doses should be administered at 3 to 6 hour intervals based on the individual response of the animal.

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Rev. 10-16

A-3007-04

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NET CONTENTS: 100 mL



MANUFACTURED BY

LENEXA, KS 66215 USA

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LOT NO.:

EXP. DATE:

ATROPINE SULFATE

atropine sulfate injection, solution

Product Information

Product Type	PRESCRIPTION ANIMAL DRUG	Item Code (Source)	NDC:58005-307
Route of Administration	INTRAVENOUS, INTRAMUSCULAR, SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ATROPINE SULFATE (UNII: 03J5ZE7KA5) (ATROPINE - UNII:7C0697DR9I)	ATROPINE SULFATE	15 mg in 1 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58005-307-04	100 mL in 1 VIAL		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		06/01/1997	

Labeler - Sparhawk Laboratories, Inc. (147979082)