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 INJECTABLE SOLUTION

(0.54 mg/mL)

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.

INDICATIONS

For use in Dogs and Cats as an antidote in the treatment of organophosphate insecticide poisoning; to reduce salivation, bronchial secretions or intestinal peristalsis associated with colic or diarrhea, and as a preanesthetic adjuvant.

DOSAGE AND ADMINISTRATION:

Dogs and Cats: Inject intravenously, intramuscularly, or subcutaneously, 1 mL for each 20 lbs of body weight as a preanesthetic adjuvant, or to reudce salivation, bronchial secretions, or intestinal peristalsis associated with colic or diarrhea.

As an antidote for parasympathomimetic drugs, inject 1 mL for each 5 lbs of body weight administered to effect and repeat as necessary. It is suggested that 1/4 of the dosage be injected intravenously and the remainder intramuscularly or subcutaneously.

WARNING

Poisonous alkaloid.

Antidotes: Warmth, Emetics, Cholinergics

COMPOSITION

Each mL of sterile solution contains:

Atropine Sulfate ..... 0.54 mg
Sodium Chloride ..... 9 mg
Benzyl Alcohol 1.5%
Water for Injection ..... QS

Store at controlled room temperature between 15º and 30ºC (59º-86ºF)

PROTECT FROM LIGHT

TAKE TIME OBSERVE LABEL DIRECTIONS

ATROPINE SULFATE

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Product Information			
Product Type	PRESCRIPTION ANIMAL DRUG	**Item Code (Source)**	NDC:58005-354
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	0.54 mg in 1 mL

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:58005-354-04	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)

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