# ATROPINE SULFATE - atropine sulfate injection, solution Covetrus North America

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.

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# ATROPINE SULFATE Injections

(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 peristals 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 15° and 30°C (59°-86°F)

### PROTECT FROM LIGHT

#### TAKE TIME OBSERVE LABEL DIRECTIONS





## ATROPINE SULFATE

atropine sulfate injection, solution

Product Information			
Product Type	PRESCRIPTION ANIMAL DRUG	Item Code (Source)	NDC:11695- 7000
Route of Administration	INTRAVENOUS, INTRAMUSCULAR, SUBCUTANEOUS		

Active Ingredient/Active Moiety		
Ingredient Name	<b>Basis of Strength</b>	Strength
ATROPINE SULFATE (UNII: 03J5ZE7KA5) (ATROPINE - UNII:7C0697DR9I)	ATROPINE SULFATE	0.54 mg in 1 mL

P	Packaging			
#	Item Code	Package Description	<b>Marketing Start Date</b>	Marketing End Date
1	NDC:11695-7000-1	100 mL in 1 VIAL		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		06/01/2021	

# Labeler - Covetrus North America (603750329)

Revised: 7/2021 Covetrus North America