

**ATROPINE- atropine sulfate solution/ drops**  
**Amneal Pharmaceuticals NY LLC**

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**HIGHLIGHTS OF PRESCRIBING INFORMATION**

**These highlights do not include all the information needed to use ATROPINE SULFATE OPHTHALMIC SOLUTION safely and effectively. See full prescribing information for ATROPINE SULFATE OPHTHALMIC SOLUTION.**

**ATROPINE SULFATE ophthalmic solution, 1%,  
for topical ophthalmic use  
Initial U.S. Approval: 1960**

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**INDICATIONS AND USAGE**  
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Atropine is an anticholinergic indicated in adults and pediatric patients aged three (3) months and older for:

- Cycloplegia (1.1)
- Mydriasis (1.2)
- Penalization of the healthy eye in the treatment of amblyopia. (1.3)

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**DOSAGE AND ADMINISTRATION**  
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- Apply 1 drop topically to the cul-de-sac of the conjunctiva, forty minutes prior to the intended maximal dilation time. (2)
- In adults and pediatric patients aged 3 years and older, doses may be repeated up to twice daily as needed. (2)

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**DOSAGE FORMS AND STRENGTHS**  
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Ophthalmic solution: 1% atropine sulfate, USP. (3)

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**CONTRAINDICATIONS**  
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Hypersensitivity or allergic reaction to any ingredient in formulation. (4.1)

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**WARNINGS AND PRECAUTIONS**  
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- Photophobia and blurred vision due to pupil unresponsiveness and cycloplegia may last up to 2 weeks. (5.1)
- Risk of blood pressure increase from systemic absorption. (5.2)

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**ADVERSE REACTIONS**  
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Most common adverse reactions that have been reported are eye pain and stinging on administration, blurred vision, photophobia, decreased lacrimation, increased heart rate and blood pressure. (6)

**To report SUSPECTED ADVERSE REACTIONS, contact Amneal Pharmaceuticals LLC at 1-877-835-5472 or FDA at 1-800-FDA-1088 or [www.fda.gov/medwatch](http://www.fda.gov/medwatch).**

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**DRUG INTERACTIONS**  
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The use of atropine and monoamine oxidase inhibitors (MAOI) is generally not recommended because of the potential to precipitate hypertensive crisis. (7)

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**USE IN SPECIFIC POPULATIONS**  
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Should only be used in pregnant women if clearly needed. (8.1)

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**Revised: 11/2025**

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\* Sections or subsections omitted from the full prescribing information are not listed.

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## **FULL PRESCRIBING INFORMATION**

### **1 INDICATIONS AND USAGE**

Atropine sulfate ophthalmic solution, 1% is indicated in adults and pediatric patients aged three (3) months and older for:

#### **1.1 Cycloplegia**

#### **1.2 Mydriasis**

#### **1.3 Penalization of the Healthy Eye in the Treatment of Amblyopia**

### **2 DOSAGE AND ADMINISTRATION**

- Apply 1 drop topically to the cul-de-sac of the conjunctiva, forty minutes prior to the

intended maximal dilation time.

- In adults and pediatric patients aged 3 years and older, doses may be repeated up to twice daily as needed.

### **3 DOSAGE FORMS AND STRENGTHS**

Ophthalmic Solution: 1% atropine sulfate, USP.

### **4 CONTRAINDICATIONS**

#### **4.1 Hypersensitivity**

Atropine sulfate ophthalmic solution, 1% is contraindicated in patients who have demonstrated a previous hypersensitivity or known allergic reaction to any ingredient of the formulation.

### **5 WARNINGS AND PRECAUTIONS**

#### **5.1 Photophobia and Blurred Vision**

Photophobia and blurred vision due to pupil unresponsiveness and cycloplegia may last up to 2 weeks.

#### **5.2 Elevation of Blood Pressure**

Elevations in blood pressure from systemic absorption has been reported following conjunctival instillation of recommended doses of atropine sulfate ophthalmic solution, 1%.

#### **5.3 Risk of Contamination**

Do not touch the dropper tip to the eye, eyelids, or any other surface as this may contaminate the solution.

### **6 ADVERSE REACTIONS**

The following serious adverse reactions are described below and elsewhere in the labeling:

- Photophobia and Blurred Vision [*see Warnings and Precautions (5.1)*]
- Elevation in Blood Pressure [*see Warnings and Precautions (5.2)*]

The following adverse reactions were identified in clinical studies or post-marketing reports following use of atropine sulfate ophthalmic solution. Because these reactions are reported voluntarily from a population of uncertain size, it is not always possible to reliably estimate their frequency or establish a causal relationship to drug exposure.

#### **6.1 Ocular Adverse Reactions**

Eye pain and stinging occurs upon instillation. Other commonly occurring adverse

reactions include, blurred vision, photophobia, superficial keratitis and decreased lacrimation. Allergic reactions such as papillary conjunctivitis, contact dermatitis and lid edema may also occur less commonly.

## **6.2 Systemic Adverse Reactions**

Systemic effects of atropine are related to its anti-muscarinic activity. Systemic adverse events reported include dryness of skin, mouth and throat from decreased secretions from mucus membranes; restlessness, irritability or delirium from stimulation of the central nervous system; tachycardia; flushed skin of the face and neck.

## **7 DRUG INTERACTIONS**

### **7.1 Monoamine Oxidase Inhibitors (MAOI)**

The use of atropine and monoamine oxidase inhibitors (MAOI) is generally not recommended because of the potential to precipitate hypertensive crisis.

## **8 USE IN SPECIFIC POPULATIONS**

### **8.1 Pregnancy**

#### Risk Summary

There are no adequate and well-controlled studies of atropine sulfate administration in pregnant women to inform a drug-associated risk. Adequate animal development and reproduction studies have not been conducted with atropine sulfate. In humans, 1% atropine sulfate is systemically bioavailable following topical ocular administration [see *Clinical Pharmacology (12.3)*].

Atropine sulfate ophthalmic solution, 1% should only be used during pregnancy if the potential benefit justifies the potential risk to the fetus.

### **8.2 Lactation**

#### Risk Summary

There is no information to inform risk regarding the presence of atropine in human milk following ocular administration of atropine sulfate to the mother. The effects on breastfed infants and the effects on milk production are also unknown. The developmental and health benefits of breastfeeding should be considered along with the mother's clinical need for atropine sulfate and any potential adverse effects on the breastfed child from atropine sulfate.

### **8.4 Pediatric Use**

The safety and effectiveness of atropine sulfate ophthalmic solution, 1% have been established in pediatric patients aged 3 months and older. Use of atropine sulfate ophthalmic solution, 1% is supported by evidence from adequate and well-controlled trials with additional safety data from published literature.

Due to the potential for systemic absorption, the use of atropine sulfate ophthalmic solution, 1% in pediatric patients aged 3 months to 3 years should be limited to no more

than one drop per eye per day.

Use in pediatric patients younger than 3 months of age is not recommended.

### 8.5 Geriatric Use

No overall differences in safety and effectiveness have been observed between elderly and younger adult patients.

## 10 OVERDOSAGE

In the event of accidental ingestion or toxic overdosage with atropine sulfate ophthalmic solution, supportive care may include a short acting barbiturate or diazepam as needed to control marked excitement and convulsions. Large doses for sedation should be avoided because central depressant action may coincide with the depression occurring late in atropine poisoning. Central stimulants are not recommended.

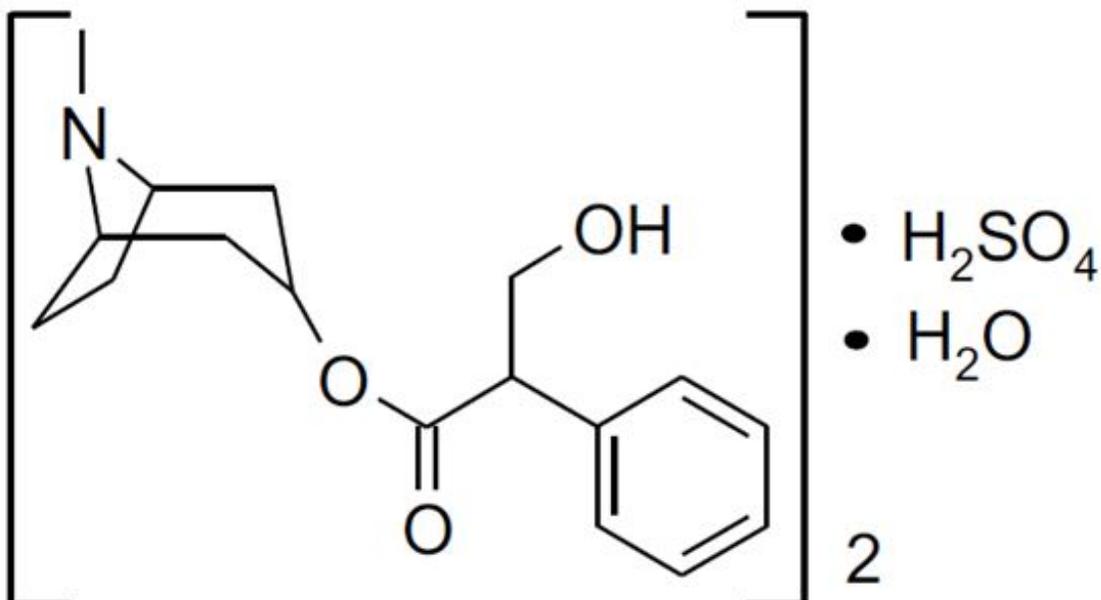
Physostigmine, given by slow intravenous injection of 1 mg to 4 mg (0.5 mg to 1 mg in pediatric populations), rapidly abolishes delirium and coma caused by large doses of atropine. Since physostigmine is rapidly destroyed, the patient may again lapse into coma after one to two hours, and repeated doses may be required.

Artificial respiration with oxygen may be necessary. Cooling measures may be needed to help to reduce fever, especially in pediatric populations.

The fatal adult dose of atropine is not known. In pediatric patients, 10 mg or less may be fatal.

## 11 DESCRIPTION

Atropine Sulfate Ophthalmic Solution USP, 1% contains atropine, an anticholinergic, in a sterile colorless, clear solution for topical ophthalmic use. The active ingredient is represented by the chemical structure:



**Chemical Name:** Benzeneacetic acid,  $\alpha$ -(hydroxymethyl)-, 8-methyl-8-azabicyclo-[3.2.1]oct-3-yl ester, *endo* -(±), sulfate (2:1) (salt), monohydrate.

**Molecular Formula:** (C<sub>17</sub>H<sub>23</sub>NO<sub>3</sub>)<sub>2</sub> • H<sub>2</sub>SO<sub>4</sub> • H<sub>2</sub>O

**Molecular Weight:** 694.83 g/mol

Each mL of atropine sulfate ophthalmic solution USP, 1% contains:

**Active:** atropine sulfate, USP 10 mg equivalent to 8.3 mg of atropine.

**Inactives:** benzalkonium chloride 0.1 mg (0.01%), dibasic sodium phosphate, edetate disodium, hypromellose (2910), monobasic sodium phosphate, hydrochloric acid and/or sodium hydroxide may be added to adjust pH (3.5 to 6.0) and water for injection USP.

## 12 CLINICAL PHARMACOLOGY

### 12.1 Mechanism of Action

Atropine is a reversible antagonist of muscarine-like actions of acetyl-choline and is therefore classified as an antimuscarinic agent. Atropine is relatively selective for muscarinic receptors. Its potency at nicotinic receptors is much lower, and actions at non-muscarinic receptors are generally undetectable clinically. Atropine does not distinguish among the M1, M2 and M3 subgroups of muscarinic receptors.

The pupillary constrictor muscle depends on muscarinic cholinceptor activation. This activation is blocked by topical atropine resulting in unopposed sympathetic dilator activity and mydriasis. Atropine also weakens the contraction of the ciliary muscle, or cycloplegia. Cycloplegia results in loss of the ability to accommodate such that the eye cannot focus for near vision.

### 12.2 Pharmacodynamics

The onset of action after administration of atropine sulfate ophthalmic solution, 1%, is usually within 40 minutes with maximal effect being reached in about 2 hours. The effect can last for up to 2 weeks in a normal eye.

### 12.3 Pharmacokinetics

The bioavailability of atropine sulfate ophthalmic solution, 1% was assessed in six healthy subjects, 24 to 29 years of age. Subjects received either 0.3 mg atropine sulfate administered as bolus intravenous injection or 0.3 mg administered as 30  $\mu$ L instilled unilaterally in the cul-de-sac of the eye. Plasma l-hyoscyamine concentrations were determined over selected intervals up to eight hours after dose administration.

The mean bioavailability of topically applied atropine was  $63.5 \pm 29\%$  (range 19% to 95%) with large inter-individual differences. Mean maximum observed plasma concentration for the ophthalmic solution was  $288 \pm 73$  pg/mL. Maximum concentration was reached in  $28 \pm 27$  min after administration. Terminal half-life of l-hyoscamine was not affected by route of administration and was calculated to be  $3 \pm 1.2$  hours (intravenous) and  $2.5 \pm 0.8$  hours (topical ophthalmic).

In another placebo-controlled study, the systemic exposure to l-hyoscyamine, and the anti-cholinergic effects of atropine were investigated in eight ocular surgery patients 56

to 66 years of age, following single topical ocular 0.4 mg atropine dose (given as 40 microliters of atropine sulfate ophthalmic solution, 1%). The mean ( $\pm$  standard deviation (SD))  $C_{\max}$  of l-hyoscyamine in these patients was  $860 \pm 402$  pg/mL, achieved within 8 minutes of eyedrop instillation.

Following intravenous administration, the mean ( $\pm$  SD) elimination half-life ( $t_{1/2}$ ) of atropine was reported to be longer in pediatric subjects under 2 years ( $6.9 \pm 3.3$  hours) and in geriatric patients 65 to 75 years ( $10.0 \pm 7.3$  hours), compared to in children over 2 years ( $2.5 \pm 1.2$  hours) and in adults 16 to 58 years ( $3.0 \pm 0.9$  hours) [see *Use in Specific Populations (8.4)*].

Atropine is destroyed by enzymatic hydrolysis, particularly in the liver; from 13% to 50% is excreted unchanged in the urine. Traces are found in various secretions, including milk. The major metabolites of atropine are noratropine, atropin-n-oxide, tropine and tropic acid. Atropine readily crosses the placental barrier and enters the fetal circulation, but is not found in amniotic fluid.

Atropine binds poorly (about 44%) to plasma protein, mainly to  $\alpha$ -1 acid glycoprotein; age has no effect on the serum protein binding of atropine. Atropine binding to  $\alpha$ -1 acid glycoprotein was concentration dependent (2 mcg/mL to 20 mcg/mL) and nonlinear *in vitro* and *in vivo*. There is no gender effect on the pharmacokinetics of atropine administered by injection.

## **13 NONCLINICAL TOXICOLOGY**

### **13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility**

#### Carcinogenesis

Long term studies in animals have not been performed to evaluate the carcinogenic potential of atropine sulfate.

#### Mutagenesis

Atropine sulfate was negative in the salmonella/microsome mutagenicity test.

#### Impairment of Fertility

Studies to evaluate impairment of fertility have not been conducted.

## **14 CLINICAL STUDIES**

Topical administration of atropine sulfate ophthalmic solution, 1% results in cycloplegia and mydriasis which has been demonstrated in several controlled clinical studies in adults and pediatric patients. Maximal mydriasis usually occurs in about 40 minutes and maximal cycloplegia is usually achieved in about 60 to 90 minutes after single administration. Full recovery usually occurs in approximately one week, but may last a couple of weeks.

## **16 HOW SUPPLIED/STORAGE AND HANDLING**

Atropine Sulfate Ophthalmic Solution USP, 1% is supplied sterile in white opaque low density polyethylene plastic bottles and natural low density polyethylene tips with red

high density polyethylene caps in the following sizes:

2 mL in 5 mL Container: NDC 60219-1748-2

5 mL in 5 mL Container: NDC 60219-1749-3

15 mL in 15 mL Container: NDC 60219-1750-8

**Storage:** Store at 20° to 25°C (68° to 77°F) [See USP Controlled Room Temperature]. Keep tightly closed. After opening, Atropine Sulfate Ophthalmic Solution can be used until the expiration date on the bottle.

## **17 PATIENT COUNSELING INFORMATION**

### Risk of Contamination

Advise patients to not touch the dropper tip to the eye, eyelids, or any other surface as this may contaminate the solution. Keep the bottle tightly closed.

### Instillation Site Pain

Advise patients that drops will sting upon instillation.

### Sensitivity to Light and Blurred Vision

Advise patients that they will experience sensitivity to light and blurred vision which may last for a couple of weeks.

Manufactured by:

**Amneal Pharmaceuticals Pvt. Ltd.**

Ahmedabad 382213, INDIA

Distributed by:

**Amneal Pharmaceuticals LLC**

Bridgewater, NJ 08807

Rev. 11-2025-01

## **PRINCIPAL DISPLAY PANEL**

**NDC 60219-1748-2**

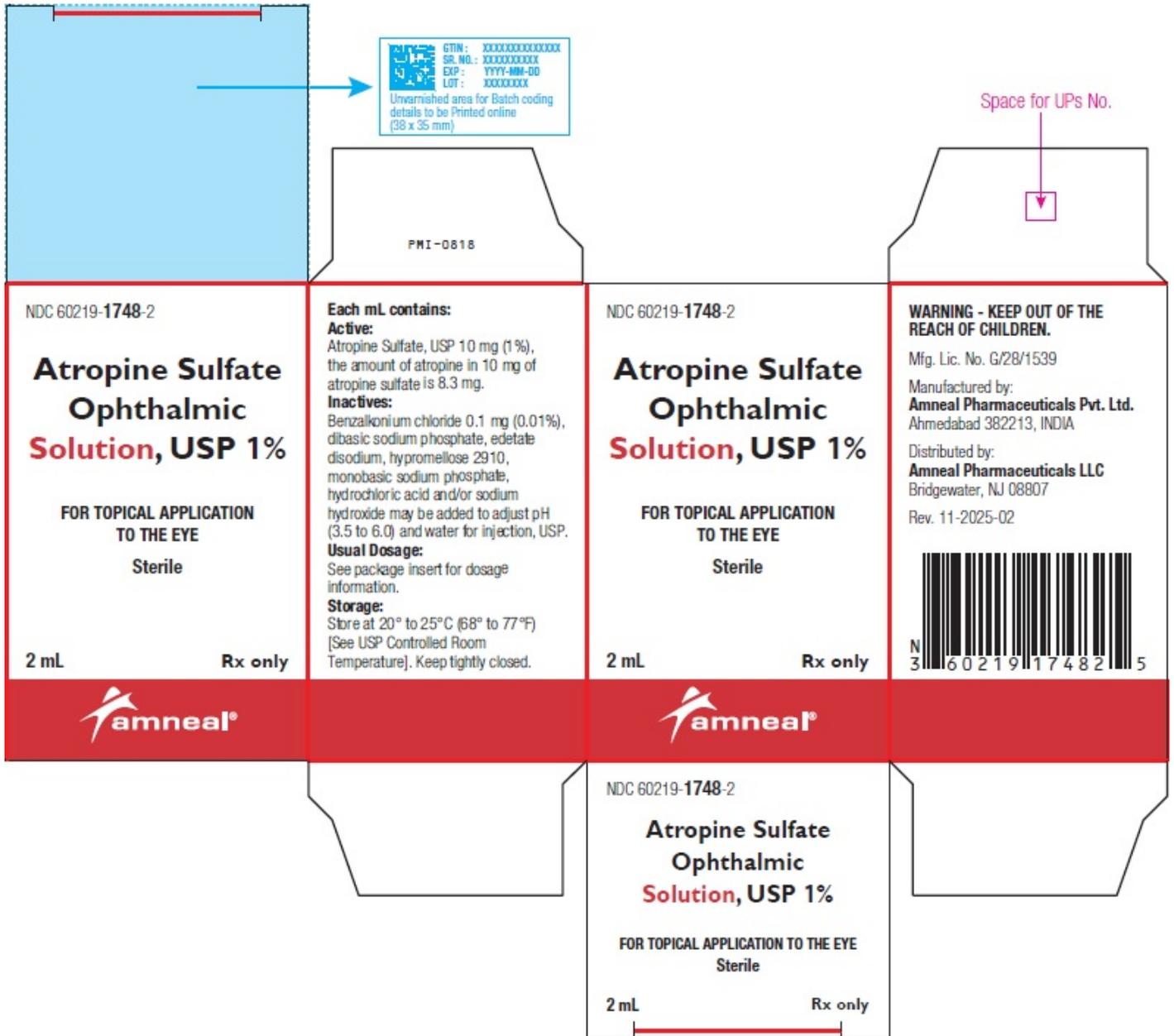
**Atropine Sulfate Ophthalmic Solution USP, 1%**

**2 mL**

**Rx only**

**Amneal Pharmaceuticals LLC**

Unvarnished Area for, (6 x 14 mm)	NDC 60219-1748-2 <b>Rx only</b>	<b>Each mL contains:</b>	 Made in INDIA Rev. 04-2024-03 Mfg. Lic. No. G/28/1539 (01) 0 0360219 17482 5	Unvarnished Area for, Lot. & Exp. (10 x 14 mm)
	<b>Atropine Sulfate Ophthalmic Solution, USP 1%</b> FOR TOPICAL APPLICATION TO THE EYE 2 mL <b>Sterile</b>	<b>Active:</b> Atropine Sulfate, USP 10 mg (1%), the amount of atropine in 10 mg of atropine sulfate is 8.3 mg. <b>Preservative:</b> Benzalkonium Chloride 0.1 mg (0.01%). <b>Storage:</b> Store at 20° to 25°C (68° to 77°F) [See USP Controlled Room Temperature]. Keep tightly closed. <b>WARNING - KEEP OUT OF THE REACH OF CHILDREN.</b> Distributed by: <b>Amneal Pharmaceuticals LLC</b>		



**NDC 60219-1749-3**  
**Atropine Sulfate Ophthalmic Solution USP, 1%**  
**5 mL**  
**Rx only**  
**Amneal Pharmaceuticals LLC**

Unvarnished Area for, (6 x 14 mm)

NDC 60219-1749-3    Rx only

**Atropine Sulfate Ophthalmic Solution, USP 1%**

FOR TOPICAL APPLICATION TO THE EYE

5 mL    Sterile

**Each mL contains:**  
**Active:** Atropine Sulfate, USP 10 mg (1%), the amount of atropine in 10 mg of atropine sulfate is 8.3 mg.  
**Preservative:** Benzalkonium Chloride 0.1 mg (0.01%).  
**Storage:** Store at 20° to 25°C (68° to 77°F) [See USP Controlled Room Temperature]. Keep tightly closed.  
**WARNING - KEEP OUT OF THE REACH OF CHILDREN.**  
 Distributed by: Amneal Pharmaceuticals LLC

Made in INDIA  
 Rev. 04-2024-03  
 Mfg. Lic. No. G/28/1539

Unvarnished Area for, Lot. & Exp. (10 x 14 mm)

Unvarnished Area for, (6 x 14 mm)

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 EXP : YYYY-MM-DD  
 LOT : XXXXXXXXX

Unvarnished area for Batch coding details to be Printed online (38 x 35 mm)

PMI-0820

Space for UPs No.

NDC 60219-1749-3

**Atropine Sulfate Ophthalmic Solution, USP 1%**

FOR TOPICAL APPLICATION TO THE EYE

Sterile

5 mL    Rx only

**Each mL contains:**  
**Active:** Atropine Sulfate, USP 10 mg (1%), the amount of atropine in 10 mg of atropine sulfate is 8.3 mg.  
**Inactives:** Benzalkonium chloride 0.1 mg (0.01%), dibasic sodium phosphate, edetate disodium, hypromellose 2910, monobasic sodium phosphate, hydrochloric acid and/or sodium hydroxide may be added to adjust pH (3.5 to 6.0) and water for injection, USP.  
**Usual Dosage:** See package insert for dosage information.  
**Storage:** Store at 20° to 25°C (68° to 77°F) [See USP Controlled Room Temperature]. Keep tightly closed.

NDC 60219-1749-3

**Atropine Sulfate Ophthalmic Solution, USP 1%**

FOR TOPICAL APPLICATION TO THE EYE

Sterile

5 mL    Rx only

**WARNING - KEEP OUT OF THE REACH OF CHILDREN.**

Mfg. Lic. No. G/28/1539

Manufactured by:  
**Amneal Pharmaceuticals Pvt. Ltd.**  
 Ahmedabad 382213, INDIA

Distributed by:  
**Amneal Pharmaceuticals LLC**  
 Bridgewater, NJ 08807

Rev. 11-2025-02

NDC 60219-1749-3

**Atropine Sulfate Ophthalmic Solution, USP 1%**

FOR TOPICAL APPLICATION TO THE EYE

Sterile

5 mL    Rx only

**NDC 60219-1750-8**  
**Atropine Sulfate Ophthalmic Solution USP, 1%**  
**15 mL**  
**Rx only**  
**Amneal Pharmaceuticals LLC**

NDC 60219-1750-8

Rx only

Each mL contains:

**Active:** Atropine Sulfate, USP 10 mg (1%),  
the amount of atropine in 10 mg of  
atropine sulfate is 8.3 mg.

**Preservative:** Benzalkonium Chloride  
0.1 mg (0.01%).

**Usual Dosage:** See package insert for  
dosage information.

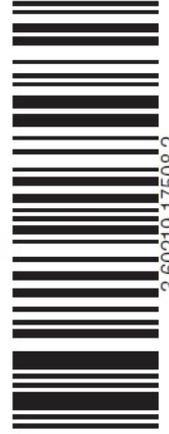
**Storage:** Store at 20° to 25°C (68° to 77°F)  
[See USP Controlled Room Temperature].  
Keep tightly closed.

**WARNING - KEEP OUT OF THE REACH OF  
CHILDREN.**

Mfg. Lic. No. G/28/1539

Distributed by: **Amneal Pharmaceuticals LLC**  
Made in INDIA.

Rev. 04-2024-03



Unvarnished Area for,  
Lot. & Exp.  
(10 x 28 mm)

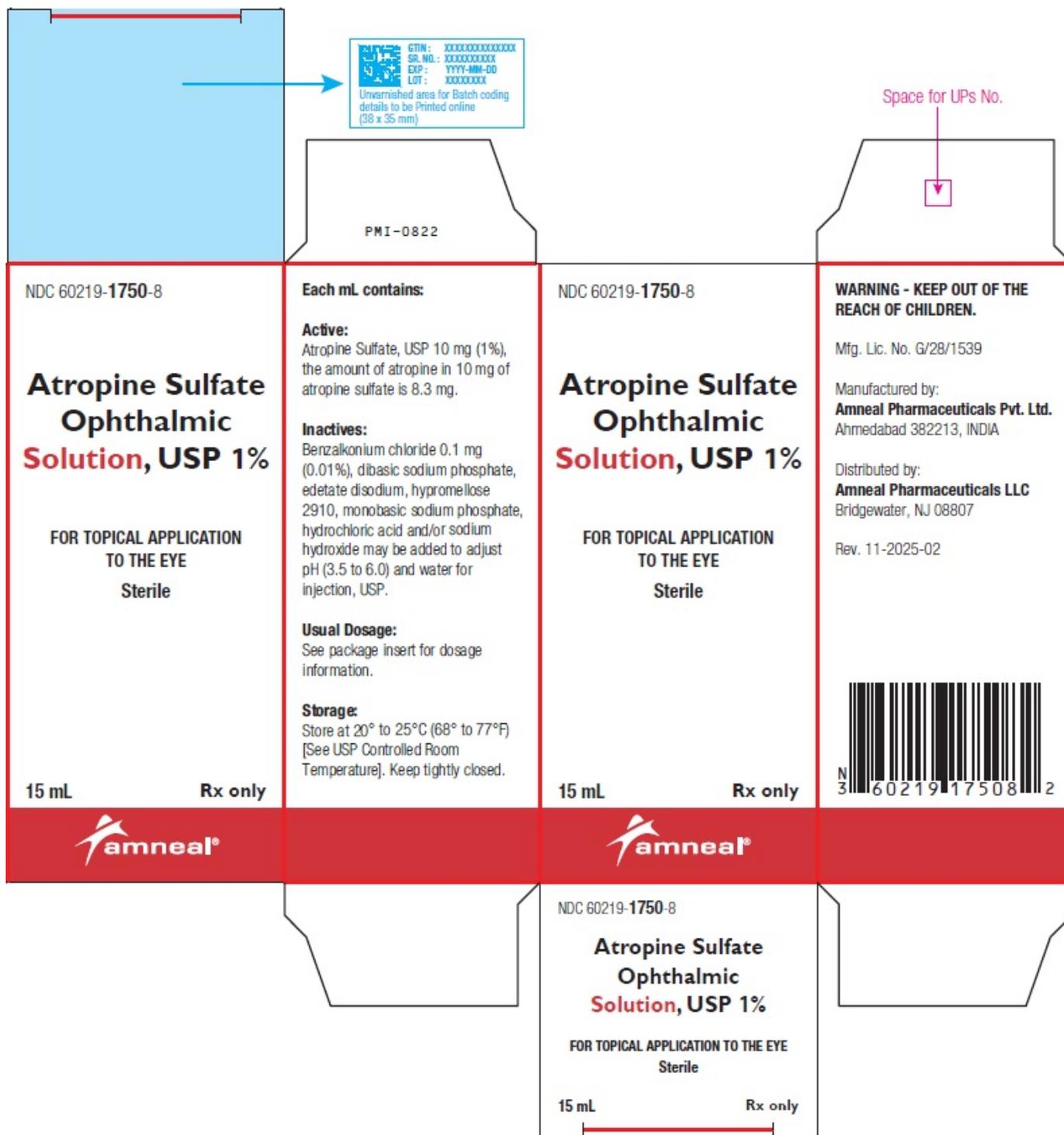
Unvarnished Area for,  
(6 x 28 mm)

# Atropine Sulfate Ophthalmic Solution, USP 1%

**FOR TOPICAL APPLICATION  
TO THE EYE**

**15 mL**

**Sterile**



## ATROPINE

atropine sulfate solution/ drops

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:60219-1748
<b>Route of Administration</b>	OPHTHALMIC		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>ATROPINE SULFATE</b> (UNII: 03J5ZE7KA5) (ATROPINE - UNII:7C0697DR9I)	ATROPINE SULFATE	10 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7)	
<b>SODIUM PHOSPHATE, DIBASIC</b> (UNII: GR686LBA74)	
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	
<b>HYPROMELLOSE 2910 (4000 MPA.S)</b> (UNII: RN3152OP35)	
<b>SODIUM PHOSPHATE, MONOBASIC</b> (UNII: 3980JIH2SW)	
<b>HYDROCHLORIC ACID</b> (UNII: QTT17582CB)	
<b>SODIUM HYDROXIDE</b> (UNII: 55X04QC32I)	
<b>WATER</b> (UNII: 059QF0KO0R)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60219-1748-2	1 in 1 CARTON	07/19/2022	
1		2 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA214752	07/19/2022	

## ATROPINE

atropine sulfate solution/ drops

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:60219-1749
<b>Route of Administration</b>	OPHTHALMIC		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>ATROPINE SULFATE</b> (UNII: 03J5ZE7KA5) (ATROPINE - UNII:7C0697DR9I)	ATROPINE SULFATE	10 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7)	

<b>SODIUM PHOSPHATE, DIBASIC</b> (UNII: GR686LBA74)
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)
<b>HYPROMELLOSE 2910 (4000 MPA.S)</b> (UNII: RN3152OP35)
<b>SODIUM PHOSPHATE, MONOBASIC</b> (UNII: 3980JH2SW)
<b>HYDROCHLORIC ACID</b> (UNII: QTT17582CB)
<b>SODIUM HYDROXIDE</b> (UNII: 55X04QC32I)
<b>WATER</b> (UNII: 059QF0KO0R)

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60219-1749-3	1 in 1 CARTON	07/19/2022	
1		5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA214752	07/19/2022	

## ATROPINE

atropine sulfate solution/ drops

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:60219-1750
<b>Route of Administration</b>	OPHTHALMIC		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>ATROPINE SULFATE</b> (UNII: 03J5ZE7KA5) (ATROPINE - UNII:7C0697DR9I)	ATROPINE SULFATE	10 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7)	
<b>SODIUM PHOSPHATE, DIBASIC</b> (UNII: GR686LBA74)	
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	
<b>HYPROMELLOSE 2910 (4000 MPA.S)</b> (UNII: RN3152OP35)	
<b>SODIUM PHOSPHATE, MONOBASIC</b> (UNII: 3980JH2SW)	
<b>HYDROCHLORIC ACID</b> (UNII: QTT17582CB)	
<b>SODIUM HYDROXIDE</b> (UNII: 55X04QC32I)	
<b>WATER</b> (UNII: 059QF0KO0R)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60219-1750-8	1 in 1 CARTON	07/19/2022	
1		15 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA214752	07/19/2022	

**Labeler** - Amneal Pharmaceuticals NY LLC (123797875)

## Establishment

Name	Address	ID/FEI	Business Operations
Amneal Pharmaceuticals Private Limited		860156658	analysis(60219-1748, 60219-1749, 60219-1750) , manufacture(60219-1748, 60219-1749, 60219-1750) , pack(60219-1748, 60219-1749, 60219-1750) , sterilize(60219-1748, 60219-1749, 60219-1750)

Revised: 12/2025

Amneal Pharmaceuticals NY LLC