

**OLOPATADINE HYDROCHLORIDE - olopatadine hydrochloride solution**  
**Amerisource Bergen**

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***Drug Facts***

***Active ingredient***

Olopatadine (0.2%)  
(equivalent to olopatadine hydrochloride, USP 0.222%)

**Purpose**

Antihistamine

***Use***

temporarily relieves itchy eyes due to pollen, ragweed, grass, animal hair and dander

***Warnings***

**For external use only**

**Do not use**

- if solution changes color or becomes cloudy
- if you are sensitive to any ingredient in this product
- to treat contact lens related irritation

**When using this product**

- do not touch tip of container to any surface to avoid contamination
- remove contact lenses before use
- wait at least 10 minutes before reinserting contact lenses after use
- do not wear a contact lens if your eye is red

**Stop use and ask a doctor if you experience:**

- eye pain
- changes in vision
- increased redness of the eye
- itching worsens or lasts for more than 72 hours

**Keep out of reach of children.**

If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

**Directions**

- **adults and children 2 years of age and older:**
  - put 1 drop in the affected eye(s) once daily, no more than once per day
  - if using other ophthalmic products while using this product, wait at least 5 minutes between each product
  - replace cap after each use
- **children under 2 years of age:**

consult a doctor

**Other information**

- only for use in the eye
- store between 2° to 25°C (36° to 77°F)

**Inactive ingredients**

benzalkonium chloride 0.01%, dibasic sodium phosphate, edetate disodium, hydrochloric acid/sodium hydroxide (adjust pH), povidone, sodium chloride and water for injection

**Questions?**

©1-855-274-4122

Distributed By  
AmerisourceBergen  
1 West First Avenue  
Conshohocken, PA 1942

Made in India  
Code: TS/DRUGS/13/2010

**PACKAGE LABEL-PRINCIPAL DISPLAY PANEL-0.2% (2.5 mL Container)**

**GOOD  
NEIGHBOR  
PHARMACY®**

**NDC 46122-671-27**

**Olopatadine Hydrochloride  
Ophthalmic Solution USP, 0.2%  
Antihistamine**

***Eye Allergy Relief***

**STERILE**

**2.5 mL (0.085 FL OZ)**



NDC 46122-671-27

**ONCE DAILY**

Only for use in the eye.  
Store between 2° to 25°C (36° to 77°F)  
**TAMPER EVIDENT:**  
Do not use if ring at bottom of cap is broken or missing.

Distributed By: AmerisourceBergen  
1 West First Avenue Conshohocken, PA 19428  
Questions or Concerns?

[www.mygnp.com](http://www.mygnp.com) LM-4643 P1428603

**Olopatadine Hydrochloride  
Ophthalmic Solution USP, 0.2%**

Antihistamine

**Eye Allergy Relief**

STERILE

2.5 mL (0.085 FL OZ)

Code: TS/DRUGS/13/2010  
Made in India



**PACKAGE LABEL-PRINCIPAL DISPLAY PANEL-0.2% (2.5 mL Container Carton)**

\*Compare to the Active Ingredient  
in Pataday® Once Daily Relief

**GOOD  
NEIGHBOR  
PHARMACY®**

**NDC 46122-671-27**

**NOW AVAILABLE without a prescription**

**Olopatadine Hydrochloride  
Ophthalmic Solution, USP  
0.2%**

**Antihistamine**

***Eye Allergy Relief***

**Works in Minutes**

**Relief from Allergens:**

- Pet Dander • Pollen
- Grass • Ragweed

**ONCE  
DAILY**

**STERILE**

**2.5 mL (0.085 FL OZ)**



## OLOPATADINE HYDROCHLORIDE

olopatadine hydrochloride solution

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:46122-671
<b>Route of Administration</b>	OPHTHALMIC		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>OLOPATADINE HYDROCHLORIDE</b> (UNII: 2XG66W44KF) (OLOPATADINE - UNII:D27V6190PM)	OLOPATADINE HYDROCHLORIDE	2 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
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<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7)	
<b>SODIUM PHOSPHATE, DIBASIC, UNSPECIFIED FORM</b> (UNII: GR686LBA74)	
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	
<b>HYDROCHLORIC ACID</b> (UNII: QTT17582CB)	
<b>SODIUM HYDROXIDE</b> (UNII: 55X04QC32I)	
<b>POVIDONE, UNSPECIFIED</b> (UNII: FZ989GH94E)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>WATER</b> (UNII: 059QF0KO0R)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:46122-671-27	1 in 1 CARTON	01/19/2021	
1		2.5 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA209995	01/19/2021	

**Labeler** - Amerisource Bergen (007914906)

**Registrant** - Aurobindo Pharma Limited (650082092)

### Establishment

Name	Address	ID/FEI	Business Operations
Eugia Pharma Specialities Limited		650498244	ANALYSIS(46122-671) , MANUFACTURE(46122-671)

Revised: 7/2023

Amerisource Bergen