

**OLOPATADINE HYDROCHLORIDE - olopatadine hydrochloride solution**  
**Aurohealth LLC**

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***Drug Facts***

***Active ingredient***

Olopatadine (0.1%).  
(equivalent to olopatadine hydrochloride, USP 0.111%)

***Purpose***

Antihistamine and Redness Reliever

***Uses***

temporarily relieves itchy and red 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) twice daily, every 6 to 8 hours, no more than twice 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 4° to 25°C (39° to 77°F)

**Inactive ingredients**

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

**Questions?**

☎1-855-274-4122

Distributed by:

**AUROHEALTH LLC**  
2572 Brunswick Pike  
Lawrenceville, NJ 08648

Made in India

**PACKAGE LABEL-PRINCIPAL DISPLAY PANEL-0.1% (5 mL Container)**

**Healthy Living™** NDC 58602-009-40

**Olopatadine Hydrochloride**

**Ophthalmic Solution, USP**

**0.1%**

Antihistamine and Redness Reliever

**Eye Allergy Itch & Redness Relief**

**STERILE 5 mL (0.17 FL OZ)**

 **Healthy Living™** NDC 58602-009-40

**Olopatadine Hydrochloride**

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**Eye Allergy Itch & Redness Relief**

**STERILE 5 mL (0.17 FL OZ)**

**TWICE DAILY**

Only for use in the eye.

Store between 4° to 25°C (39° to 77°F)

**TAMPER EVIDENT:**

Do not use if ring at

bottom of cap is

broken or missing.

Distributed by: **AUROHEALTH LLC**, 2572 Brunswick Pike,  
Lawrenceville, NJ 08648

Made in India

Code: TS/DRUGS/13/2010

LM-4256

P1426561



**PACKAGE LABEL-PRINCIPAL DISPLAY PANEL-0.1% (5 mL Container Carton)**

**Healthy Living™** NDC 58602-009-40

\*Compare to the Active Ingredient in

Pataday® Twice Daily Relief

**NOW AVAILABLE without a prescription**

**Olopatadine Hydrochloride**

**Ophthalmic Solution, USP**

**0.1%**

Antihistamine and Redness Reliever  
**Eye Allergy Itch & Redness Relief**

Works in Minutes  
 Relief from Allergens:

- Pet Dander • Pollen
- Grass • Ragweed

**TWICE  
 DAILY  
 STERILE**

**5 mL (0.17 FL OZ)**



**OLOPATADINE HYDROCHLORIDE**

olopatadine hydrochloride solution

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:58602-009
<b>Route of Administration</b>	OPHTHALMIC		

**Active Ingredient/Active Moiety**

Ingredient Name		Basis of Strength	Strength	
OLOPATADINE HYDROCHLORIDE (UNII: 2XG66W44KF) (OLOPATADINE - UNII:D27V6190PM)		OLOPATADINE HYDROCHLORIDE	1 mg in 1 mL	
Inactive Ingredients				
Ingredient Name			Strength	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)				
SODIUM PHOSPHATE, DIBASIC, UNSPECIFIED FORM (UNII: GR686LBA74)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
HYDROCHLORIC ACID (UNII: QTT17582CB)				
SODIUM HYDROXIDE (UNII: 55X04QC32I)				
WATER (UNII: 059QF0K00R)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58602-009-40	1 in 1 CARTON	07/15/2020	
1		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	ANDA204812	07/15/2020		

**Labeler** - Aurohealth LLC (078728447)

### Establishment

Name	Address	ID/FEI	Business Operations
Aurobindo Pharma Limited		650498244	ANALYSIS(58602-009) , MANUFACTURE(58602-009)

Revised: 8/2020

Aurohealth LLC