

**OLOPATADINE HYDROCHLORIDE- olopatadine hydrochloride solution**  
**Rising Pharma Holdings, Inc.**

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

**ACTIVE INGREDIENT(S)**

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

**PURPOSE**

Antihistamine

**USE(S)**

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

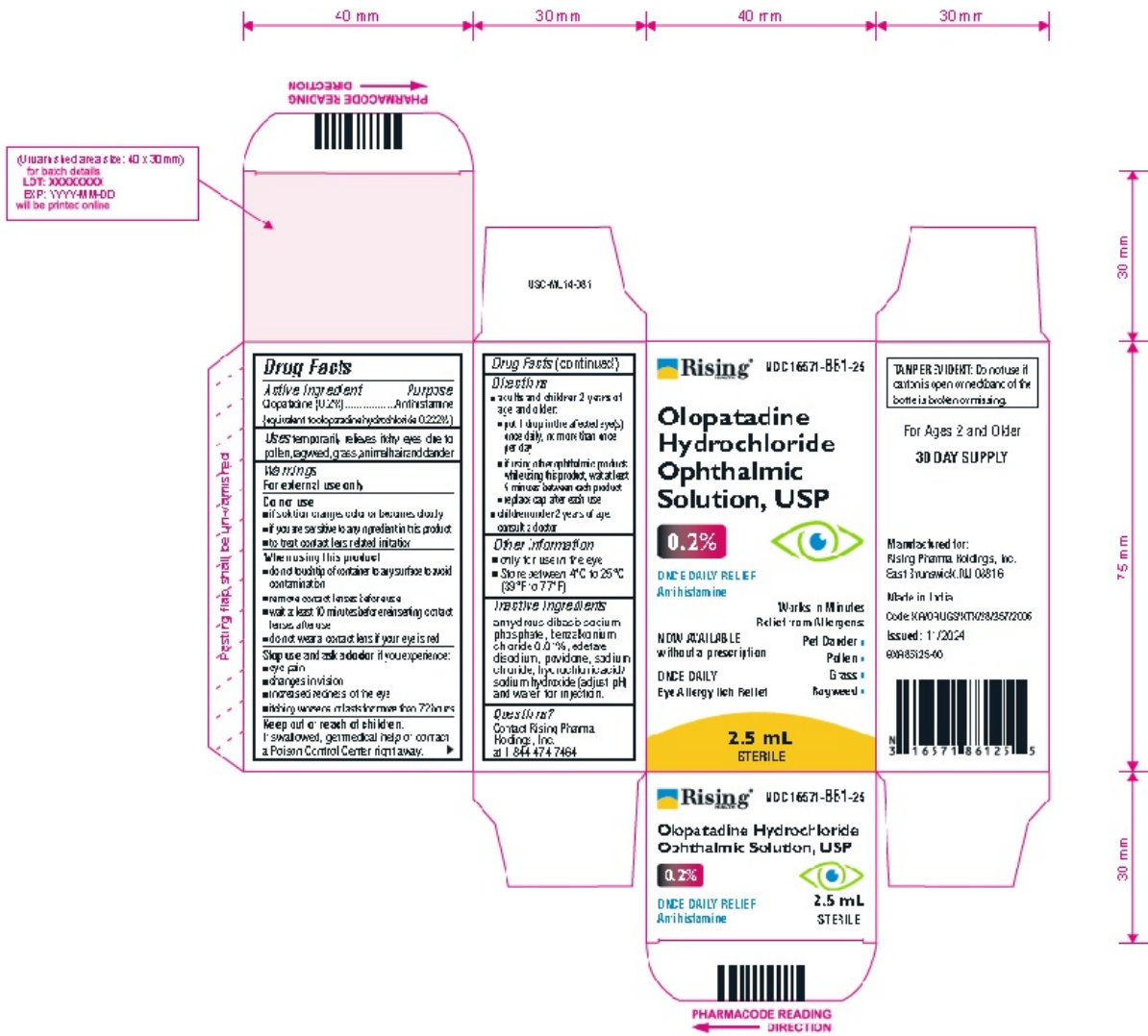
**STOP USE AND ASK DOCTOR IF**

you experience:

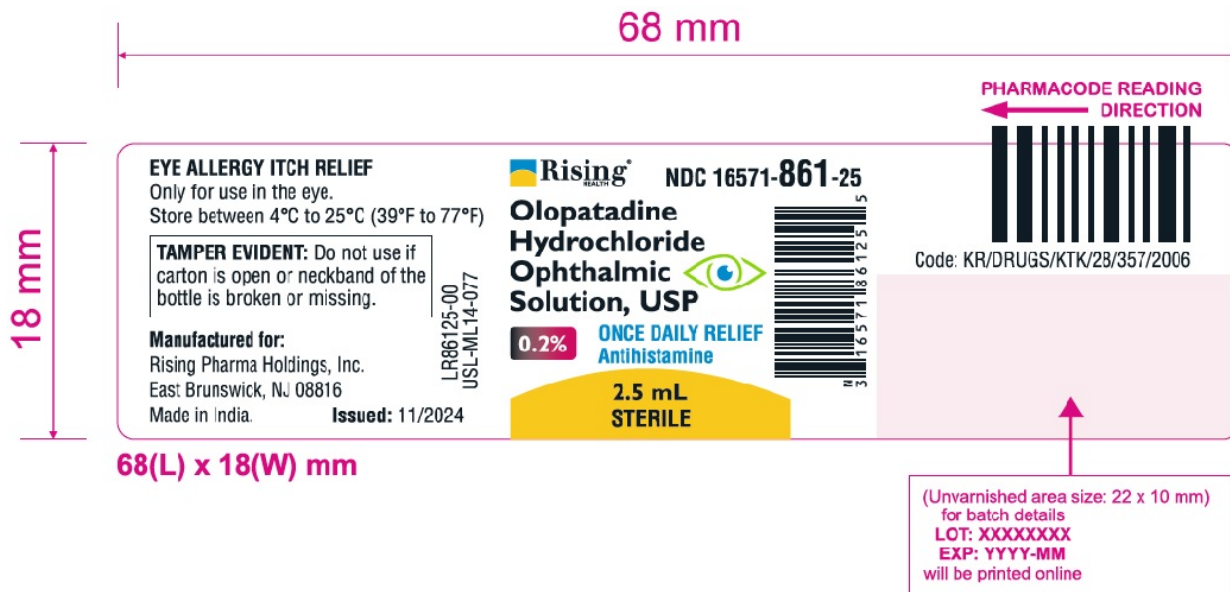
- eye pain
- changes in vision
- increased redness of the eye
- itching worsens or lasts for more than 72 hours

**KEEP OUT OF REACH OF CHILDREN**





Rising NDC 16571-861-25  
 Olopatadine Hydrochloride Ophthalmic Solution, USP  
 0.2%  
 ONCE DAILY RELIEF Antihistamine  
 Eye Allergy Itch Relief  
 ONCE DAILY  
 2.5 mL  
 STERILE



## OLOPATADINE HYDROCHLORIDE

olopatadine hydrochloride solution

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:16571-861
<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	2 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7)	
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	
<b>SODIUM PHOSPHATE, DIBASIC, ANHYDROUS</b> (UNII: 22ADO53M6F)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>HYDROCHLORIC ACID</b> (UNII: QTT17582CB)	
<b>SODIUM HYDROXIDE</b> (UNII: 55X04QC32I)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>POVIDONE K30</b> (UNII: U725QWY32X)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date

<b>1</b>	NDC:16571-861-25	1 in 1 CARTON	03/07/2025	
<b>1</b>		2.5 mL in 1 CONTAINER; Type 0: Not a Combination Product		
<b>Marketing Information</b>				
<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>		<b>Marketing Start Date</b>	<b>Marketing End Date</b>
ANDA	ANDA204620		03/07/2025	

**Labeler** - Rising Pharma Holdings, Inc. (116880195)

<b>Establishment</b>			
<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
Micro Labs Limited		677600482	ANALYSIS(16571-861) , LABEL(16571-861) , MANUFACTURE(16571-861) , PACK(16571-861)

Revised: 3/2025

Rising Pharma Holdings, Inc.