

OLOPATADINE HYDROCHLORIDE OPHTHALMIC SOLUTION- olopatadine hydrochloride ophthalmic solution
CVS Pharmacy

ACTIVE INGREDIENT

Olopatadine (0.7%) (equivalent to olopatadine hydrochloride 0.776%)

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 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 right away(1-800-222-1222).

DIRECTIONS

- **adults and children 2 years of age and older:**
- put **1 drop** in the affected eye(s) **once daily**
- **do not use more than 1 drop in each eye 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 0.015% (preservative), boric acid, hydrochloric acid/sodium hydroxide (to adjust pH), hydroxypropyl gamma cyclodextrin, hypromellose, mannitol, polyethylene glycol 400, povidone and purified water.

QUESTIONS?

call toll-free weekdays 8 AM to 8 PM EST at **1-888-375-3784**

Distributed by:

CVS Pharmacy Inc.

One CVS Drive

Woonsocket, RI 02895

Made in India

PRINCIPAL DISPLAY PANEL

Carton Label:

EXTRA STRENGTH

Olopatadine

Hydrochloride

Ophthalmic

Solution, USP

0.7%

Antihistamine

ONCE DAILY RELIEF

Eye Allergy Itch Relief
2.5 mL (0.085 fl oz)
Sterile

Placeholder Image|

Bottle Label:

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OLOPATADINE HYDROCHLORIDE OPHTHALMIC SOLUTION

olopatadine hydrochloride ophthalmic solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51316-567
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OLOPATADINE HYDROCHLORIDE (UNII: 2XG66W44KF) (OLOPATADINE - UNII:D27V6190PM)	OLOPATADINE	7 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BORIC ACID (UNII: R57ZHV85D4)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
HYDROXYPROPYL .GAMMA.-CYCLODEXTRIN (UNII: P6BYU725IU)	
HYPROMELLOSE 2910 (4000 MPA.S) (UNII: RN3152OP35)	
MANNITOL (UNII: 3OWL53L36A)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
POVIDONE K30 (UNII: U725QWY32X)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51316-567-01	1 in 1 CARTON	01/15/2026	
1		2.5 mL in 1 BOTTLE; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA213514	01/15/2026	

