

OLOPATADINE HYDROCHLORIDE- olopatadine hydrochloride solution
WALGREEN COMPANY

Walgreens Olopatadine Hydrochloride Ophthalmic Solution USP, 0.2% 3.5 mL and 2 x 3.5 mL Twin Pack

Active ingredient

Olopatadine (0.2%)

(equivalent to olopatadine hydrochloride 0.222%)

Purpose

Antihistamine

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 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 4-25 °C (39-77 °F)
- protect from light

Inactive Ingredients

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

Questions?

Call 1-800-459-6906

Package/Label Principal Display Panel

91.5



OLOPATADINE HYDROCHLORIDE

olopatadine hydrochloride solution

Product Information

| | | | |
|-------------------------|----------------|--------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:0363-0869 |
| Route of Administration | OPHTHALMIC | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|-------------------|----------|
| OLOPATADINE HYDROCHLORIDE (UNII: 2XG66W44KF) (OLOPATADINE - | OLOPATADINE | 2 mg |

| | | | | |
|--|------------------|--|----------------------|--------------------|
| UNII:D27V6190PM) | | | CLOFATADINE | in 1 mL |
| | | | | |
| Inactive Ingredients | | | | |
| Ingredient Name | | | | Strength |
| POVIDONE, UNSPECIFIED (UNII: FZ989GH94E) | | | | |
| SODIUM PHOSPHATE, DIBASIC, UNSPECIFIED FORM (UNII: GR686LBA74) | | | | |
| SODIUM CHLORIDE (UNII: 451W47IQ8X) | | | | |
| EDETATE DISODIUM (UNII: 7FLD91C86K) | | | | |
| BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) | | | | |
| HYDROCHLORIC ACID (UNII: QTT17582CB) | | | | |
| SODIUM HYDROXIDE (UNII: 55X04QC32I) | | | | |
| WATER (UNII: 059QF0KO0R) | | | | |
| | | | | |
| Packaging | | | | |
| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
| 1 | NDC:0363-0869-01 | 1 in 1 CARTON | 12/22/2025 | |
| 1 | | 3.5 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | | |
| 2 | NDC:0363-0869-02 | 2 in 1 CARTON | 12/22/2025 | |
| 2 | | 3.5 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 | | ANDA206087 | 12/22/2025 | |

Labeler - WALGREEN COMPANY (008965063)

| | | | |
|----------------------------|----------------|---------------|----------------------------|
| Establishment | | | |
| Name | Address | ID/FEI | Business Operations |
| Bausch & Lomb Incorporated | | 079587625 | manufacture(0363-0869) |

Revised: 12/2025

WALGREEN COMPANY