

**OLOPATADINE HYDROCHLORIDE- olopatadine hydrochloride solution**  
**YYBA Corp**

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**Olopatadine 2%**

Active Ingredients - Purpose - Olopatadine 0.2% (equivalent to olopatadine hydrochloride 0.222%) Antihistamine

| Active Ingredients   | Purpose       |
|--|---------------|
| Olopatadine 0.2%<br>(equivalent to olopatadine hydrochloride 0.222%) | 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**

- 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) 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°C to 25°C (36°F to 77°F)

### **Inactive ingredients**

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

### **Questions?**

In the U.S., call toll free weekdays 9 AM to 6 PM EST at 1 (866) 933-6337.

Distributed by:

Wellspring Meds USA

Airmont, NY 10952

Olopatadine hydrochloride  
ophthalmic solution 0.2%

Antihistamine  
2.5 mL  
STERILE

EYE ALLERGY ITCH RELIEF

|   |  |   |
|---|--|---|
| <p>Only for use in the eye.<br/>Store between 2°C to 25°C (36°F to 77°F).<br/><b>TAMPER EVIDENT:</b> For your protection, this bottle has a tamper-evident ring attached to the bottle cap. Do not use if seal is broken or missing.</p> <p>Distributed by<br/><b>Wellspring</b><br/>Airmont, NY 10952 U.S.A.<br/>866-933-6337<br/>wellspringmeds.com</p> | <p></p> <p><b>OLOPATADINE<br/>HCl OPHTHALMIC<br/>SOLUTION, USP</b></p> <p><b>Eye Allergy Itch Relief</b></p> <p><b>ANTIHISTAMINE</b> <b>STERILE</b></p> <p>2.5 mL (0.085 FL OZ)</p> | <p><b>Product of Spain</b> Code No.: 1335</p> <p>LOT</p> <p>EXP</p> <p></p> <p>3 73581 00087 6</p> |
|---|--|---|

Original Prescription Strength

ONCE DAILY RELIEF

Olopatadine hydrochloride ophthalmic solution 0.2%  
Antihistamine

Eye Allergy Itch Relief

STERILE  
2.5 mL

BOTTLE 2x2.5 ML  
SAME SIZE ARTWORK  
CRT SIZE: 50 mm x 31 mm x 96 mm



| OLOPATADINE HYDROCHLORIDE          |                |                    |               |
|------------------------------------|----------------|--------------------|---------------|
| olopatadine hydrochloride solution |                |                    |               |
| Product Information                |                |                    |               |
| Product Type                       | HUMAN OTC DRUG | Item Code (Source) | NDC:73581-709 |
| Route of Administration            | OPHTHALMIC     |                    |               |

**Active Ingredient/Active Moiety**

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

**Inactive Ingredients**

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

**Packaging**

| # | Item Code        | Package Description                                   | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:73581-709-02 | 2 in 1 CARTON   | 01/01/2026           |                    |
| 1 |                  | 2.5 mL in 1 BOTTLE; Type 0: Not a Combination Product |                      |                    |

**Marketing Information**

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| ANDA               | ANDA219557                               | 01/01/2026           |                    |

**Labeler** - YYBA Corp (006339772)**Registrant** - YYBA Corp (006339772)