**OLOPATADINE HCL- olopatadine hcl solution/ drops**
**Sola Pharmaceuticals**

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**Olopatadine Hcl**

***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 our 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 opthalmic 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)

***Inactive ingredients***

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

***Questions?***

Call 1-866-747-7365

**Principle Display Panel**

Manufactured for:

SOLA Pharmaceuticals LLC,

Baton Rouge, LA 70810

Made in India

Code No: DD/DRUGS/DD/292

Olopatadine Hcl Opthalmic Solution 0.1% Bottle Label:

Olopatadine Hcl Opthalmic Solution 0.1% Carton Label:


# OLOPATADINE HCL

olopatadine hcl solution/ drops

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:70512-520 |
| **Route of Administration** | OPHTHALMIC | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **OLOPATADINE HYDROCHLORIDE** (UNII: 2XG66W44KF) (OLOPATADINE - | OLOPATADINE | 1 mg |

| | | |
|---|---|---|
| UNII:D27V6190PM) | HYDROCHLORIDE | in 1 mL |

## Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **BENZALKONIUM CHLORIDE** (UNII: F5UM2KM3W7) | |
| **WATER** (UNII: 059QF0KO0R) | |
| **SODIUM HYDROXIDE** (UNII: 55X04QC32I) | |
| **SODIUM CHLORIDE** (UNII: 451W47IQ8X) | |
| **SODIUM PHOSPHATE, DIBASIC** (UNII: GR686LBA74) | |
| **HYDROCHLORIC ACID** (UNII: QTT17582CB) | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:70512-520-05 | 1 in 1 CARTON | 07/05/2022 | |
| **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 | ANDA203152 | 07/05/2022 | |

## Labeler - Sola Pharmaceuticals (080121345)

Revised: 5/2024 Sola Pharmaceuticals