OLOPATADINE HCL- olopatadine hcl solution/ drops
Sola Pharmaceuticals

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: 11/2023 Sola Pharmaceuticals