

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 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)

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 Ophthalmic Solution 0.1% Bottle Label:

TWICE DAILY
Only for use in the eye.
Store between 4°-25°C
(39°-77°F).

TAMPER EVIDENT:
For your protection, this
bottle has a seal around
the neck. Do not use if
seal is damaged or
missing at time of
purchase.

NDC 70512-0520-05
Olopatadine HCl
Ophthalmic Solution,
USP 0.1%

LOT:
EXP.:

Antihistamine and Redness Reliever
Eye Allergy Itch & Redness Relief

5 mL (0.17 FL OZ) STERILE

Manufactured for:
SOLA PHARMACEUTICALS LLC,
Baton Rouge, LA 70810
Made in India
Code No.: DD/DRUGS/DD/292

3016698

Olopatadine Hcl Ophthalmic Solution 0.1% Carton Label:

3016699

1

Drug Facts

Active ingredient	Purpose
Olopatadine (0.1%) (equivalent to olopatadine hydrochloride 0.111%)	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

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When using this product

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Drug Facts (continued)

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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.

Manufactured for:
SOLA PHARMACEUTICALS LLC,
Baton Rouge, LA 70810

Made in India
Code No.: DD/DRUGS/DD/292

NDC 70512-0520-05
Original Prescription Strength

Olopatadine HCl Ophthalmic Solution, USP 0.1%

Antihistamine and Redness Reliever

Eye Allergy Itch & Redness Relief

TWICE DAILY
Works in minutes
Relief from Allergens:
• Pet Dander • Pollen
• Grass • Ragweed



5 mL (0.17 FL OZ) • STERILE

Olopatadine HCl Ophthalmic Solution, USP 0.1%
Eye Allergy Itch & Redness Relief



Works in Minutes For Ages 2 and Older 30 DAY SUPPLY

TAMPER EVIDENT:
For your protection, this bottle has a seal around the neck.
— Fill Line —

Do not use if seal is damaged or missing at time of purchase.

ACTUAL SIZE



7 05120 52005 5

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