

OLOPATADINE HYDROCHLORIDE OPHTHALMIC SOLUTION- olopatadine hydrochloride ophthalmic solution/ drops
Chain Drug Marketing Association INC

ACTIVE INGREDIENT

Olopatadine (0.1%) (equivalent to olopatadine hydrochloride 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 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° to 25°C (39° to 77°F)

INACTIVE INGREDIENTS

Benzalkonium chloride 0.01%, Dibasic sodium phosphate, Hydrochloric acid and /or Sodium hydroxide (to adjust pH), Sodium chloride and Water for Injection.

QUESTIONS?

Call 1-888-375-3784

PRINCIPAL DISPLAY PANEL

NDC 63868-822-05

**Olopatadine
Hydrochloride
Ophthalmic
Solution, USP
Bottle Label:**

	<p>Each mL contains: Olopatadine (0.1%) (equivalent to olopatadine hydrochloride 0.111%) Store between 4°-25°C (39°-77°F) Packaged with Tamper-Evident bottle cap. Do Not Use if breakable ring is separated or missing. Code: AP/DRUGS/103/97 LAB-020866-00 REV: 06/20</p>	<p>Olopatadine Hydrochloride Ophthalmic Solution, USP 0.1% Antihistamine and Redness Reliever TWICE DAILY RELIEF Only for use in the eye Eye Allergy Itch & Redness Relief STERILE 5 mL</p>	 <p>(01)00343598765075</p> <p>LOT: _____ EXP: _____</p>	<p>DISTRIBUTED BY: Dr. Reddy's Laboratories, Inc. Princeton, NJ 08540 Made in India</p>
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Carton Label:



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

NDC 63868-822-05



Compare to the active ingredient in Pataday®
Twice Daily Relief*

NOW AVAILABLE without a prescription
Olopatadine Hydrochloride Ophthalmic Solution USP, 0.1%
Antihistamine and Redness Reliever

Eye Allergy Itch & Redness Relief

TWICE DAILY RELIEF

Works in Minutes

- Relief from Allergens:
- Pet Dander • Pollen
 - Grass • Ragweed



TWICE DAILY

Packaged with Tamper-Evident bottle cap. Do Not Use if breakable ring is separated or missing.

Olopatadine Hydrochloride Ophthalmic Solution, USP

0.1%

TWICE DAILY RELIEF

FILL LINE

30 DAY SUPPLY

Eye Allergy Itch & Redness Relief
Works in Minutes
For Ages 2 and Older

Actual Size Bottle



Distributed by C.D.M.A., Inc®
43157 W 9 Mile Rd
Novi, MI 48375
www.cdma@tychochoice.com
Questions: 800-955-2362

Made in India
Code: AP/DRUGS/103/97 REV: 10/20

Drug Facts

Active ingredient Purpose
Olopatadine (0.1%).....Antihistamine and Redness Reliever (equivalent to olopatadine hydrochloride 0.111%)

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 (1-800-222-1222).

Drug Facts (continued)

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
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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 and /or Sodium hydroxide (to adjust pH), Sodium chloride and Water for Injection.

Questions?
call 1-888-375-3784

*This product is not manufactured or distributed by Alcon Laboratories Inc., distributor of Pataday® Twice Daily Relief. Pataday® is a registered trademark of Novartis AG.



5 mL STERILE



LOT
EXP

OLOPATADINE HYDROCHLORIDE OPHTHALMIC SOLUTION

olopatadine hydrochloride ophthalmic solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63868-822(NDC:43598-765)
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OLOPATADINE HYDROCHLORIDE (UNII: 2XG66W44KF) (OLOPATADINE - UNII:D27V6190PM)	OLOPATADINE	1 mg in 1 mL

Inactive Ingredients

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

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-822-05	1 in 1 CARTON	01/01/2021	
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	ANDA209619	01/01/2021	

Labeler - Chain Drug Marketing Association INC (011920774)

Establishment			
Name	Address	ID/FEI	Business Operations
Reed Lane Inc		001819879	repack(63868-822)

Revised: 12/2025

Chain Drug Marketing Association INC