

OLOPATADINE HYDROCHLORIDE OPHTHALMIC SOLUTION- olopatadine hydrochloride ophthalmic solution/ drops
Dr. Reddy's Laboratories 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



Dr.Reddy's 

Olopatadine Hydrochloride Ophthalmic Solution, USP
0.1%

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

Drug Facts

Active ingredient Purpose
Olopatadine (0.1%).....Antihistamine and Redness Reliever (equivalent to olopatadine hydrochloride 0.11%)

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
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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
- 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 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.

Dr.Reddy's  NDC 43598-765-07
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

TWICE DAILY RELIEF
Eye Allergy Itch & Redness Relief

Works in Minutes
Relief from Allergens:

- Pet Dander
- Pollen
- Grass
- Ragweed

TWICE DAILY





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



5 mL **STERILE**



LOT

EXP

DIST. BY: Dr. Peddy's Laboratories, Inc.
Princeton, NJ 08540
Made in India
Code: AP/DRUGS/103/97 REV: 06/20

OLOPATADINE HYDROCHLORIDE OPHTHALMIC SOLUTION			
olopatadine hydrochloride ophthalmic solution/ drops			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	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
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:43598-765-07	1 in 1 CARTON	09/01/2020	
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	09/01/2020	

Labeler - Dr. Reddy's Laboratories Inc. (802315887)

Revised: 12/2025

Dr. Reddy's Laboratories Inc.