

**EYE ALLERGY ITCH RELIEF ONCE DAILY RELIEF- olopatadine hydrochloride
ophthalmic solution
Dr. Reddy's Laboratories Inc.**

ACTIVE INGREDIENT

Olopatadine (0.2%) (equivalent to olopatadine hydrochloride 0.222%)

PURPOSE

Antihistamine

USE

temporarily relieves itchy 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) once daily, no more than once 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 2° to 25°C (36° to 77°F)

INACTIVE INGREDIENTS

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

QUESTIONS?

Call 1-888-375-3784

PRINCIPAL DISPLAY PANEL

Olopatadine Hydrochloride
Ophthalmic Solution, USP
0.2%
Single pack:



Drug Facts

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Purpose Antihistamine (equivalent to olopatadine hydrochloride 0.222%)

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Drug Facts (continued)

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

HealthCare  Aisle

Eye Allergy Itch Relief

Olopatadine Hydrochloride Ophthalmic Solution USP, **0.2%**

HealthCare  Aisle

NDC 43598-827-02

NOW AVAILABLE Without A Prescription

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

ONCE DAILY RELIEF

Eye Allergy Itch Relief

Olopatadine Hydrochloride Ophthalmic Solution, USP

0.2%

Antihistamine

Works in Minutes

Relief from Allergens:

- Pet Dander
- Pollen
- Grass
- Ragweed



STERILE

2.5 mL

ONCE DAILY



3 43598 82702 5

LOT

EXP

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

Eye Allergy Itch Relief

Olopatadine Hydrochloride Ophthalmic Solution, USP

0.2%

ONCE DAILY RELIEF

FILL LINE

Works in Minutes
For Ages 2 and Older
30 DAY SUPPLY

Actual Size Bottle

DISTRIBUTED BY:
Dr. Reddy's Laboratories, Inc.
Princeton, NJ 08540
Made in India

Code: AP/DRUGS/103/97 REV: 01/21



EYE ALLERGY ITCH RELIEF ONCE DAILY RELIEF

olopatadine hydrochloride ophthalmic solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:43598-827(NDC:43598-764)
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
POVIDONE K30 (UNII: U725QWY32X)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:43598-827-02	1 in 1 CARTON	10/01/2020	
1		2.5 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:43598-827-25	2 in 1 CARTON	10/01/2020	
2		2.5 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	ANDA209752	10/01/2020	

Revised: 4/2021

Dr. Reddy's Laboratories Inc.