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

ACTIVE INGREDIENT

Olopatadine (0.7%) (equivalent to olopatadine hydrochloride 0.776%)

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(1-800-222-1222).

DIRECTIONS

- **adults and children 2 years of age and older:**
- put **1 drop** in the affected eye(s) **once daily**
- **do not use more than 1 drop in each eye 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 0.015% (preservative), boric acid, hydrochloric acid/sodium hydroxide (to adjust pH), hydroxypropyl gamma-cyclodextrin, hypromellose, mannitol, polyethylene glycol 400, povidone and purified water.

QUESTIONS?

call toll-free weekdays 8 AM to 8 PM EST at **1-888-375-3784**

Distributed by:

Dr. Reddy's Laboratories, Inc.

Princeton, NJ 08540

Made in India

REV: 01/26

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

PRINCIPAL DISPLAY PANEL

Carton Label:

NDC 75907-285-02

Extra Strength

Olopatadine Hydrochloride
Ophthalmic Solution USP, 0.7%
Antihistamine

ONCE DAILY RELIEF

Eye Allergy Itch Relief

Full 24HR

Works in Minutes

Sterile

2.5 mL (0.085 fl oz)

Compare to the active ingredient in Pataday Once Daily Relief - Extra Strength *

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

For Ages 2 and Older

30 DAY SUPPLY

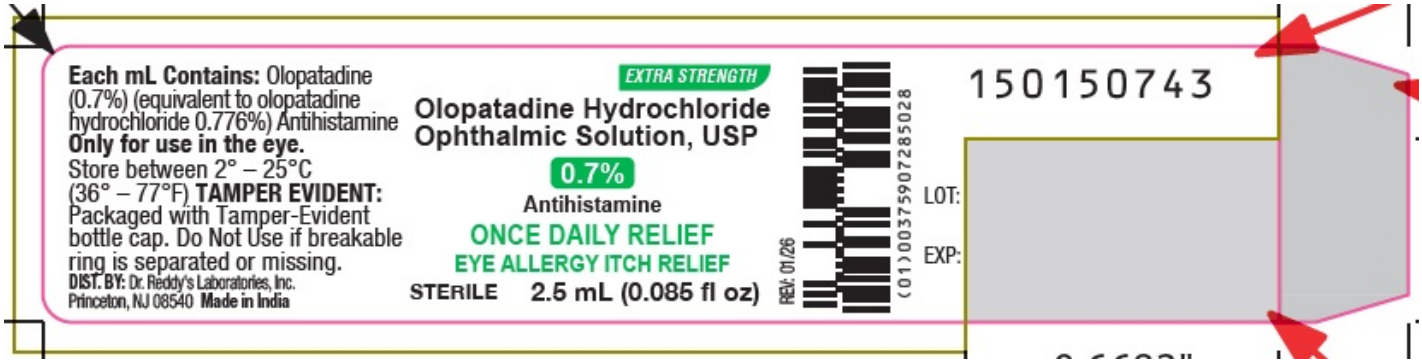
Bottle Label:

Extra Strength

Eye Allergy Itch Relief

Olopatadine Hydrochloride
Ophthalmic Solution USP, 0.7%
Antihistamine
ONCE DAILY RELIEF

Sterile
2.5 mL (0.085 fl oz)



OLOPATADINE HYDROCHLORIDE OPHTHALMIC SOLUTION

olopatadine hydrochloride solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75907-285
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OLOPATADINE HYDROCHLORIDE (UNII: 2XG66W44KF) (OLOPATADINE - UNII:D27V6190PM)	OLOPATADINE	7 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BORIC ACID (UNII: R57ZHV85D4)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
HYDROXYPROPYL .GAMMA.-CYCLODEXTRIN (UNII: P6BYU725IU)	
HYPROMELLOSE 2910 (4000 MPA.S) (UNII: RN3152OP35)	
MANNITOL (UNII: 3OWL53L36A)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
POVIDONE K30 (UNII: U725QWY32X)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75907-285-02	1 in 1 CARTON	04/15/2026	

1		2.5 mL in 1 BOTTLE, DROPPER; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		
2	NDC:75907-285-25	2 in 1 CARTON	04/15/2026	
2		2.5 mL in 1 BOTTLE, DROPPER; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA213514	04/15/2026	

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

Revised: 4/2026 Dr. Reddy's Laboratories Inc.