

OLOPATADINE HYDROCHLORIDE OPHTHALMIC SOLUTION ONCE DAILY-
olopatadine hydrochloride ophthalmic solution
DOLGENCORP, LLC

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%

591980091



SEAL AREA
INK FREE AREA

Drug Facts

Active ingredient Purpose
Olopatadine (0.2%).....Antihistamine
(equivalent to olopatadine hydrochloride 0.222%)

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 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) 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°-25°C (36°-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

*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.

DG health

Eye Allergy Itch Relief

Olopatadine Hydrochloride Ophthalmic Solution USP, 0.2% Antihistamine

DG health

NDC 55910-137-25
Compare to the active ingredient of Pataday® Once Daily Relief*

Original Prescription Strength

Eye Allergy Itch Relief

Olopatadine Hydrochloride Ophthalmic Solution USP, 0.2% Antihistamine



Once Daily Relief

Works in Minutes Relief from Allergens:
• Pet Dander • Pollen
• Grass • Ragweed

Sterile

2.5 mL (0.085 FL OZ) Once Daily

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



DISTRIBUTED BY OLD EAST MAIN CO.
100 MISSION RIDGE, GOODLETTSVILLE, TN 37072
MADE IN INDIA
Code: AP/DRUGS/103/97 REV:08/21

SEAL AREA
INK FREE AREA



150086163

A1550



LOT
EXP

no coating area

Each mL contains: Olopatadine (0.2%) (equivalent to olopatadine hydrochloride 0.222%)
Store between 2°-25°C (36°-77°F)
Packaged with Tamper-Evident bottle cap. Do Not Use if breakable ring is separated or missing.
DIST. BY: Dr. Reddy's Laboratories, Inc. Princeton, NJ 08540 Made in India

Olopatadine Hydrochloride Ophthalmic Solution, USP

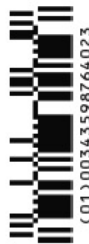
0.2%

Antihistamine
ONCE DAILY RELIEF
Only for use in the eye
Eye Allergy Itch Relief

STERILE

2.5 mL

REV:01/21



(01)00343598764023

LOT:
EXP:

150084228

OLOPATADINE HYDROCHLORIDE OPHTHALMIC SOLUTION ONCE DAILY

olopatadine hydrochloride ophthalmic solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55910-137(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:55910-137-25	1 in 1 CARTON	12/01/2021	
1		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	12/01/2021	

Labeler - DOLGENCORP, LLC (068331990)

Revised: 8/2023

DOLGENCORP, LLC