

**CVS EYE ALLERGY ITCH RELIEF ONCE DAILY- olopatadine hydrochloride
ophthalmic solution
CVS Health Corp**

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%**

6 2 8 1 8 0 0 9 1
CO FPO 28
 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.



ONCE DAILY
Olopatadine HCl
 OPHTHALMIC
 SOLUTION USP, 0.2%

**NOW AVAILABLE
 WITHOUT A PRESCRIPTION**



Compare to
 Pataday® Once
 Daily Relief*
 NDC 69842-269-25

ONCE DAILY
Olopatadine HCl
 OPHTHALMIC
 SOLUTION USP, 0.2%
 Antihistamine
EYE ALLERGY ITCH RELIEF

Works in Minutes
 Relief from Allergens:
 Pet dander; Pollen;
 Grass; Ragweed



ONCE DAILY
 RELIEF

STERILE

Actual Bottle
 Size on Side Panel

2.5 mL (0.085 FL OZ)

Packaged with tamper-evident
 bottle cap. Do not use if breakable
 ring is separated or missing.

Distributed by: CVS Pharmacy, Inc.
 One CVS Drive, Woonsocket, RI 02895
 © 2020 CVS/pharmacy
 CVS.com® 1-800-SHOP CVS
 Made in India V-37111



30 DAY SUPPLY

For ages 2 and older

— FILL LINE —

Actual Size Bottle

Code: AP/DRUGS/103/97 REV: 07/20

SEAL AREA
 INK FREE AREA
CO FPO 28
 1 5 0 0 8 1 3 2 9

#403924
 FPO 80%
 UPC# 050428629451
 X XXXXXX XXXXXX X

LOT
 EXP
 no coating
 area

CVS EYE ALLERGY ITCH RELIEF ONCE DAILY

olopatadine hydrochloride ophthalmic solution

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|------------------------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:69842-269(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:69842-269-25 | 1 in 1 CARTON | 09/20/2020 | |
| 1 | | 2.5 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | | |
| 2 | NDC:69842-269-50 | 2 in 1 CARTON | 09/20/2020 | |
| 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 | 09/20/2020 | |

Labeler - CVS Health Corp (062312574)

Revised: 4/2021

CVS Health Corp