

ONCE DAILY RELIEF- olopatadine hydrochloride ophthalmic solution solution H E B

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%



H-E-B
100% GUARANTEE promise
 If you aren't completely pleased with this product, we'll be happy to replace it or refund your money. You have our word on it.

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

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

MADE WITH PRIDE AND CARE FOR H-E-B®, SAN ANTONIO, TEXAS 78204 • MADE IN INDIA
 *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.

Compare to Pataday® Once Daily Relief active ingredient*

NOW AVAILABLE NDC 37808-045-25 without a prescription

H-E-B

Olopatadine Hydrochloride Ophthalmic Solution USP, 0.2%

Antihistamine
Once Daily Relief

Eye Allergy Itch Relief

Works In Minutes
 Relief From Allergens:
 • Pet Dander
 • Pollen
 • Grass
 • Ragweed

Once Daily **STERILE**

2.5 mL (0.085 FL OZ)

Olopatadine Hydrochloride Ophthalmic Solution, USP

0.2%

ONCE DAILY RELIEF

Eye Allergy Itch Relief

— FILL LINE —

Works in Minutes For Ages 2 and Older 30 DAY SUPPLY

Actual Size Bottle



Code: AP/DRUGS/103/97 REV: 12/21



16882-2112

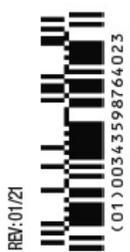
0 41220 53050 2

LOT: _____

EXP: _____

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**



150084228

LOT: _____

EXP: _____

ONCE DAILY RELIEF

olopatadine hydrochloride ophthalmic solution solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:37808-045(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:37808-045-25	1 in 1 CARTON	03/07/2022	
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	03/07/2022		

Labeler - H E B (007924756)

Revised: 8/2023

H E B