EYE ALLERGY ITCH RELIEF ONCE DAILY RELIEF- olopatadine hydrochloride ophthalmic solution Strategic Sourcing Services 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 833-358-6431 Monday to Friday 9:00am to 7:00pm EST

PRINCIPAL DISPLAY PANEL

Olopatadine Hydrochloride Ophthalmic Solution, USP 0.2%



EYE ALLERGY ITCH RELIEF ONCE DAILY RELIEF

olopatadine hydrochloride ophthalmic solution

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70677-1155
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)			

l	Packaging				
	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
	1 NDC:70677- 1155-1	1 in 1 CARTON	08/04/2023		
l	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	03/15/2021		

Labeler - Strategic Sourcing Services LLC (116956644)

Revised: 7/2023 Strategic Sourcing Services LLC