OLOPATADINE HYDROCHLORIDE OPHTHALMIC SOLUTION- olopatadine hydrochloride ophthalmic solution/ drops Walgreens Company

ACTIVE INGREDIENT

Olopatadine (0.1%) (equivalent to olopatadine hydrochloride 0.111%)

PURPOSE

Antihistamine and redness reliever

USES

temporarily relieves itchy and red 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) twice daily, every 6 to 8 hours, no more than twice 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 4° to 25°C (39° to 77°F)

INACTIVE INGREDIENTS

Benzalkonium chloride 0.01%, Dibasic sodium phosphate, Hydrochloric acid and /or Sodium hydroxide (to adjust pH), Sodium chloride and Water for Injection.

QUESTIONS?

Call 1-888-375-3784

PRINCIPAL DISPLAY PANEL

NDC 0363-7130-05





OLOPATADINE HYDROCHLORIDE OPHTHALMIC SOLUTION

olopatadine hydrochloride ophthalmic solution/ drops

aradiant Nar

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:0363-7130(NDC:43598-765) Route of Administration OPHTHALMIC Active Ingredient/Active Moiety

Basis of

іпутешені маше	Strength	Strength
OLOPATADINE HYDROCHLORIDE (UNII: 2XG66W44KF) (OLOPATADINE - UNII: D27V6190PM)	OLOPATADINE	1 mg in 1 mL

Inactive Ingredients			
Ingredient Name	Strength		
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)			
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)			
HYDROCHLORIC ACID (UNII: QTT17582CB)			
SODIUM CHLORIDE (UNII: 451W47IQ8X)			
SODIUM HYDROXIDE (UNII: 55X04QC32I)			
WATER (UNII: 059QF0KO0R)			

F	Packaging						
#	t Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:0363- 7130-05	1 in 1 CARTON	09/15/2020				
1		5 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product					

Marketing I	Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
ANDA	ANDA209619	09/15/2020			

Labeler - Walgreens Company (008965063)

Revised: 4/2021 Walgreens Company