

OLOPATADINE- olopatadine hydrochloride solution/ drops

Akorn

Drug Facts

Active ingredient

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

Purpose

Antihistamine and redness reliever

Use

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 the 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.

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 at 4° to 25°C (39° to 77°F).

Inactive ingredients

benzalkonium chloride 0.01%, dibasic sodium phosphate, hydrochloric acid and/or sodium hydroxide (adjust pH), sodium chloride, and water for injection

Questions?

Call toll-Free 1-800-932-5676 weekdays, 7:00 AM - 5:30 PM CST

Principal Display Panel Text for Container Label:

NDC 17478-308-05

Olopatadine HCl Ophthalmic

Solution, USP 0.1%

Antihistamine and Redness Reliever

Eye Allergy Itch & Redness Relief

STERILE 5 mL (0.17 FL OZ)

The image shows a simulated container label for Olopatadine HCl Ophthalmic Solution, USP 0.1%. The label is enclosed in a red rounded rectangular border. On the left, there is a box labeled 'Manufacturing Code'. The main text includes: 'Only for use in the eye. TWICE DAILY. Each mL contains: Active: Olopatadine (0.1%) (equivalent to olopatadine hydrochloride 0.111%). Storage: Store at 4° to 25°C (39° to 77° F). OLPAL Rev. 03/21'. The product name 'Olopatadine HCl Ophthalmic Solution, USP 0.1%' is highlighted in a teal box. Below it, 'Antihistamine and Redness Reliever Eye Allergy Itch & Redness Relief' is written. At the bottom, it says 'STERILE 5 mL (0.17 FL OZ)'. On the right, there is a 'TAMPER EVIDENT' warning, 'Keep out of reach of children.', and 'Manufactured by: Akorn, Inc. Lake Forest, IL 60045'. A vertical box on the right contains 'RSS Barcode'. At the bottom right, 'Lot: Exp:' is indicated.

Principal Display Panel Text for Carton Label:

Original NDC 17478-308-05

Prescription

Strength Akorn logo

Olopatadine

HCl Ophthalmic

Solution, USP

0.1%

Antihistamine and Redness Reliever

Eye Allergy Itch & Redness Relief

Works in Minutes

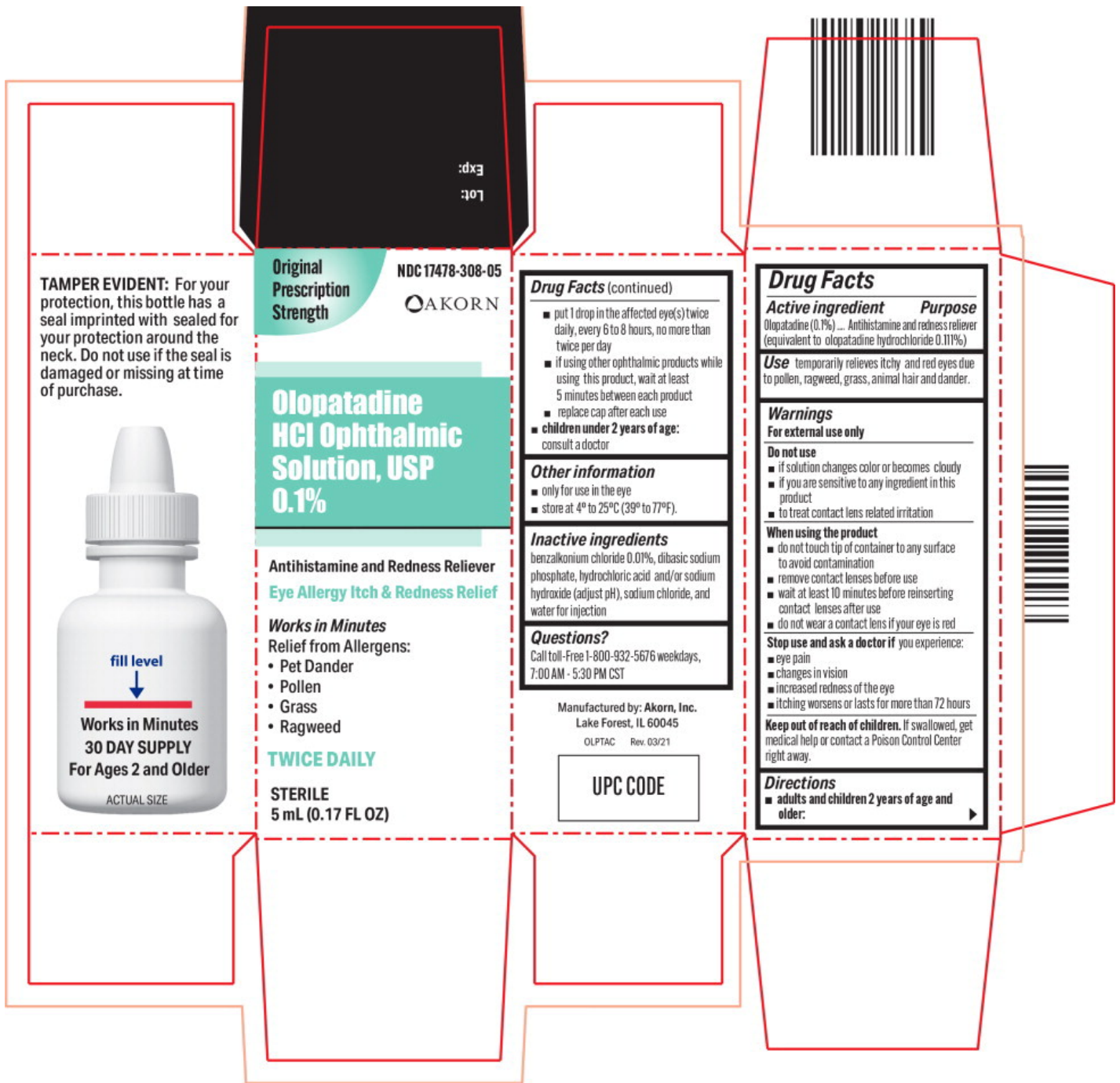
Relief from Allergens:

- Pet Dander
- Pollen
- Grass
- Ragweed

TWICE DAILY

STERILE

5 mL (0.17 FL OZ)



OLOPATADINE

olopatadine hydrochloride solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:17478-308
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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Olopatadine Hydrochloride (UNII: 2XG66W44KF) (Olopatadine - UNII:D27V6190PM)	Olopatadine	1 mg in 1 mL
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Inactive Ingredients

Ingredient Name	Strength
benzalkonium chloride (UNII: F5UM2KM3W7)	
sodium phosphate, dibasic, unspecified form (UNII: GR686LBA74)	
hydrochloric acid (UNII: QTT17582CB)	
sodium hydroxide (UNII: 55X04QC32I)	
sodium chloride (UNII: 451W47IQ8X)	
water (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:17478-308-05	1 in 1 CARTON	09/01/2021	
1		5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA204532	09/01/2021	

Labeler - Akorn (117696770)

Registrant - AKORN OPERATING COMPANY LLC (117693100)

Establishment

Name	Address	ID/FEI	Business Operations
Akorn		117696840	LABEL(17478-308) , MANUFACTURE(17478-308) , PACK(17478-308) , ANALYSIS(17478-308) , STERILIZE(17478-308)

Revised: 2/2022

Akorn