OLOPATADINE HYDROCHLORIDE- olopatadine hydrochloride solution
Bausch & Lomb Incorporated

Drug Facts

Active ingredient

Olopatadine (0.2%)

(equivalent to olopatadine hydrochloride 0.222%)

Purpose

Antihistamine

Uses

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 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 4 —25 °C (39 —77 °F)
- protect from light

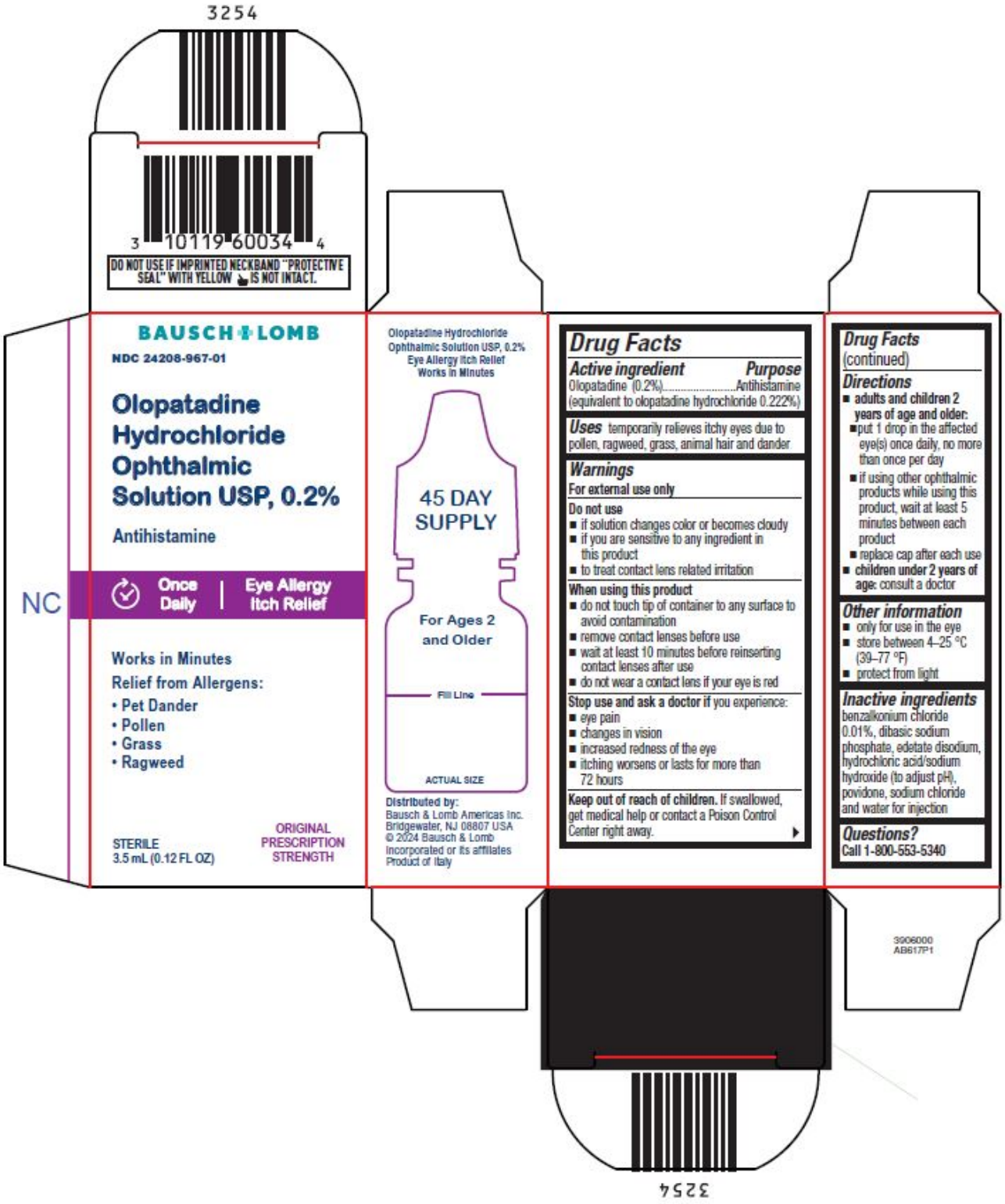
Inactive ingredients

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

Questions?

Call 1-800-553-5340

Principle Display Panel



BAUSCH + LOMB

NDC 24208-967-01

Olopatadine

Hydrochloride

Ophthalmic

Solution USP, 0.2%

Antihistamine

Once Daily | Eye Allergy
Itch Relief

Works in Minutes
Relief from Allergens:

- Pet Dander
- Pollen
- Grass
- Ragweed

STERILE

3.5 ml (0.12 FL OZ)

ORIGINAL

PRESCRIPTION

STRENGTH

3906000
AB617P1

OLOPATADINE HYDROCHLORIDE

olopatadine hydrochloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:24208-967
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**
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
SODIUM PHOSPHATE, DIBASIC, UNSPECIFIED FORM (UNII: GR686LBA74)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

EDETATE DISODIUM (UNII: 7FLD91C86K)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:24208-967-01	1 in 1 CARTON	12/02/2024	
1		3.5 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:24208-967-02	2 in 1 CARTON	12/02/2024	
2		3.5 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:24208-967-03	3 in 1 CARTON	12/02/2024	
3		3.5 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	ANDA206087	12/02/2024	

Labeler - Bausch & Lomb Incorporated (196603781)

Establishment

Name	Address	ID/FEI	Business Operations
Bausch & Lomb Incorporated		079587625	manufacture(24208-967)

Revised: 10/2024 Bausch & Lomb Incorporated