

**OLOPATADINE HYDROCHLORIDE- olopatadine hydrochloride
ophthalmic solution
Golden State Medical Supply, Inc.**

**OLOPATADINE HYDROCHLORIDE.
Drug Facts**

Active Ingredient	Purpose
Olopatadine (0.1%). (equivalent to olopatadine hydrochloride 0.111%)	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 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 between 4°C to 25°C (39°F 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 or comments?

Call weekdays 9 AM to 6 PM EST at **1 (888) 721-7115.**

Distributed by:

Glenmark Therapeutics Inc., USA

Elmwood Park, NJ 07407

Product of Spain

07/25

Marketed by:

GSMS, Inc.

Camarillo, CA 93012 USA

PRINCIPAL DISPLAY PANEL

NDC 51407-938-05

Olopatadine HCl Ophthalmic Solution, USP 0.1%

Antihistamine and Redness Reliever

Twice Daily Relief

Eye Allergy Itch & Redness Relief

STERILE

5 mL (0.17 FL OZ)



OLOPATADINE HCl OPTHALMIC SOLUTION, USP 0.1%

Antihistamine and Redness Reliever
Eye Allergy Itch & Redness Relief



STERILE
 5 mL (0.17 FL OZ)

Original Prescription Strength

NDC 51407-938-05

[†]Compare to the active ingredient in Pataday[®] Twice Daily Relief

OLOPATADINE HCl OPTHALMIC SOLUTION, USP 0.1%

Antihistamine and Redness Reliever

Eye Allergy Itch & Redness Relief

Relief from Allergens:

- Pet Dander
- Pollen
- Grass
- Ragweed



TWICE DAILY
 STERILE
 5 mL (0.17 FL OZ)

For Ages 2 And Older
 Works in Minutes
 30 Day Supply

OLOPATADINE HCl OPTHALMIC SOLUTION, USP 0.1%

Antihistamine and Redness Reliever

TAMPER EVIDENT:

For your protection, this bottle has a tamper-evident ring attached to the bottle cap.

Do not use if seal is broken or missing.

Fill Line

ACTUAL SIZE

Drug Facts (continued)

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°C to 25°C (39°F 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 weekdays 9 AM to 6 PM EST at 1 (888) 721-7115.

[†]This product is not manufactured or distributed by Alcon Laboratories, Inc., distributor of Pataday[®] Twice Daily Relief, and/or Novartis AG, owner of the registered trademark Pataday[®], respectively. [Neither Alcon nor Novartis make or license this product.]

Marketed by: GSMS, Incorporated
 Camarillo, CA 93012 USA
 Product of Spain Code No.: 1335 01/26

Drug Facts

Active ingredient Purpose
 Olopatadine (0.1%)..... Antihistamine and redness reliever
 (equivalent to olopatadine hydrochloride 0.111%)

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 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. ➔



LOT
 EXP

NDC 51407-938-05

Olopatadine HCl Ophthalmic Solution, USP 0.1%

Antihistamine and Redness Reliever

Twice Daily

STERILE

5 mL (0.17 FL OZ)

Only for use in the eye.

Store between 4°C to 25°C (39°F to 77°F).

TAMPER EVIDENT: For your protection, this bottle has a tamper-evident ring attached to the bottle cap. Do not use if seal is broken or missing.

Marketed by: **GSMS**, Incorporated
Camarillo, CA 93012 USA

NDC 51407-938-05

OLOPATADINE HCl OPTHALMIC SOLUTION, USP **0.1%**

Antihistamine and Redness Reliever
Eye Allergy Itch & Redness Relief

gsms

STERILE TWICE
5 mL (0.17 FL OZ) DAILY

Product of Spain
01/26

Code No.: 1335

LOT

EXP



OLOPATADINE HYDROCHLORIDE

olopatadine hydrochloride ophthalmic solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51407-938
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OLOPATADINE HYDROCHLORIDE (UNII: 2XG66W44KF) (OLOPATADINE - UNII:D27V6190PM)	OLOPATADINE	1 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51407-938-05	1 in 1 CARTON	03/03/2026	
1		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	ANDA200810	06/28/2017	

Labeler - Golden State Medical Supply, Inc. (603184490)

Establishment

Name	Address	ID/FEI	Business Operations
Sam Chun Dang Pharm Co, Ltd		687792325	manufacture(51407-938)

Revised: 4/2026

Golden State Medical Supply, Inc.