PATADAY TWICE A DAY RELIEF- olopatadine hydrochloride solution
Alcon Laboratories, Inc.

Drug Facts

Active Ingredients	***Purpose***
Olopatadine (0.1%). (equivalent to olopatadine hydrochloride 0.111%)	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 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°-25°C (39°-77°F)

Inactive ingredients

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

Questions?

In the U.S., call 1-800-757-9195 or email alcon.medinfo@alcon.com

Alcon Laboratories, Inc.
6201 South Freeway
Fort Worth, Texas 76134
Country of Origin: Japan

NDC: 0065-4274-01
9017919-1219

PRINCIPAL DISPLAY PANEL

Original Prescription Strength

Pataday®
TWICE DAILY RELIEF
Olopatadine hydrochloride
ophthalmic solution 0.1%
Antihistamine and Redness Reliever

Eye Allergy Itch & Redness Relief
TWICE DAILY

Works in Minutes
Relief from Allergens:
- Pet Dander
- Pollen
- Grass
- Ragweed

STERILE
5 mL (0.17 FL OZ)

Alcon

Pataday®

TWICE DAILY RELIEF
Eye Allergy Itch & Redness Relief
Works in Minutes

For Ages 2 and Older
30 DAY SUPPLY

TAMPER EVIDENT: For your protection, this bottle has a seal imprinted with Alcon around the neck.

_____Fill Line_____

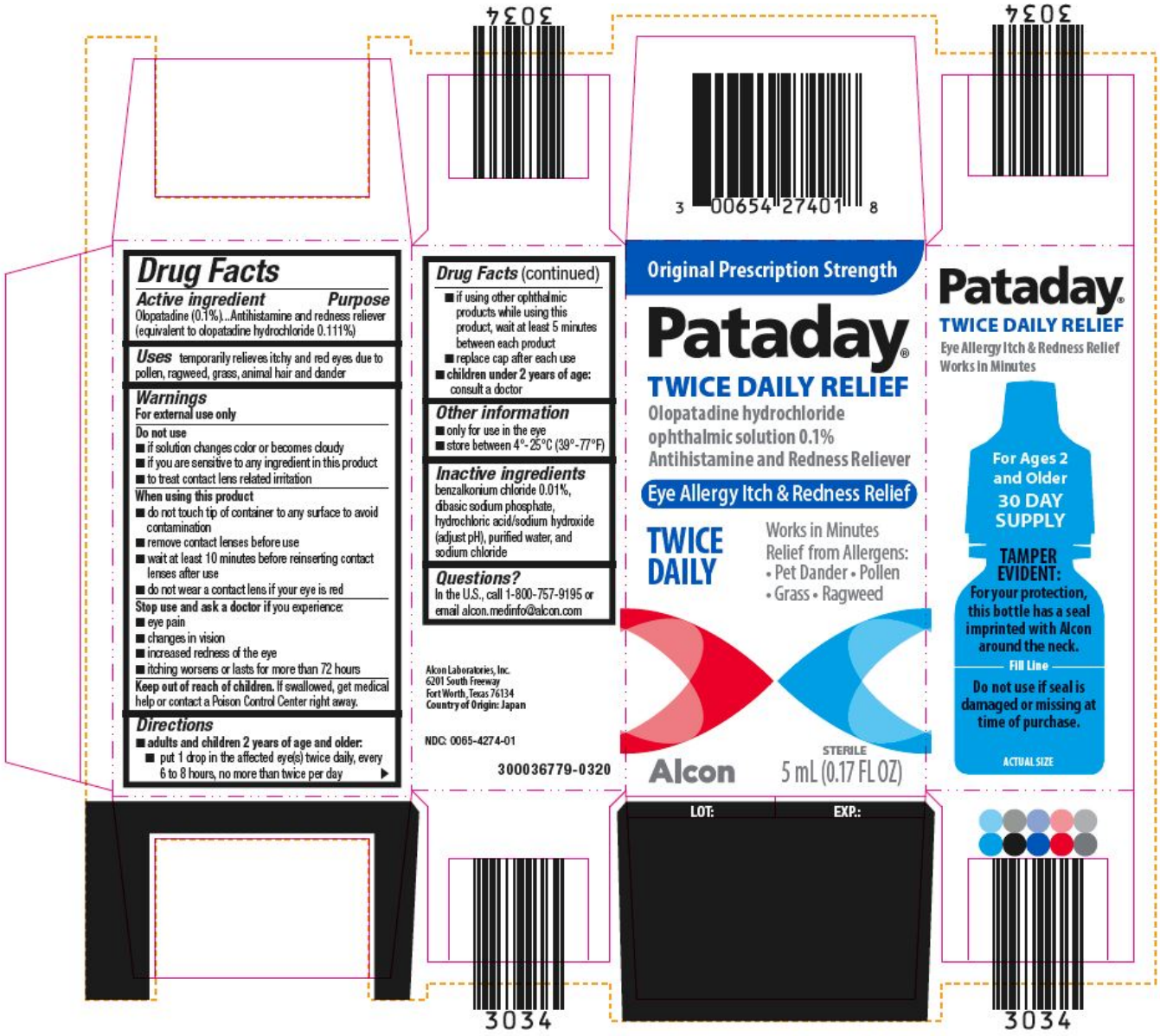
Do not use if seal is damaged or missing at time of purchase.

ACTUAL SIZE

Alcon Laboratories, Inc.
6201 South Freeway
Fort Worth, Texas 76134
Country of Origin: Japan

NDC: 0065-4274-01
300036779-0320

LOT: EXP.:



Pataday®
TWICE DAILY RELIEF
Olopatadine hydrochloride
ophthalmic solution 0.1%
Antihistamine and Redness Reliever
Eye Allergy Itch & Redness Relief

5 mL (0.17 FL OZ)
STERILE

Only for use in the eye.
Store between 4° – 25° C (39° – 77° F)

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6201 South Freeway
Fort Worth, TX 76134

LOT: EXP.:

H15723-1219

PATADAY TWICE A DAY RELIEF

olopatadine hydrochloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0065-4274
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**
Sodium Phosphate, Dibasic, Unspecified Form (UNII: GR686LBA74)	
Sodium Chloride (UNII: 451W47IQ8X)	
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:0065-4274-01	1 in 1 CARTON	02/25/2020	
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
NDA	NDA020688	02/25/2020	

Labeler - Alcon Laboratories, Inc. (008018525)

Establishment

Name	Address	ID/FEI	Business Operations
Alcon Research LLC		007672236	manufacture(0065-4274)

Revised: 12/2023 Alcon Laboratories, Inc.