PATADAY ONCE DAILY RELIEF- olopatadine hydrochloride solution
Alcon Laboratories, Inc.

Drug Facts

Active Ingredients	***Purpose***
Olopatadine 0.2% (equivalent to olopatadine hydrochloride 0.222%)	Antihistamine

Use 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 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) 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 2°-25°C (36°-77°F)

Inactive ingredients

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

Questions?

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

PRINCIPAL DISPLAY PANEL

Pataday®
ONCE DAILY RELIEF
Olopatadine hydrochloride
ophthalmic solution 0.2% Antihistamine

2.5 mL (0.085 FL OZ)
STERILE

EYE ALLERGY ITCH RELIEF
Only for use in the eye. Store between 2°– 25° C (36°– 77° F)

TAMPER EVIDENT: For your protection, this bottle has a seal imprinted with Alcon around the neck. Do not use if seal is damaged or missing at time of purchase.

Alcon Laboratories, Inc.
Fort Worth, TX 76134

LOT: EXP.:

H15725-219

Original Prescription Strength

Pataday

ONCE DAILY RELIEF

Olopatadine hydrochloride
ophthalmic solution 0.2%
Antihistamine

Eye Allergy Itch Relief

ONCE DAILY

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

STERILE

2.5 mL (0.085 FL OZ)

Alcon

Pataday

ONCE DAILY RELIEF

Eye Allergy Itch Relief
Works in Minutes

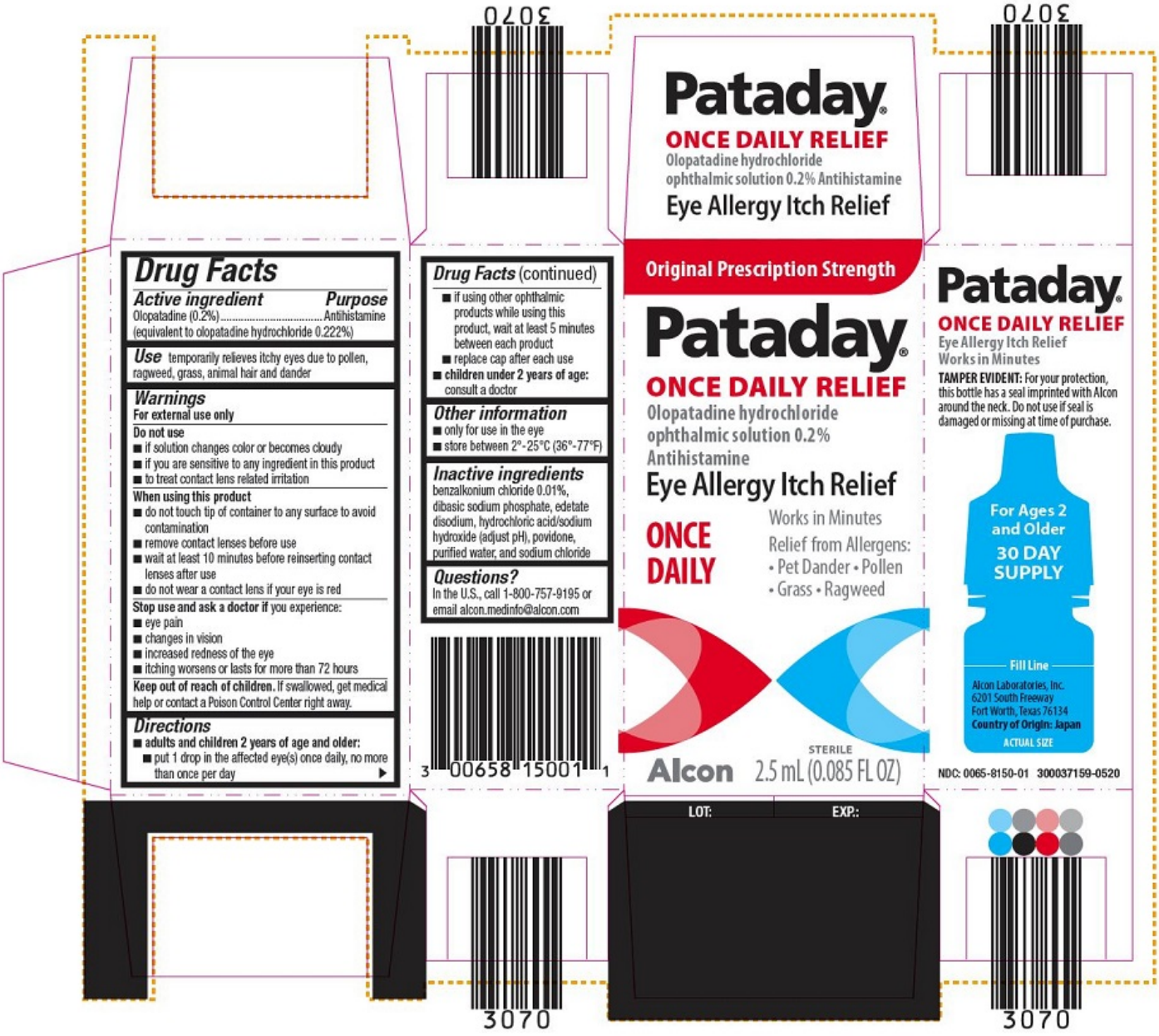
TAMPER EVIDENT: For your protection, this bottle has a seal imprinted with Alcon around the neck. Do not use if seal is damaged or missing at time of purchase.

For Ages 2
and Older
30 DAY
SUPPLY
_____Fill Line_____

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

ACTUAL SIZE

NDC: 0065-6150-01 300037159-0520



PATADAY ONCE DAILY RELIEF

olopatadine hydrochloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0065-8150
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:0065-8150-01	1 in 1 CARTON	02/28/2020	
1		2.5 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:0065-8150-03	2 in 1 CARTON	02/28/2020	
2		2.5 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:0065-8150-04	1 in 1 CARTON	02/28/2020	
3		0.5 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
4	NDC:0065-8150-07	3 in 1 CARTON	01/15/2021	
4		2.5 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	NDA021545	02/28/2020	

Labeler - Alcon Laboratories, Inc. (008018525)

Establishment

Name	Address	ID/FEI	Business Operations
Alcon Research LLC		007672236	manufacture(0065-8150)