ALAWAY- ketotifen fumarate solution/ drops
Bausch & Lomb Incorporated

Drug Facts

Active ingredient

Ketotifen 0.025%
(equivalent to ketotifen fumerate 0.035%)

Purpose

Antihistamine

Uses

For the temporary relief of itchy eyes due to ragweed, pollen, grass, animal hair and dander.

Warnings

For external use only

Do not use

- if you are sensitive to any ingredient in this product
- if solution changes color or becomes cloudy
- to treat contact lens related irritation

When using this product

- remove contact lenses before use
- wait at least 10 minutes before re-inserting contact lenses after use
- do not touch tip of container to any surface to avoid contamination
- replace cap after each use

Stop use and ask doctor if you experience any of the following:

- eye pain
- changes in vision
- redness of the eyes
- itching that worsens or lasts 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 3 years and older: put 1 drop in the affected eye(s) twice daily, every 8-12 hours, no more than twice per day.

Children under 3 years of age: consult a doctor

Other Information

Store at 4-25°C (39-77°F)

Inactive ingredients

benzalkonium chloride 0.01%, glycerin, hydrochloric acid and/or sodium hydroxide, water for injection

Questions or comments?

Toll Free Product Information

Call: 1-800-553-5340

Distributed by: Bausch + Lomb, a division of Bausch Health US, LLC
Bridgewater, NJ 08807
Product of Italy

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Package/Label Principal Display Panel

NDC 24208-601-10

BAUSCH + LOMB

Alaway®

ketotifen fumarate ophthalmic solution 0.035%

ANTIHISTAMINE EYE DROPS

UP TO

12

HOURS

EYE ITCH RELIEF

WORKS IN MINUTES!

- *Original Prescription Strength*
- *For ages 3 years and older*

60 DAY SUPPLY

STERILE 0.34 FL OZ (10 mL)

ALAWAY

ketotifen fumarate solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:24208-601
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
KETOTIFEN FUMARATE (UNII: HBD503WORO) (KETOTIFEN - UNII:X49220T18G)	KETOTIFEN	0.25 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:24208-601-10	1 in 1 CARTON	12/01/2006	
1		10 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
2	NDC:24208-601-95	1 in 1 CARTON	12/01/2006	09/30/2015

2		1 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
3	NDC:24208-601-05	1 in 1 CARTON	12/01/2006	
3		5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
4	NDC:24208-601-90	2 in 1 CARTON	12/01/2006	
4		10 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
NDA	NDA021996	12/01/2006	

Labeler - Bausch & Lomb Incorporated (196603781)

Establishment

Name	Address	ID/FEI	Business Operations
Bausch & Lomb Incorporated		079587625	MANUFACTURE(24208-601) , PACK(24208-601) , LABEL(24208-601)

Revised: 1/2020 Bausch & Lomb Incorporated