

LUMIFY REDNESS RELIEVER EYE DROPS- brimonidine tartrate solution/ drops
Bausch & Lomb Incorporated

Drug Facts

Active ingredient

Brimonidine tartrate (0.025%)

Purpose

Redness reliever

Use

- relieves redness of the eye due to minor eye irritations

Warnings

For external use only

Do not use

- if solution changes color or becomes cloudy

Stop use and ask a doctor if

- you experience eye pain, changes in vision, continued redness or irritation of the eye
- condition worsens or persists for more than 3 days

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- adults and children 5 years of age and over:
- instill 1 drop in the affected eye(s) every 6-8 hours
- do not use more than 4 times daily
- remove contact lenses before use
- wait at least 10 minutes before re-inserting contact lenses after use
- if using other ophthalmic products while using this product, wait at least 5 minutes between each product
- to avoid contamination, do not touch tip of container to any surface
- replace cap after each use

- children under 5 years of age: consult a doctor

Other information

- store at 15-25°C (59-77°F)

Inactive ingredients

benzalkonium chloride, boric acid, calcium chloride dihydrate, glycerin, potassium chloride, sodium borate decahydrate, sodium chloride, water for injection. Hydrochloric acid and/or sodium hydroxide may be used to adjust pH.

Questions or comments?

Call: **1-800-553-5340**

PRINCIPAL DISPLAY PANEL - 7.5 mL Carton

BAUSCH + LOMB

NDC 24208-537-25

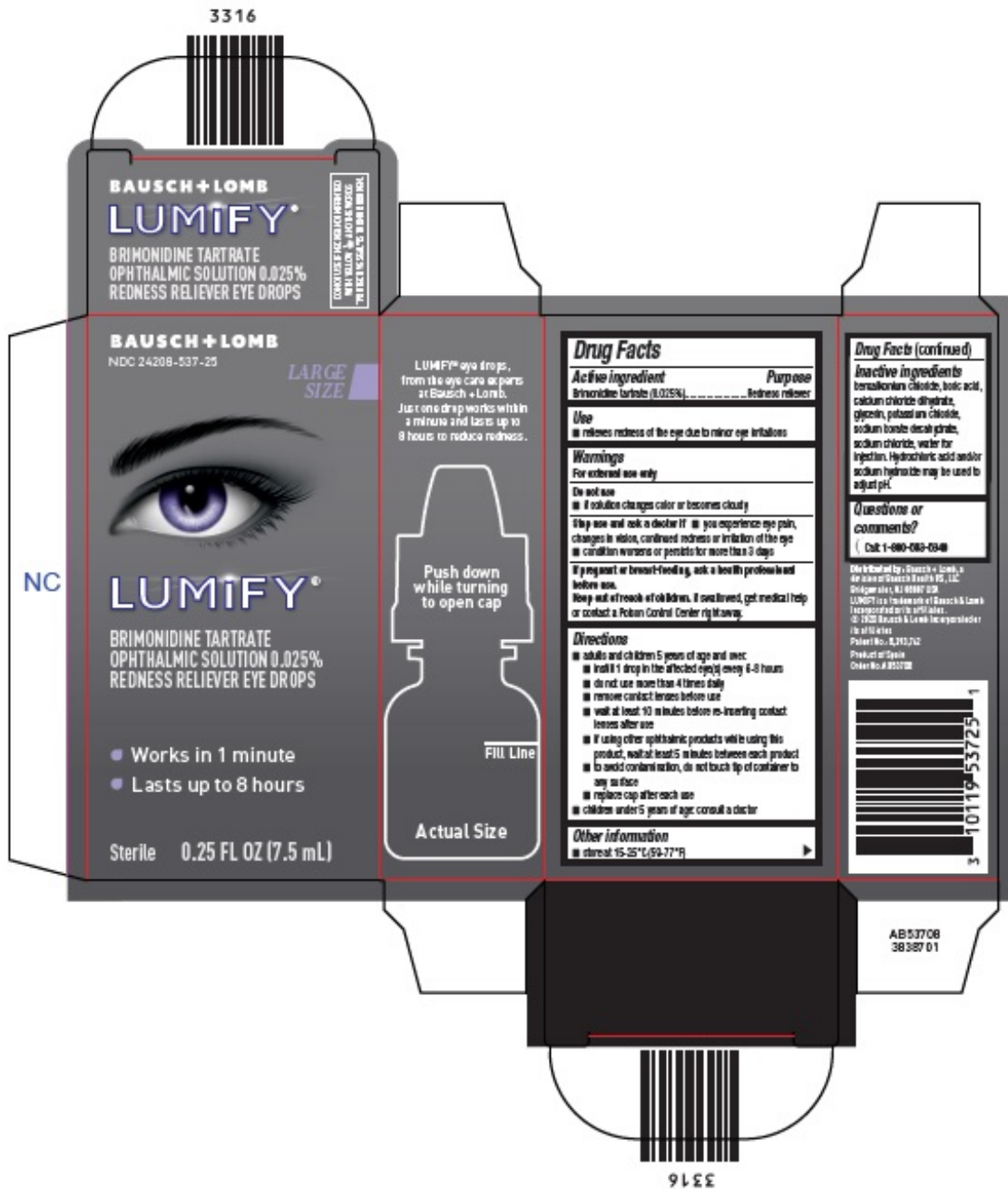
**LARGE
SIZE**

LUMIFY®

**BRIMONIDINE TARTRATE
OPHTHALMIC SOLUTION 0.025%
REDNESS RELIEVER EYE DROPS**

- **Works in 1 minute**
- **Lasts up to 8 hours**

Sterile 0.25 FL OZ (7.5 mL)



LUMIFY REDNESS RELIEVER EYE DROPS

brimonidine tartrate solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:24208-537
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BRIMONIDINE TARTRATE (UNII: 4S9CL2DY2H) (BRIMONIDINE - UNII:E6GNX3HHT E)	BRIMONIDINE TARTRATE	0.25 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BORIC ACID (UNII: R57ZHV85D4)	
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	
GLYCERIN (UNII: PDC6A3C0OX)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:24208-537-25	1 in 1 CARTON	12/22/2017	
1		7.5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
2	NDC:24208-537-08	1 in 1 CARTON	12/22/2017	
2		2.5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
3	NDC:24208-537-99	1 in 1 CARTON	12/22/2017	
3		2.5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
4	NDC:24208-537-75	1 in 1 CARTON	12/22/2017	
4		7.5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
5	NDC:24208-537-10	1 in 1 CARTON	05/01/2019	
5		0.4 mL in 1 AMPULE; Type 0: Not a Combination Product		
6	NDC:24208-537-01	1 in 1 CARTON	05/01/2019	
6		0.4 mL in 1 AMPULE; Type 0: Not a Combination Product		
7	NDC:24208-537-15	2 in 1 CARTON	04/30/2019	
7		7.5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
8	NDC:24208-537-05	1 in 1 CARTON	09/25/2020	
8		5 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
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NDA	NDA208144	12/22/2017	
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Labeler - Bausch & Lomb Incorporated (196603781)

Establishment

Name	Address	ID/FEI	Business Operations
Bausch & Lomb Incorporated		079587625	MANUFACTURE(24208-537)

Establishment

Name	Address	ID/FEI	Business Operations
Laboratoire Unither		574139809	MANUFACTURE(24208-537)

Revised: 7/2021

Bausch & Lomb Incorporated