

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

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***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**

3641



3641

BAUSCH + LOMB  
NDC 24208-537-25

**LARGE  
SIZE**

**LUMIFY**<sup>®</sup>

**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)**

AB53708

3838702

**LUMIFY REDNESS RELIEVER EYE DROPS**

brimonidine tartrate solution/ drops

**Product Information**

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

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>BRIMONIDINE TARTRATE</b> (UNII: 4S9CL2DY2H) (BRIMONIDINE - UNII:E6GNX3HHTE)	BRIMONIDINE TARTRATE	0.25 mg in 1 mL

**Inactive Ingredients**

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

**Packaging**

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
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		

4		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		
9	NDC:24208-537-35	1 in 1 CARTON	05/02/2023	
9		3.5 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
NDA	NDA208144	12/22/2017	

**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: 11/2023

Bausch & Lomb Incorporated