SOOTHE- glycerin and propylene glycol solution/ drops Bausch & Lomb Incorporated

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Soothe Preservative Free Drug Facts

Active ingredients

Glycerin (0.6%)

Propylene glycol (0.6%)

Purpose

Lubricant

Lubricant

Uses

- relieves dryness of the eye
- prevents further irritation

Warnings

For external use only Do not use

- if solution changes color or becomes cloudy
- if single-unit dispenser is not intact

When using this product

- do not touch tip of container to any surface to avoid contamination
- once opened, discard
- do not reuse

Stop use and ask a doctor if

- you experience eye pain, changes in vision, continued redness or irritation of the eve
- condition worsens or persists 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

- to open, completely twist off tab
- instill 1 to 2 drops in the affected eye(s) as needed
- discard dispenser immediately after use

Other information

store at 15 – 25 °C (59 – 77 °F)

Inactive ingredients

boric acid, hydroxyalkylphosphonate, purified water, sodium alginate, sodium borate

Questions?

[phone icon] Call: 1-800-553-5340

Marketed by:

Bausch & Lomb Americas Inc. Bridgewater, NJ 08807 USA

Soothe is a trademark of Bausch & Lomb Incorporated or its affiliates.

© 2022 Bausch & Lomb Incorporated or its affiliates Product of France

Package/Label Principal Display Panel



Bausch & Lomb Soothe®

Lubricant Eye Drops

PRESERVATIVE FREE

Dry Eye Therapy

Moisturizes Comforts Lasts Recommended for LASIK dryness* 30 Single-Use Vials

Sterile 0.02 FL OZ EA (0.6 mL EA)

3839601

SootheEyeDrops.com

Order No. 622219

SOOTHE

glycerin and propylene glycol solution/ drops

Product	Inform	ation
Product		ation

Product Type HUMAN OTC DRUG Item Code (Source) NDC:24208-495

Route of Administration OPHTHALMIC

Active Ingredient/Active Moiety Ingredient Name Basis of Strength GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX) PROPYLENE GLYCOL (UNII: 6DC9Q167V3) (PROPYLENE GLYCOL - UNII:6DC9Q167V3) PROPYLENE GLYCOL 6 mg in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
BORIC ACID (UNII: R57ZHV85D4)	
WATER (UNII: 059QF0KO0R)	
SODIUM ALGINATE (UNII: C269C4G2ZQ)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:24208- 495-28	28 in 1 CARTON	08/26/2010	06/30/2022
1		0.6 mL in 1 VIAL, SINGLE-USE; Type 0: Not a Combination Product		
2	NDC:24208- 495-30	30 in 1 CARTON	05/01/2020	
2		0.6 mL in 1 VIAL, SINGLE-USE; Type 0: Not a Combination Product		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part349	08/26/2010		

Labeler - Bausch & Lomb Incorporated (196603781)

Establishment				
Name	Address	ID/FEI	Business Operations	

Bausch & Lomb Incorporated 114406598 MANUFACTURE(24208-495)

Revised: 1/2023 Bausch & Lomb Incorporated