ADVANCED EYE RELIEF DRY EYE REJUVENATION- glycerin 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.

Drug Facts

Active ingredients

Glycerin (0.3%)

Propylene glycol (1.0%)

Purpose

Lubricant

Lubricant

Uses

- temporary relief of burning and irritation due to dryness of the eye
- prevents further irritation

Warnings

For external use only

Do not use

- if solution changes color or becomes cloudy

When using this product

- do not touch tip of container to any surface to avoid contamination
- remove contact lenses before using
- replace cap after use

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 72 hours

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- instill 1 or 2 drops in the affected eye(s) as needed

Other information

- store at 15–25°C (59–77 °F)
- keep tightly closed
- use before expiration date marked on the carton and bottle

Inactive ingredients

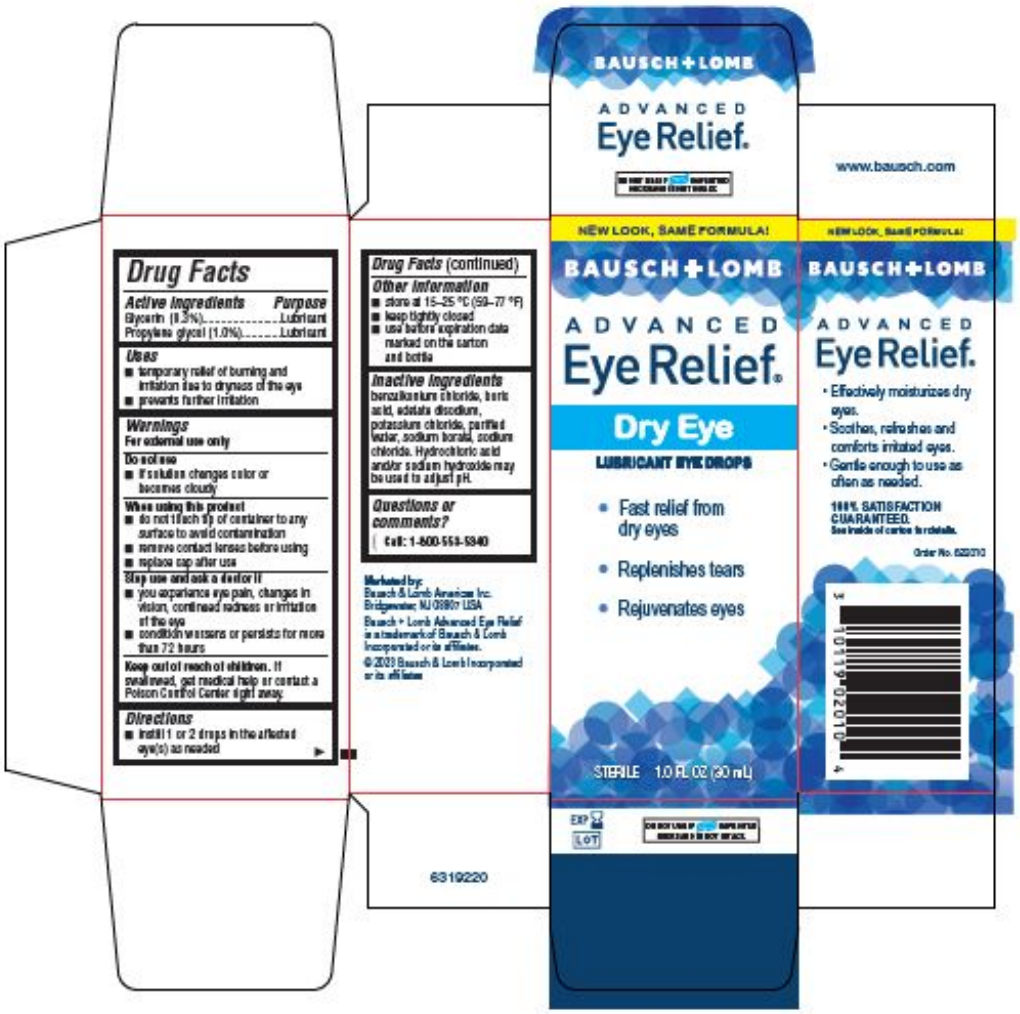
benzalkonium chloride, boric acid, edetate disodium, potassium chloride, purified water, sodium borate, sodium chloride. Hydrochloric acid and/or sodium hydroxide may be used to adjust pH.

Questions or comments?

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

Marketed by:
Bausch & Lomb Americas Inc.
Bridgewater, NJ 08807 USA

Package/Label Principal Display Panel



NEW LOOK, SAME FORMULA!

BAUSCH + LOMB

ADVANCED
Eye Relief®

Dry Eye
LUBRICANT EYE DROPS

- Fast relief from dry eyes
- Replenishes tears
- Rejuvenates eyes

STERILE 1.0 FL OZ (30 mL)

6319220

ADVANCED EYE RELIEF DRY EYE REJUVENATION

glycerin propylene glycol solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:24208-454
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	3 mg in 1 mL
PROPYLENE GLYCOL (UNII: 6DC9Q167V3) (PROPYLENE GLYCOL - UNII:6DC9Q167V3)	PROPYLENE GLYCOL	10 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BORIC ACID (UNII: R57ZHV85D4)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:24208-454-32	1 in 1 CARTON	09/01/2010	
1		15 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:24208-454-30	1 in 1 CARTON	05/01/2017	
2		30 mL in 1 BOTTLE; 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	09/01/2010	

Labeler - Bausch & Lomb Incorporated (196603781)

Establishment

Name	Address	ID/FEI	Business Operations
Bausch & Lomb Incorporated		114406598	MANUFACTURE(24208-454)