BLINK TEARS- polyethylene glycol 400 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

OTC - ACTIVE INGREDIENT SECTION

Polyethylene Glycol 400 0.25%

Purpose

Eye lubricant

Uses

- For the temporary relief of burning, irritation and discomfort due to dryness of the eye or exposure to wind or sun.
- May be used as a protectant against further irritation.

WARNINGS

- For external use only.
- To avoid contamination, do not touch tip of container to any surface. Replace cap after using.
- Do not use if solution changes color or becomes cloudy.

Stop use and ask doctor if:

You experience eye pain, changes in vision, continued redness or irritation of the eye, or if the condition worsens or persists for more than 72 hours.

OTC - KEEP OUT OF REACH OF CHILDREN SECTION

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

Directions

Instill 1 or 2 drops in the affected eye(s) as needed.

Other Information

Use only if tape seals on top and bottom flaps are intact.

RETAIN THIS CARTON FOR FUTURE REFERENCE.

INACTIVE INGREDIENT SECTION

Boric acid, calcium chloride, magnesium chloride, potassium chloride, purified water, sodium borate, sodium chloride, sodium chlorite (OcuPure[®] Brand) as a preservative; Sodium Hyaluronate.

OTC - QUESTIONS SECTION

In the U.S. call 1-800-347-5005 www.yourhealthyeyes.com

Principal Display Panel - 30 mL Bottle Carton

\$3 SAVINGS see inside

blink[®] tears

Dry Eye Lubricating Eye Drops Mild-Moderate

Long-Lasting Relief

Instantly Soothes & Hydrates

1 FL OZ (30 mL) STERILE



BLINK TEARS

polyethylene glycol 400 solution/ drops

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:29943-002	
Route of Administration	OPHTHALMIC			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
Polyethylene Glycol 400 (UNII: B697894SGQ) (Polyethylene Glycol 400 - UNII: B697894SGQ)	Polyethylene Glycol 400	2.5 mg in 1 mL		

Inactive Ingredients		
Ingredient Name	Strength	
Boric Acid (UNII: R57ZHV85D4)		
Calcium Chloride (UNII: M4I0D6VV5M)		
Magnesium Chloride (UNII: 02F3473H9O)		
Potassium Chloride (UNII: 660YQ98I10)		
Water (UNII: 059QF0KO0R)		
SODIUM BORATE (UNII: 91MBZ8H3QO)		
Sodium Chloride (UNII: 451W47IQ8X)		
SODIUM CHLORITE (UNII: G538EBV4VF)		
HYALURONATE SODIUM (UNII: YSE9PPT4TH)		

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:29943- 002-15	1 in 1 CARTON	02/01/2008	
1		15 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
2	NDC:29943- 002-30	1 in 1 CARTON	02/01/2008	
2		30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
3	NDC:29943- 002-02	1 in 1 CARTON	02/01/2008	
3		2 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	
OTC monograph final	part349	02/01/2008		

Labeler - Bausch & Lomb Incorporated (196603781)

Revised: 10/2023 Bausch & Lomb Incorporated