GNP LUBRICANT EYE - polyethylene glycol 400 solution/ drops Amerisource Bergen

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 Purpose

Polyethylene Glycol 400 (0.4%)......Lubricant

Propylene Glycol (0.3%)....Lubricant

Uses

• For the temporary relief of burning and irritation due to dryness of the eye.

Warnings For external use only.

Do not use

- if this product changes color or becomes cloudy.
- if you are sensitive to any ingredient in this product.

When using this product

- Do not touch tip of container to any surface to avoid contamination.
- Replace cap after each use.

Stop use and ask a doctor if

- you feel eye pain.
- changes in vision occur.
- redness or irritation of the eye(s) get worse or lasts 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 to 2 drops in the affected eye(s) as needed
- Children under 6 years of age: Ask a doctor.

Other information

- Store at room temperature.
- Do not use if carton is open or neckband on the bottle is broken or missing.

Inactive ingredients: Boric Avid, Calcium Chloride, Chlorhexidine Gluconate, Hydrochloric Acid, Hypromellose 2910, Magnesium Chloride, Potassium Chloride, Purified Water, Sodim Chloride, Sodium Hydroxide, Zinc Chloride

Distributed By:

Amerisource Bergen Corporation

1300 Morris Drive

Chesterbrook, PA 19087

Made in Korea



Lubricant

Eye Drops

Dry Eye Relief

Long Lasting , Soothing Moisture

0.5 fl oz (15 mL)



ompare to Systane^s active ingredients

NDC 46122-0018-05

STERILE



Lubricant eye drops are specially formulated to protect, refresh, and soothe dry eyes. It offers long lasting relief.

This product is not manufactured or distributed by Alcon, owner of the registered trademark Systane Lubricant Eye Drops.

ABC# 004 - 044



Active ingredients Purpose
Polyethylene Glycol 400 (0.4%) Lubricant
Propylene Glycol (0.3%) Lubricant

Uses

For the temporary relief of burning and irritation due to dryness of the eye.

Warnings For external use only.

Do not use

- if this product changes color or becomes cloudy.
- if you are sensitive to any ingredient in this product.

When using this product

- Do not touch tip of container to any surface to avoid contamination.
- Replace cap after each use.

Stop use and ask a doctor if

- you feel eye pain
- changes in vision occur.
- redness or irritation of the eye(s) gets worse or lasts 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 to 2 drops in the affected eye(s) as needed
 Children under 6 years of age: Ask a doctor.
- Other information
- Store at room temperature.
 Do not use if carton is open or neckband
- Do not use if carton is open or neckband on the bottle is broken or missing.

Inactive ingredients: Boric Acid, Calcium Chloride, Chlorhexidine Gluconate, Hydrochloric Acid, Hypromelloss 2910, Magnesium Chloride, Potassium Chloride, Purified Water, Sodium Chloride, Sodium Hydroxide, Zinc Chloride



Lubricant Eye Drops



Side effects occur. You may report side effects to FDA at 1-800-FDA-1088.

Distributed by: AmerisourceBergen Corporation 1300 Morris Drive Chesterbrook, PA 19087

www.goodneighborpharmacy.com Made in Korea

LOT & EXP.

GNP LUBRICANT EYE

polyethylene glycol 400 solution/ drops

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:46 122-0 18

Route of Administration OPHTHALMIC

Active Ingredient/Active Moiety						
Ingredient Name	Basis of Strength	Strength				
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ) (POLYETHYLENE GLYCOLS - UNII: 3WJQ0SDW1A)	POLYETHYLENE GLYCOL 400	0.4 mg in 1 mg				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3) (PROPYLENE GLYCOL - UNII:6DC9Q167V3)	PROPYLENE GLYCOL	0.3 mg in 1 mg				

Inactive Ingredients		
Ingredient Name	Strength	
BORIC ACID (UNII: R57ZHV85D4)		
CALCIUM CHLO RIDE (UNII: M4I0 D6 VV5M)		
CHLORHEXIDINE GLUCONATE (UNII: MOR8 4MUD8 E)		
HYDRO CHLO RIC ACID (UNII: QTT17582CB)		
MAGNESIUM CHLO RIDE (UNII: 02F3473H9O)		
POTASSIUM CHLORIDE (UNII: 660 YQ98 I10)		
WATER (UNII: 059QF0KO0R)		
SODIUM CHLORIDE (UNII: 451W47IQ8X)		
SODIUM HYDRO XIDE (UNII: 55X04QC32I)		
ZINC CHLORIDE (UNII: 86Q357L16B)		

P	Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:46122-018-05	1 in 1 CARTON					
1		15 mg in 1 BOTTLE					

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph final	part349	06/30/2013				

Labeler - Amerisource Bergen (007914906)

Revised: 6/2013 Amerisource Bergen