GOOD NEIGHBOR PHARMACY LUBRICANT EYE DROPS - polyethylene glycol 400 solution
HANLIM PHARM. CO., LTD.

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 eyes.
- For protection against further irritation.

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 on the bottle is broken or missing

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

Distributed by:

AmerisourceBergen Corporation

1300 Morris Drive

Chesterbrook, PA 19087

Side effects occur.

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

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GOOD NEIGHBOR PHARMACY LUBRICANT EYE DROPS
polyethylene glycol 400 solution

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11716-4060
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ) (POLYETHYLENE GLYCOL - UNII:3WJQ0SDW1A)	POLYETHYLENE GLYCOL 400	0.4 mL in 1 mL
PROPYLENE GLYCOL (UNII: 6DC9Q167V3) (PROPYLENE GLYCOL - UNII:6DC9Q167V3)	PROPYLENE GLYCOL	0.3 mL in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BORIC ACID (UNII: R57ZHV85D4)	
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	
CHLORHEXIDINE GLUCONATE (UNII: MOR84MUD8E)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
HYPROMELLOSE 2910 (4000 CPS) (UNII: RN3152OP35)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
ZINC CHLORIDE (UNII: 86Q357L16B)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11716-4060-9	1 in 1 CARTON		
1		15 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part349	09/20/2010	

Labeler - HANLIM PHARM. CO., LTD. (687986034)

Registrant - UNITED EXCHANGE CORP. (840130579)

Establishment

Name	Address	ID/FEI	Business Operations
HANLIM PHARM. CO., LTD.		687986034	manufacture