GOODSENSE ULTRA LUBRICANT- polyethylene glycol 400, and propylene glycol solution/ drops
Geiss, Destin & Dunn, Inc

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.

Active ingredients **Purpose**

Polyethlene glycol 400 0.4%.....Eye lubricant

Propylene glycol 0.3%.....Eye 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 containation
- 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, persists 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

- shake well before using
- instill 1 or 2 drops in the affected eye(s) as needed

Other information

- store at room temperature

Inactive ingredients

boric acid, calcium chloride, chlorhexidine gluconate, hydrochloric acid, hypromellose, magnesium chloride, potassium chloride, purified water, sodium chloride, sodium hydroxide, zinc chloride

Distributed By:

Geiss, Destin & Dunn, Inc.

Peachtree City, GA 30269

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Made in Korea

GOODSENSE ULTRA LUBRICANT

polyethylene glycol 400, and propylene glycol solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50804-821
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	4 mg in 1 mL
PROPYLENE GLYCOL (UNII: 6DC9Q167V3) (PROPYLENE GLYCOL - UNII:6DC9Q167V3)	PROPYLENE GLYCOL	3 mg 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, UNSPECIFIED (UNII: 3NXW29V3WO)	
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:50804-821-15	1 in 1 BOX	05/18/2017	
1		15 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	05/18/2017	

Labeler - Geiss, Destin & Dunn, Inc (076059836)

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