LUBRICANT EYE DROPS LONG LASTING- polyethylene glycol 400, propylene glycol liquid Nexus Optical 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.

Lubricant Eye Drops Long Lasting

Drug Facts

Active ingredients

Polyethylene glycol 400 0.4%

Propylene glycol 0.3%

Purposes

Lubricant

Use

• 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

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

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

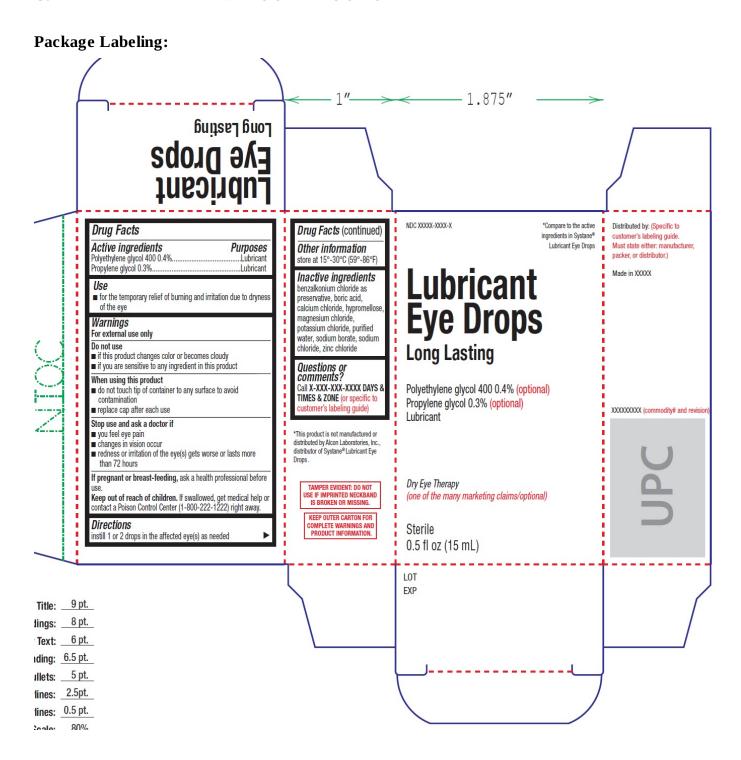
Other information

Inactive ingredients

benzalkonium chloride as preservative, boric acid, calcium chloride, hypromellose, magnesium chloride, potassium chloride, purified water, sodium borate, sodium chloride, zinc chloride

Questions or comments?

Call X-XXX-XXXX DAYS & TIMES & ZONE





LUBRICANT EYE DROPS LONG LASTING

polyethylene glycol 400, propylene glycol liquid

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73450-001	
Route of Administration	ОРНТНАЬМІС			

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	
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7)		
BORIC ACID (UNII: R57ZHV85D4)		
CALCIUM CHLO RIDE (UNII: M4I0 D6 VV5M)		
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29 V3WO)		
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)		
POTASSIUM CHLORIDE (UNII: 660 YQ98 I10)		
WATER (UNII: 059QF0KO0R)		
SODIUM BORATE (UNII: 91MBZ8H3QO)		
SODIUM CHLORIDE (UNII: 451W47IQ8X)		
ZINC CHLORIDE (UNII: 86Q357L16B)		

P	ackaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date

1 NDC:73450-001-15	1 in 1 BOX	04/20/2020		
1	15 mL in 1 BOTTLE; Type 0: Not a Combination Product			
Marketing Information				
Marketing Catego	ry Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not fi	nal part349	04/20/2020		

Labeler - Nexus Optical Inc (107289657)

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