LUBRICANT EYE DROPS ULTRA- polyethylene glycol, propylene glycol liquid Discount Drug Mart

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

Polyethylene Glycol 400

Propylene Glycol

Purpose

Lubricant

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, 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

 $benzal konium\,chloride,\,boric\,acid,\,hydrochloric\,acid,\,hydroxyethyl\,cellulose,\,potassium\,chloride,\,purified\,water,\,sodium\,chloride,\,sodium\,hydroxide$

Package label

02401 DDM Ultra Lub Eye Drop .33oz 2pk Ind.box (dim. 65 x 32 x 108 mm)





LOT PU16001 EXP 06/2018



Read Carton For Uses And All Warnings. Active ingredients: Polyethylene Glycol 400 04%, Prop/ene Glycol 0.3% Warnings: Keep and of reach of bildren. If swellowed, per metical help or contact a Directions: Brake well before using. = Shake well before using.

LUBRICANT EYE DROPS ULTRA

polyethylene glycol, propylene glycol liquid

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53943-240	
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	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)		
BORIC ACID (UNII: R57ZHV85D4)		
HYDRO CHLORIC ACID (UNII: QTT17582CB)		
POTASSIUM CHLORIDE (UNII: 660 YQ98110)		
SODIUM CHLORIDE (UNII: 451W47IQ8X)		
SO DIUM HYDRO XIDE (UNII: 55X04QC32I)		
WATER (UNII: 059QF0KO0R)		

I	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:53943-240- 02	2 in 1 BOX	11/11/20 16		
1		10 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product			

Marketing Info	Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph final	part349	11/11/2016			

Labeler - Discount Drug Mart (047741335)

Revised: 4/2017 Discount Drug Mart