

CVS LUBRICANT EYE DROPS ORIGINAL- polyethylene glycol 400, propylene glycol solution
CVS

CVS Lubricant Eye Drops Original 30 ct. (PLD)

Active ingredients

Polyethylene glycol 400.....0.4%
Propylene glycol.....0.3%

Purposes

Polyethylene glycol 400....Lubricant
Propylene glycol.....Lubricant

Use

- for the temporary relief of burning and irritation of the eye due to dryness of the eye

Warnings

For external use only

Do not use

- if this solution changes 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

- store at 15°-25° (59°-77 F)
- use only if single-use container is intact
- use before expiration date marked on container
- **RETAIN THIS CARTON FOR FUTURE REFERENCE**

Inactive ingredients

boric acid, hydrochloric acid**, hypromellose, potassium chloride, purified water, sodium chloride, sodium hydroxide**

**May contain these ingredients to adjust pH.

CVS LUBRICANT EYE DROPS ORIGINAL			
polyethylene glycol 400, propylene glycol solution			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69842-746
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	0.4 g in 100 mL
PROPYLENE GLYCOL (UNII: 6DC9Q167V3) (PROPYLENE GLYCOL - UNII:6DC9Q167V3)		PROPYLENE GLYCOL	0.3 g in 100 mL
Inactive Ingredients			
Ingredient Name			Strength
HYDROCHLORIC ACID (UNII: QTT17582CB)			
SODIUM HYDROXIDE (UNII: 55X04QC32I)			

WATER (UNII: 059QF0KO0R)	
BORIC ACID (UNII: R57ZHV85D4)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69842-746-01	30 in 1 CARTON	02/12/2021	
1		0.4 mL in 1 VIAL, DISPENSING; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M018	02/12/2021	

Labeler - CVS (062312574)

Registrant - K.C. Pharmaceuticals, Inc. (174450460)

Establishment

Name	Address	ID/FEI	Business Operations
Unimed Pharmaceuticals, Inc.		689852052	manufacture(69842-746) , pack(69842-746) , label(69842-746)