

SUNMARK LUBRICANT EYE DROPS- polypropylene glycol 400, propylene glycol solution/ drops
Strategic Sourcing Services LLC

Sunmark Lubricant Eye Drops (PLD)

Active Ingredients

Polyethylene glycol 400 0.4%, Propylene glycol 0.3%

Purpose

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 the tip of the 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 or 2 drops in the affected eye(s) as needed
- children under 6 years of age: ask a doctor

Other information

- **RETAIN THIS CARTON FOR FUTURE REFERENCE**

- Store at room temperature

Inactive ingredients

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



SUNMARK LUBRICANT EYE DROPS

polypropylene glycol 400, propylene glycol solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49348-947
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
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BORIC ACID (UNII: R57ZHV85D4)	
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
ZINC CHLORIDE (UNII: 86Q357L16B)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49348-947-29	1 in 1 CARTON	12/01/2010	
1		15 mL in 1 BOTTLE, PLASTIC; 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	12/01/2010	

Labeler - Strategic Sourcing Services LLC (116956644)

Registrant - KC Pharmaceuticals, Inc (174450460)

Establishment

Name	Address	ID/FEI	Business Operations
KC Pharmaceuticals, Inc.		174450460	manufacture(49348-947) , pack(49348-947) , label(49348-947)

Revised: 12/2023

Strategic Sourcing Services LLC