DG LUBRICANT EYE DROPS- polyethylene glycol 400, propylene glycol solution
Dolgencorp LLC

DG Lubricant Eye Drops (PLD)

Active ingredients

Polyethylene glycol 400 0.4%

Propylene glycol 0.3%

Active ingredients

Polyethylene glycol .....Lubricant

Propylene glycol.....Lubricant

Uses

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

Warnngs

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 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

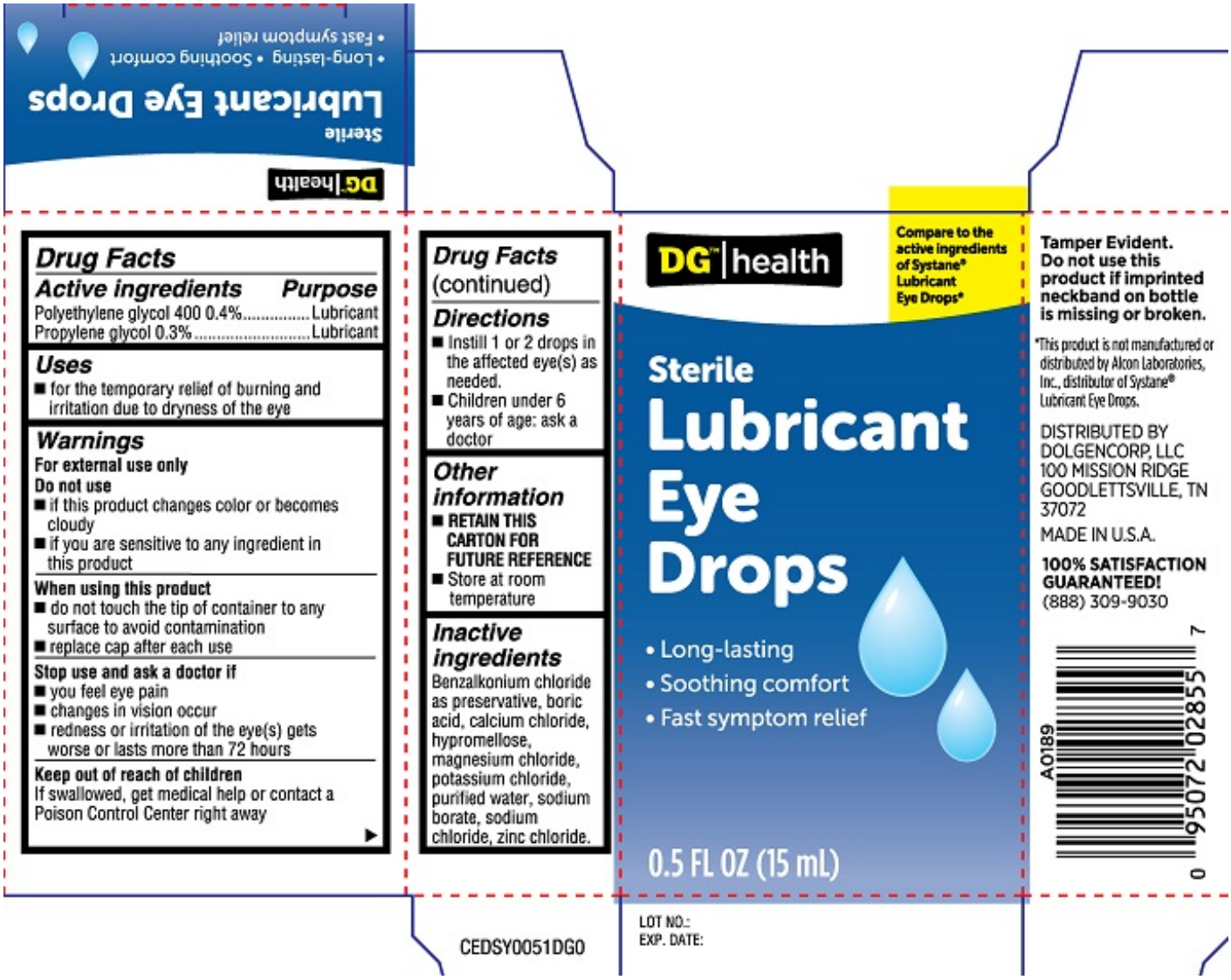
- Instill 1 or 2 drops in the affected eye(s) as needed
- Children under 6 years of age: ask a doctor

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



DG LUBRICANT EYE DROPS

polyethylene glycol 400, propylene glycol solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55910-364
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
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
BORIC ACID (UNII: R57ZHV85D4)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
ZINC CHLORIDE (UNII: 86Q357L16B)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55910-364-01	1 in 1 CARTON	01/27/2020	
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 Drug	M018	01/27/2020	

Labeler - Dolgencorp LLC (068331990)

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

Establishment

Name	Address	ID/FEI	Business Operations
K.C. Pharmaceuticals, Inc.		174450460	manufacture(55910-364) , pack(55910-364) , label(55910-364)

Revised: 12/2023 Dolgencorp LLC