GOODSENSE ULTRA LUBRICANT EYE DROPS- polyethylene glycol 400, propylene glycol solution
Geiss, Destin & Dunn, Inc.

GoodSense Ultra Lubricant Eye Drops (PLD)

Active ingredients

Polyethylene glycol 400.....0.4%

Propylene glycol.....0.3%

Purpose

Polyethylene glycol 400.....Lubricant

Propylene glycol.....Lubricant

Use

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

Warnings

For external use only

When using this product

- do not touch tip of container to any surface to avoid contamination
- replace cap after each use

Do not use

- if this product changes color or becomes cloudy
- if you are sensitive to any ingredient in this product

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

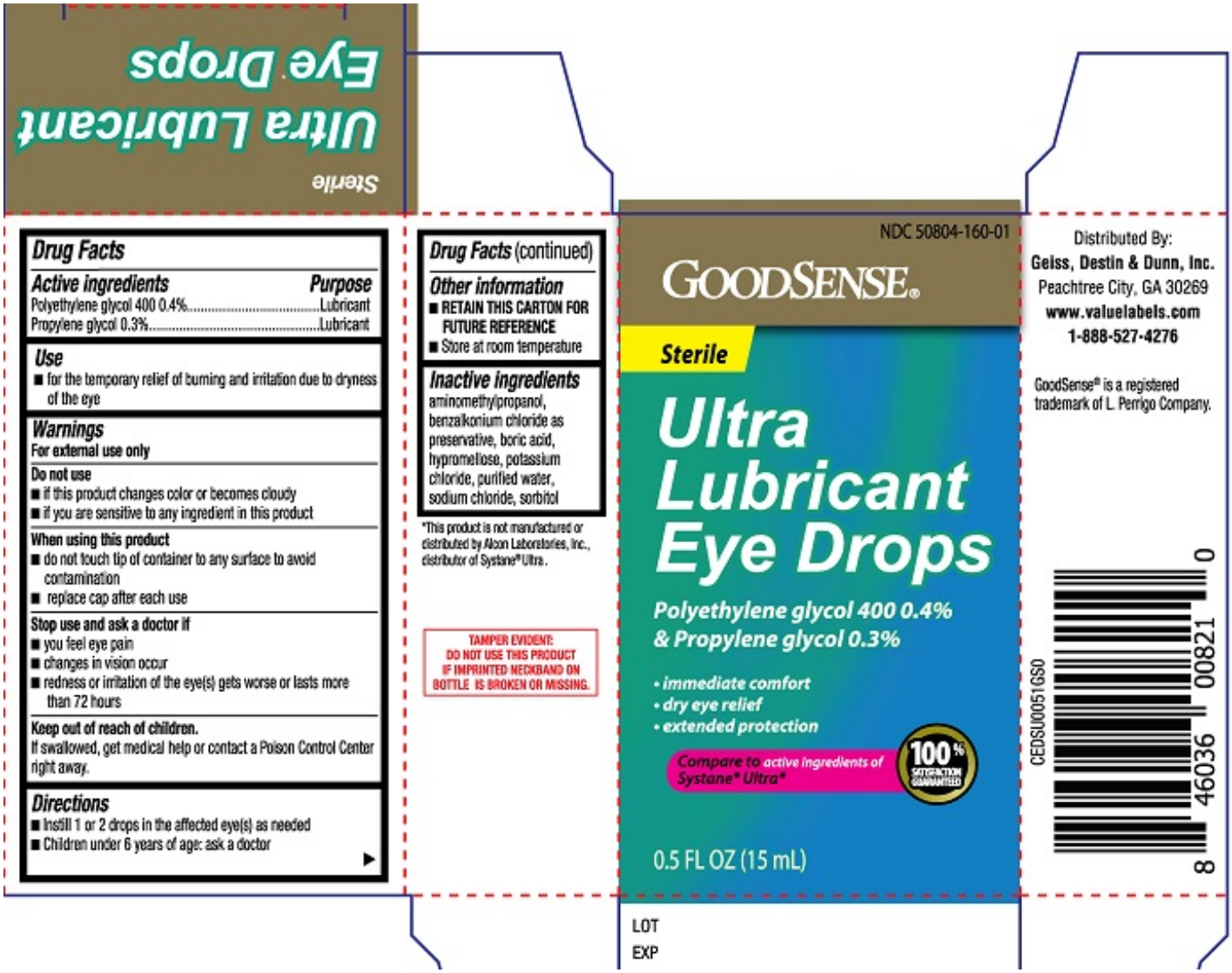
- Instill 1 or 2 drops in the affected eye(s) as needed
- Children under 6 years of age: ask a doctor

Other information

- RETAIN THIS CARTON FOR FUTURE REFERENCE
- Store at room temperature

Inactive ingredients

aminomethylpropanol, benzalkonium chloride as preservative, boric acid, hypromellose, potassium chloride, sorbitol



GOODSENSE ULTRA LUBRICANT EYE DROPS

polyethylene glycol 400, propylene glycol solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50804-160
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
PROPYLENE GLYCOL (UNII: 6DC9Q167V3) (PROPYLENE GLYCOL - UNII:6DC9Q167V3)	PROPYLENE GLYCOL	0.3 g in 100 mL
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ) (POLYETHYLENE GLYCOL 400 - UNII:B697894SGQ)	POLYETHYLENE GLYCOL 400	0.4 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**

AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BORIC ACID (UNII: R57ZHV85D4)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SORBITOL (UNII: 506T60A25R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50804-160-01	1 in 1 CARTON	01/03/2020	
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	01/03/2020	

Labeler - Geiss, Destin & Dunn, Inc. (076059836)

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

Establishment

Name	Address	ID/FEI	Business Operations
K.C. Pharmaceuticals, Inc		174450460	manufacture(50804-160) , pack(50804-160) , label(50804-160)

Revised: 12/2025 Geiss, Destin & Dunn, Inc.