# LUBRICANT EYE DROPS- propylene gycol 0.6% solution/ drops Solis Pharmaceuticals Inc.

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#### **Sterile Lubricant Eye Drops**

### Active ingredient

Propylene Glycol 0.6%

## Purpose

Lubricant

#### Uses

- for the temporary relief of burning and irritation due to dryness of the eye
- for the temporary relief of discomfort due to minor irritations of the eye or to exposure to wind or sun
- for use as a protectant against further irritation or to relieve dryness of the eye
- for use as a lubricant to prevent further irritation or to relieve 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 experience any of the following:

- eye pain,
- changes in vision,
- continued redness or irritation of the eye,
- condition worsens or persists for 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.

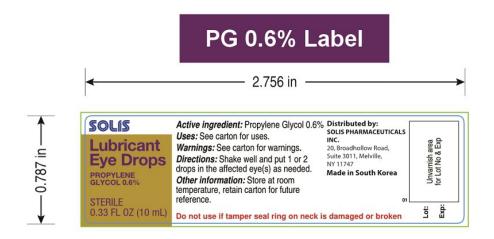
#### Other information

- store at room temperature
- retain carton for future reference

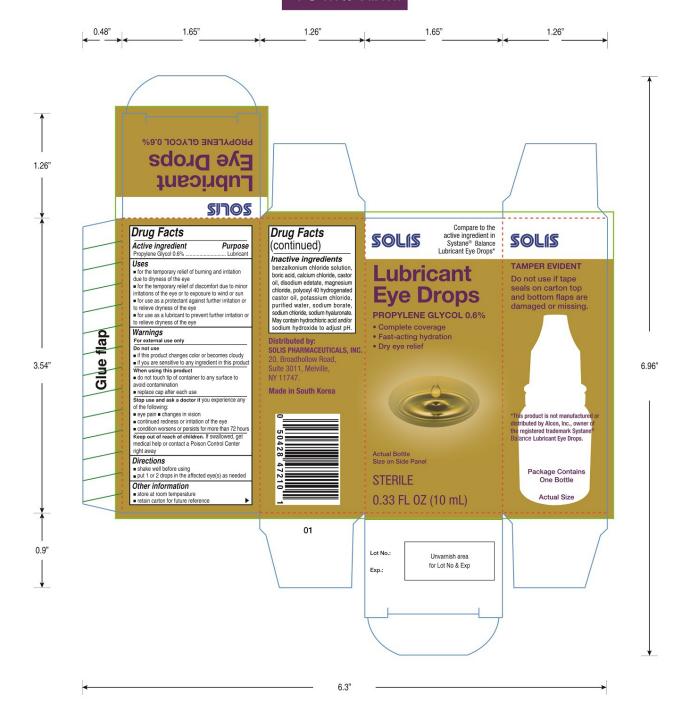
## Inactive ingredients

benzalkonium chloride solution, boric acid, calcium chloride, castor oil, disodium edetate, magnesium chloride, polyoxyl 40 hydrogenated castor oil, potassium chloride, purified water, sodium borate, sodium chloride, sodium hyaluronate. May contain hydrochloric acid and/or sodium hydroxide to adjust pH.

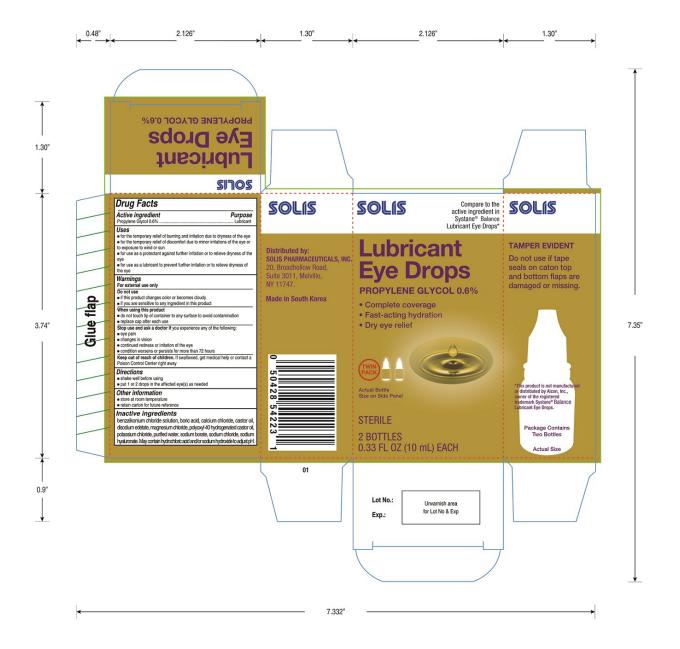
#### PRINCIPAL DISPLAY PANEL



#### PG 0.6% Carton



# PG 0.6 -Twin Pack Carton



## **LUBRICANT EYE DROPS**

propylene gycol 0.6% solution/ drops

<b>Product</b>	Inform	ation
Product		ation

Product Type HUMAN OTC DRUG Item Code (Source) NDC:85185-002

**Route of Administration** OPHTHALMIC

# Active Ingredient/Active Moiety Ingredient Name Basis of Strength PROPYLENE GLYCOL (UNII: 6DC9Q167V3) (PROPYLENE GLYCOL - UNII:6DC9Q167V3) PROPYLENE GLYCOL 6 mg in 1 mL

Inactive Ingredients				
Ingredient Name	Strength			
EDETATE DISODIUM (UNII: 7FLD91C86K)				
BORIC ACID (UNII: R57ZHV85D4)				
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)				
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)				
POTASSIUM CHLORIDE (UNII: 660YQ98I10)				
WATER (UNII: 059QF0KO0R)				
SODIUM BORATE (UNII: 91MBZ8H3QO)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
HYDROCHLORIC ACID (UNII: QTT17582CB)				
SODIUM HYDROXIDE (UNII: 55X04QC32I)				
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)				

Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:85185- 002-10	1 in 1 CARTON	03/01/2025			
1		10 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product				
2	NDC:85185- 002-20	2 in 1 CARTON	03/01/2025			
2		10 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	03/01/2021			

# Labeler - Solis Pharmaceuticals Inc. (119380963)

Revised: 3/2025 Solis Pharmaceuticals Inc.