

**LUBRICANT EYE DROPS- carboxymethylcellulose sodium solution/ drops  
KC Pharmaceuticals, Inc.**

-----  
**KC Pharmaceuticals (MFG) labeler NDC for Discount Drug Mart Lubricant Eye  
Drops 5 ct (PLD) (NDC 53943-510)**

Carboxymethylcellulose sodium 0.5%

Eye Lubricant

- for the temporary relief of burning, irritation, and discomfort due to dryness of the eye or exposure to wind or sun
- may be used as a protectant against further irritation

Keep out of reach of children. If accidentally swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) immediately.

Directions

- To open, pull and twist tab to remove
- Instill 1 or 2 drops in the affected eye(s) as needed and discard container

Calcium Chloride, \*\*Hydrochloric acid, Magnesium Chloride, Potassium Chloride, Purified water, Sodium Chloride, \*\*Sodium Hydroxide, Sodium Lactate. \*\*May contain these ingredients to adjust pH.

Warnings

**For external use only**

**Do not use this product if:**

- solution changes color or becomes cloudy

**When using this product**

- do not reuse
- once opened, discard
- to avoid contamination, do not touch tip of container to any surface
- do not touch unit-dose to eye

**Stop use and ask a doctor if you experience any of the following**

- you experience eye pain
- changes in vision occur
- redness or irritation of the eye continues
- redness or irritation of the eye gets worse or lasts more than 72 hours



## LUBRICANT EYE DROPS

carboxymethylcellulose sodium solution/ drops

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:55651-510
<b>Route of Administration</b>	OPHTHALMIC		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>CARBOXYMETHYLCELLULOSE SODIUM</b> (UNII: K679OBS311) (CARBOXYMETHYLCELLULOSE - UNII:05JZ17B19X)	CARBOXYMETHYLCELLULOSE SODIUM	5 mg in 1 mL

## Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>CALCIUM CHLORIDE</b> (UNII: M4I0D6VV5M)	
<b>MAGNESIUM CHLORIDE</b> (UNII: 02F3473H9O)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>POTASSIUM CHLORIDE</b> (UNII: 660YQ98I10)	
<b>SODIUM HYDROXIDE</b> (UNII: 55X04QC32I)	
<b>SODIUM LACTATE</b> (UNII: TU7HW0W0QT)	
<b>HYDROCHLORIC ACID</b> (UNII: QTT17582CB)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55651-510-01	5 in 1 BOX	02/25/2026	
1		0.4 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	02/25/2026	

**Labeler** - KC Pharmaceuticals, Inc. (174450460)

Revised: 2/2026

KC Pharmaceuticals, Inc.