

**CVS LUBRICANT EYE DROPS- carboxymethylcellulose sodium solution/ drops  
CVS Pharmacy, Inc.**

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**CVS Lubricant Eye Drops 70ct (PLD)**

**Active ingredient**

Carboxymethylcellulose sodium 0.5%

**Purpose**

Lubricant

**Uses**

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

**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 tip to eye

**Stop use and ask a doctor if**

- you experience eye pain
- changes in vision occur
- redness or irritation of the eye continues
- redness or irritation of the eye worsens or persists for more than 72 hours

**Keep out of the reach of children.**

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

**Directions**

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

- if used for post-operative (e.g., LASIK) dryness and discomfort, follow your eye doctor's instructions

**Other information**

- store at 15°-25°C (59°-77°F)
- use only if single-use container is intact
- use before expiration date marked on container
- RETAIN THIS CARTON FOR FUTURE REFERENCE

**Inactive ingredients**

calcium chloride, magnesium chloride, potassium chloride, purified water, sodium chloride and sodium lactate. May contain sodium hydroxide and/or hydrochloric acid to adjust pH.

**CVS Lubricant Eye Drops Preservative Free 70 ct**



## CVS LUBRICANT EYE DROPS

carboxymethylcellulose sodium solution/ drops

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:69842-804
<b>Route of Administration</b>	OPHTHALMIC		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	CARBOXYMETHYLCELLULOSE	0.5 g

(CARBOXYMETHYLCELLULOSE - UNII:05JZ17B19X) SODIUM in 100 mL

### Inactive Ingredients

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

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69842-804-01	70 in 1 BOX	07/30/2019	
1		0.4 mL in 1 VIAL, SINGLE-USE; 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	07/30/2019	

**Labeler** - CVS Pharmacy, Inc. (062312574)

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

### Establishment

Name	Address	ID/FEI	Business Operations
KC Pharmaceuticals, Inc.		174450460	pack(69842-804) , label(69842-804)

### Establishment

Name	Address	ID/FEI	Business Operations
Unimed		689852052	manufacture(69842-804)