

## **LUBRICANT EYE RITE AID- carboxymethylcellulose sodium gel RITE AID**

*Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.*

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### **Lubricant Gel Drops**

#### **Drug Facts**

##### **Active ingredient**

Carboxymethylcellulose sodium 1%

##### **Purpose**

Eye lubricant

##### **Uses**

- 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.

##### **Warnings**

- **For external use only.**
- **To avoid contamination, do not touch tip of container to any surface. Replace cap after using.**
- **If solution changes color or becomes cloudy, do not use.**

**Stop use and ask a doctor if** you experience eye pain, changes in vision, continued redness or irritation of the eye, or if the 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**

Instill 1 or 2 drops in the affected eye(s) as needed.

##### **Other information**

- Use before expiration date marked on container.
- Discard 30 days after opening.

- Store at 59°-86°F (15°-30°C).
- RETAIN THIS CARTON FOR FUTURE REFERENCE.

### Inactive ingredients

Boric acid; calcium chloride; magnesium chloride; potassium chloride; purified water; stabilized oxychloro complex; and sodium chloride.

May contain hydrochloric acid and/or sodium hydroxide to adjust pH



## LUBRICANT EYE RITE AID

carboxymethylcellulose sodium gel

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:11822-9706
<b>Route of Administration</b>	OPHTHALMIC		

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>CARBOXYMETHYLCELLULOSE SODIUM</b> (UNII: K679OBS311) (CARBOXYMETHYLCELLULOSE - UNII:05JZI7B19X)	CARBOXYMETHYLCELLULOSE SODIUM	10 mg in 1 mL

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
<b>BORIC ACID</b> (UNII: R57ZHV85D4)	
<b>CALCIUM CHLORIDE</b> (UNII: M4I0D6VV5M)	
<b>MAGNESIUM CHLORIDE</b> (UNII: 02F3473H9O)	
<b>POTASSIUM CHLORIDE</b> (UNII: 660YQ98I10)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM BORATE</b> (UNII: 91MBZ8H3QO)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>SODIUM CHLORITE</b> (UNII: G538EBV4VF)	

**Packaging**

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:11822-9706-5	1 in 1 CARTON	10/03/2020	
1		15 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		

**Marketing Information**

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC monograph final	part349	10/03/2020	

**Labeler** - RITE AID (014578892)**Registrant** - Velocity Pharma LLC (962198409)

Revised: 6/2022

RITE AID