

**CARBOXYMETHYLCELLULOSE SODIUM EYE DROPS 0.5%-  
carboxymethylcellulose sodium eye drops 0.5% for solution  
Aurolab**

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**Active ingredient**

Carboxymethyl cellulose sodium Eye Drops 0.5% w/v

**DIRECTIONS FOR USE**

- Instill 1or 2 drops in the affected eye, as needed

**INACTIVE INGREDIENT**

- 1.Boric acid
- 2.Calcium chloride
- 3.Glycerin
- 4.Magnesium chloride
- 5.Mannitol
- 6.Potassium chloride
- 7.Purified water
- 8.Stablized oxy cholro complex
- 9.Sodium tetra borate
- 10.Sodium hyaluronate
- 11.Sodium citrate

**Tamper Protection**

- For your protection a tamper evident ring is attached to the bottlecap
- Upon opening, this will separate from the cap and can be discarded
- Use only if this ring is present and attached when the bottle is first opened

**Use**

For use as a lubricant to prevent further irritation or to relieve dryness of the eye

**Questions**

Call. 1-800-103-7321

E-mail : [info@aurolab.com](mailto:info@aurolab.com)

Web : [www.aurolab.com](http://www.aurolab.com)

## **Keep out of reach of children**

If swallowed get medical help or contact a Poison Control Center right away

## **Stop use and ask a doctor if**

- 1.Excessive watering of the eye
- 2.Burning, stinging,foreign body sensation Continued redness (or) irritation of the eye, unusual eye secretions, change in vision

## **Do not use**

- 1.If you are sensitive to any ingredient in this product
- 2.If solution changes color or becomes cloudy

## **Warnings**

For external use only

## **Indication & usage**

Do not touch the nozzle tip to any surface since this may contaminate the solution  
Remove contact lenses before use Should not use at the same time as other ophthalmic drugs  
Replace cap after using

## **Dose**

Instill 1 or 2 drops in the affected eyes as needed

## **Eye lubricant**

Eye lubricant

## **Carton**



## CARBOXYMETHYLCELLULOSE SODIUM EYE DROPS 0.5%

carboxymethylcellulose sodium eye drops 0.5% for solution

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:16030-501
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311) (CARBOXYMETHYLCELLULOSE - UNII:05JZ17B19X)	CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED	5 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>BORIC ACID</b> (UNII: R57ZHV85D4)	
<b>SODIUM CHLORITE</b> (UNII: G538EBV4VF)	
<b>CALCIUM CHLORIDE</b> (UNII: M410D6VV5M)	
<b>MAGNESIUM CHLORIDE</b> (UNII: 02F3473H9O)	
<b>POTASSIUM CHLORIDE</b> (UNII: 660YQ98I10)	
<b>SODIUM CITRATE</b> (UNII: 1Q73Q2JULR)	

**MANNITOL** (UNII: 3OWL53L36A)

**GLYCERIN** (UNII: PDC6A3C0OX)

**PEG-9 DIGLYCIDYL ETHER/SODIUM HYALURONATE CROSSPOLYMER** (UNII: 788QAG3W8A)

**SODIUM BORATE** (UNII: 91MBZ8H3QO)

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:16030-501-05	5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	09/20/2022	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M018	09/20/2022	

**Labeler - Aurolab** (677319965)

### Establishment

Name	Address	ID/FEI	Business Operations
Aurolab		677319965	manufacture(16030-501)

Revised: 1/2026

Aurolab