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

Active ingredient

Carboxymethyl cellulose sodium Eye Drops 0.5% w/v

DIRECTIONS FOR USE

- lnstill 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
Web : 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			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:16030-501
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311) (CARBOXYMETHYLCELLULOSE - UNII:05JZI7B19X)	CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED	5 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
BORIC ACID (UNII: R57ZHV85D4)	
SODIUM CHLORITE (UNII: G538EBV4VF)	
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
SODIUM CITRATE (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: 4/2025 Aurolab