

**AUROVISC- hypromellose ophthalmic solution 2% w/v solution**  
**Aurolab**

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**ACTIVE INGREDIENT**

Hypromellose USP 2% w/v

**INACTIVE INGREDIENT**

1. Acetic acid 1%
2. Calcium chloride
3. Citric acid 0.1465%
4. Magnesium Chloride
5. Sodium chloride
6. Sodium acetate,
7. Sodium Citrate
8. Potassium chloride
9. Purified water.

**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**

1. Transient blurring of vision
2. Ocular discomfort or irritation
3. Matting or Stickness of eyelashes
4. Photophobia
5. Hypersensitivity or edema of the eyelids

**DO NOT USE**

1. If the solution becomes dark brown or any floating particles are observed.
2. If you are sensitive to any ingredient in this product

## **WARNINGS**

For External use only

## **INDICATIONS AND USAGE**

Do not use if package is damaged

Discard after a single use

Do not freeze

Do not resterilize

## **Purpose**

Lubricant

## **Dose**

Instill 1 or 2 drops in the affected eyes as needed

## **PACKAGE CARTON**



## AUROVISC

hypromellose ophthalmic solution 2% w/v solution

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:16030-301
<b>Route of Administration</b>	OPHTHALMIC		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>HYPROMELLOSE 2208 (15000 MPA.S)</b> (UNII: Z 78RG6M2N2) (HYPROMELLOSE 2208 (15000 MPA.S) - UNII:Z 78RG6M2N2)	HYPROMELLOSE 2208 (15000 MPA.S)	20 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>SODIUM ACETATE</b> (UNII: 4550K0SC9B)	

<b>SODIUM CITRATE</b> (UNII: 1Q73Q2JULR)	
<b>POTASSIUM CHLORIDE</b> (UNII: 660YQ98I10)	
<b>WATER</b> (UNII: 059QF0K00R)	
<b>ACETIC ACID</b> (UNII: Q40Q9N063P)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>MAGNESIUM CHLORIDE</b> (UNII: 02F3473H9O)	
<b>CALCIUM CHLORIDE</b> (UNII: M4I0D6VV5M)	
<b>CITRIC ACID ACETATE</b> (UNII: DSO12WL7AU)	

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### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:16030-301-02	2 mL in 1 SYRINGE, GLASS; Type 1: Convenience Kit of Co-Package	09/26/2022	

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### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M018	09/26/2022	

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### Labeler - Aurolab (677319965)

### Establishment

Name	Address	ID/FEI	Business Operations
Aurolab DBA Aurolab Trust		677319965	manufacture(16030-301)

Revised: 1/2026

Aurolab