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

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
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

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

INDIACATIONS AND USAGE

Do not use if package is damaged
Discard after a single use
Do not freeze
Do not resterilize

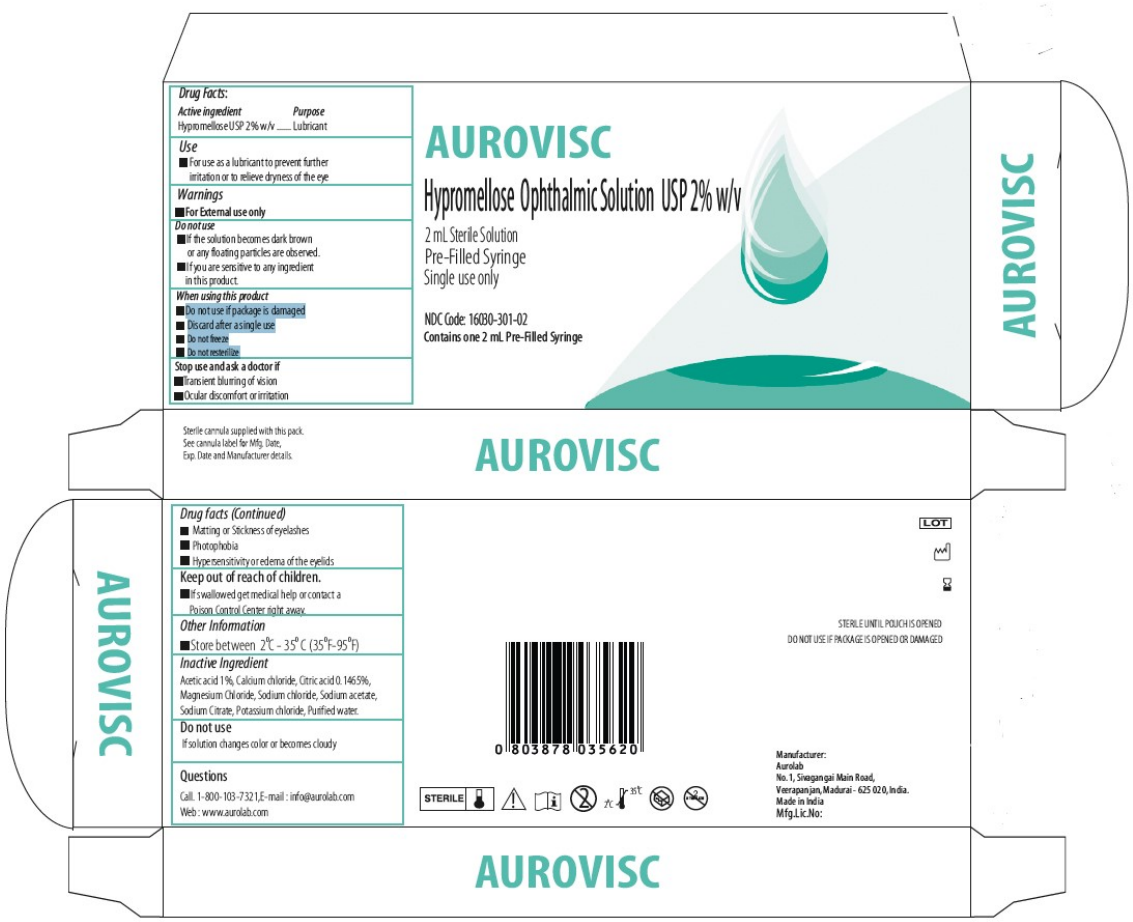
Purpose

Lubricant

Dose

Instill 1 or 2 drops in the affected eyes as needed

PACKAGE CARTON



AUROVISC OE

hypromellose ophthalmic solution 2% w/v solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:16030-304
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
HYPROMELLOSE 2208 (15000 MPA.S) (UNII: Z78RG6M2N2) (HYPROMELLOSE 2208 (15000 MPA.S) - UNII:Z78RG6M2N2)	HYPROMELLOSE 2208 (15000 MPA.S)	20 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
SODIUM ACETATE (UNII: 4550K0SC9B)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	

SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	
CITRIC ACID ACETATE (UNII: DSO12WL7AU)	
ACETIC ACID (UNII: Q40Q9N063P)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:16030-304-02	2 mL in 1 SYRINGE, GLASS; Type 1: Convenience Kit of Co-Package	04/01/2025	12/31/2027

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M018	04/01/2025	12/31/2027

Labeler - Aurolab (677319965)

Establishment

Name	Address	ID/FEI	Business Operations
Aurolab		677319965	manufacture(16030-304)

Revised: 3/2025 Aurolab