HYPROMELLOSE EYE DROPS 0.7%- hypromellose eye drops 0.7% for solution Aurolab

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.

Active Ingredient

Hypromellose 0.7% BP w/v

DIRECTIONS FOR USE

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

INACTIVE INGREDIENT

1. Benzalkonium Chloride
2. Borax
3. Boric acid
4. EDTA disodium salt
5. Potassium chloride
6. Purified water
7. Sodiumchloride

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

Email: 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

ASK DOCTOR

- If you experience eye pain
- Change in vision
- Continued redness(or) irritation of the eye
- Condition worsens or persists for more than 72 hours

DO NOT USE

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

Dosage

Instill 1 or 2 drops in the affected eyes as needed

Warnings

For External Use Only

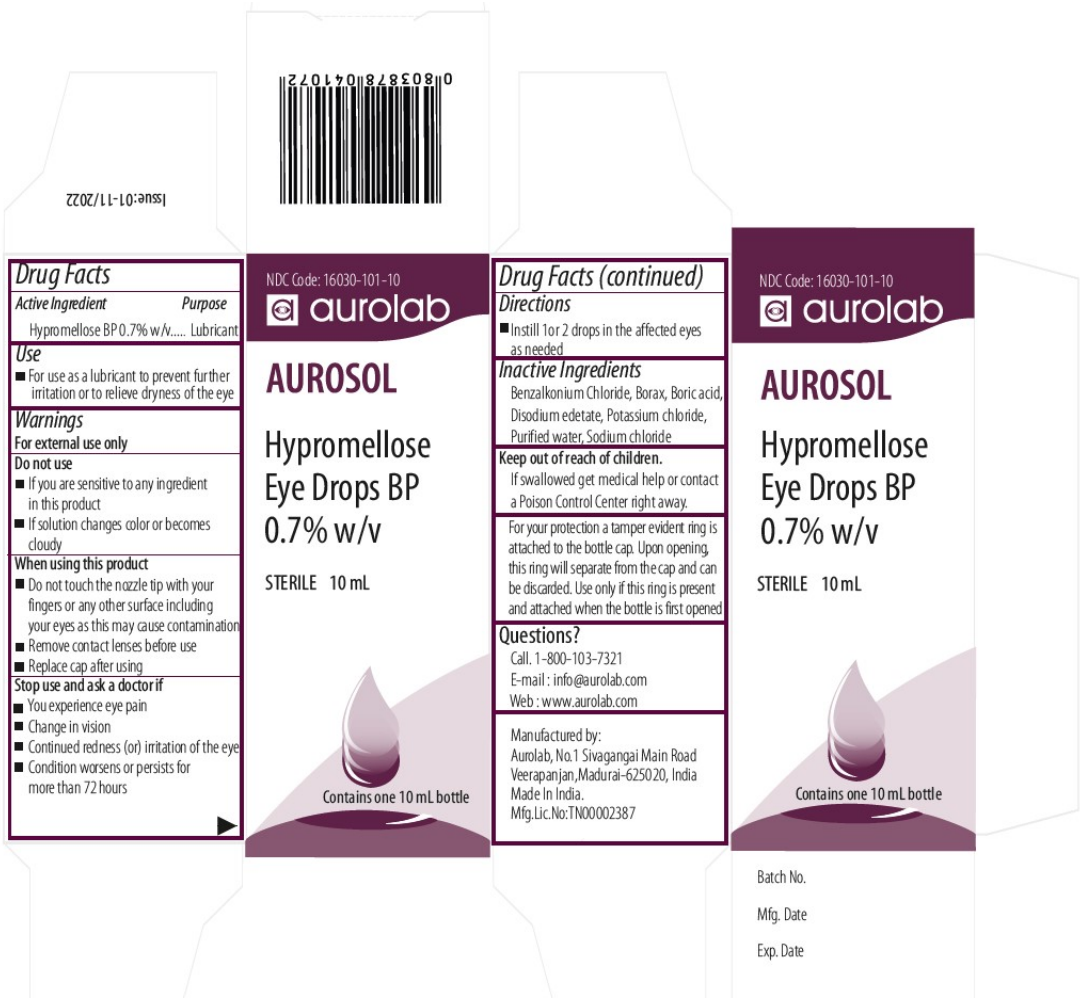
Indications and Usage

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

Eye Lubricant

Eye Lubricant

CARTON LABEL



HYPROMELLOSE EYE DROPS 0.7%

hypromellose eye drops 0.7% for solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:16030-101
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
HYPROMELLOSE 2910 (4000 MPA.S) (UNII: RN3152OP35) (HYPROMELLOSE 2910 (4000 MPA.S) - UNII:RN3152OP35)	HYPROMELLOSE 2910 (4000 MPA.S)	7 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
BORIC ACID (UNII: R57ZHV85D4)	
WATER (UNII: 059QF0KO0R)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	

SODIUM CHLORIDE (UNII: 451W47IQ8X)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:16030-101-10	10 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	09/01/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part349	09/01/2022	

Labeler - Aurolab (677319965)

Establishment

Name	Address	ID/FEI	Business Operations
Aurolab		677319965	manufacture(16030-101)

Revised: 8/2023 Aurolab