POLYVINYL ALCOHOL EYE DROPS 1,4 % W/V- polyvinyl alcohol eye drops 1,4 % w/v for solution Aurolab

Active ingredient

Polyvinyl Alcohol Eye Drops 1.4 % 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 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. If you experience eye pain
- 2. change in vision
- 3. Continued Redness or irritaion of teh eye
- 4. Condition worsens or persists for mroe than 72 hours

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





Issue:01-11/2022

Drug Facts

Active Ingredient

Purpose

Polyvinyl alcohol USP 1.4% w/v.. Lubricant

Use

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

Warnings

For external use only

Do not use

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

When using this product

- Do not touch the nozzle tip with your fingers or any other surface including your eyes as this may cause contamination
- Remove contact lenses before use
- Replace cap after using

Stop use and ask a 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

NDC Code: 16030-201-10



Polyvinyl Alcohol Eye drops 1.4% w/v

AUROLUBE

STERILE 10 mL



DITECUOTIS

Directions

■ Instill 1 or 2 drops in the affected eyes as needed

Drug Facts (continued)

Inactive Ingredients

Benzalkonium Chloride, Disodium Edetate, Povidone, Purified water, Sodium chloride

Questions?

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Keep out of reach of children.

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

Manufactured by: Aurolab, No.1 sivagangai Main Road Veerapanjan, Madurai-625020, India.

Made In India. Mfg.Lic.No:TN00002387 NDC Code: 16030-201-10



Polyvinyl Alcohol Eye drops 1.4% w/v

AUROLUBE

STERILE 10 mL



Contains one 10 mL bottle

Batch No.

Mfg.Date:

Exp. Date:

POLYVINYL ALCOHOL EYE DROPS 1,4 % W/V

polyvinyl alcohol eye drops 1,4 % w/v for solution

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:16030-201

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name

POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990) (POLYVINYL ALCOHOL, UNSPECIFIED - UNII:532B59J990)

POLYVINYL ALCOHOL, UNSPECIFIED - UNII:532B59J990)

POLYVINYL ALCOHOL, UNSPECIFIED - UNII:532B59J990)

Inactive Ingredients

Ingredient Name	Strength		
POVIDONE (UNII: FZ989GH94E)			
EDETATE SODIUM (UNII: MP1J8420LU)			
WATER (UNII: 059QF0KO0R)			
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)			
SODIUM CHLORIDE (UNII: 451W47IQ8X)			

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
		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(16030-201)		

Revised: 12/2024 Aurolab