CLEAR EYES TRIPLE RELIEF- polyvinyl alcohol and povidone and tetrahydrozoline hydrochloride liquid
Prestige Brands Holdings, Inc.

Clear Eyes Triple Relief

Drug Facts

Active ingredients

Polyvinyl alcohol 0.5%

Purpose

Lubricant

Active ingredients

Povidone 0.6%

Purpose

Lubricant

Active ingredient

Tetrahydrozoline hydrochloride 0.05%

Purpose

Redness reliever

Uses

- for the temporary relief of burning & irritation due to dryness of the eye
- for use as a protectant against further irritation or to relieve dryness of the eye
- relieves redness of the eye due to minor eye irritations

Warnings

For external use only.

Do not use if

solution changes color or becomes cloudy.

Ask a doctor before use if you have

narrow angle glaucoma.

When using this product

- to avoid contamination, do not touch tip to any surface
- replace cap after using
- overuse may produce increased redness of the eye
- pupils may become temporarily enlarged

Stop use & ask a doctor if

- you experience eye pain
- you experience changes in vision
- you experience continued redness or irritation of the eye
- the condition worsens or symptoms last for more than 72 hours

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

Instill 1 to 2 drops in the affected eye(s) up to four times daily.

Other information

- store at room temperature
- remove contact lenses before using

Inactive ingredients

benzalkonium chloride, dextrose, dibasic sodium phosphate, edetate disodium, monobasic sodium phosphate, potassium chloride, purified water, sodium bicarbonate, sodium chloride, sodium citrate

Questions?

1-877-274-1787 Cleareyes.com

PRINCIPAL DISPLAY PANEL

CLEAR EYES®

TRIPLE RELIEF
LUBRICANT/REDNESS RELIEVER EYE DROPS

STERILE 0.5 FL OZ (15 mL)

CLEAR EYES TRIPLE RELIEF

polyvinyl alcohol and povidone and tetrahydrozoline hydrochloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67172-898
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990) (POLYVINYL ALCOHOL, UNSPECIFIED - UNII:532B59J990)	POLYVINYL ALCOHOL, UNSPECIFIED	5 mg in 1 mL
POVIDONE (UNII: FZ989GH94E) (POVIDONE - UNII:FZ989GH94E)	POVIDONE	6 mg in 1 mL
TETRAHYDROZOLINE HYDROCHLORIDE (UNII: 0YZT43HS7D) (TETRAHYDROZOLINE - UNII:S9U025Y077)	TETRAHYDROZOLINE HYDROCHLORIDE	0.5 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**

BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
DEXTROSE (UNII: IY9XDZ35W2)	
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
SODIUM PHOSPHATE, MONOBASIC (UNII: 3980JIH2SW)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67172-898-01	1 in 1 BOX	03/15/2011	
1		15 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M018	03/15/2011	

Labeler - Prestige Brands Holdings, Inc. (159655021)

Revised: 10/2024 Prestige Brands Holdings, Inc.