CLEAR EYES COOLING REDNESS RELIEF- glycerin and naphazoline hydrochloride liquid Prestige Brands Holdings, Inc.

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.

Clear Eyes Cooling Redness Relief

Drug Facts

Active ingredients

Glycerin 0.5%

Purpose

Lubricant

Active ingredients

Naphazoline hydrochloride 0.03%

Purpose

Redness Reliever

Uses

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

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 of container 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 persists 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 20°-25°C (68°-77°F)
- remove contact lenses before using

Inactive ingredients

benzalkonium chloride, boric acid, cyclodextrin, edetate disodium, menthol, purified water, sodium borate

Questions?

1-877-274-1787 www.cleareyes.com

PRINCIPAL DISPLAY PANEL

CLEAR EYES®
COOLING
COMFORT
Lubricant/Redness Relief Eye Drops
STERILE 0.5 FL OZ (15 ML)



CLEAR EYES COOLING REDNESS RELIEF

glycerin and naphazoline hydrochloride liquid

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Product Type HUMAN OTC DRUG Item Code (Source) NDC:67172-393

Route of Administration OPHTHALMIC

Active Ingredient/Active Moiety					
Ingredient Name	Basis of Strength	Strength			
NAPHAZOLINE HYDROCHLORIDE (UNII: MZ1131787D) (NAPHAZOLINE - UNII: H231GF11BV)	NAPHAZ OLINE HYDROCHLORIDE	0.3 mg in 1 mL			
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	5 mg in 1 mL			

Inactive Ingredients		
	Ingredient Name	Strength

BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BORIC ACID (UNII: R57ZHV85D4)	
BETADEX (UNII: JV039JZZ3A)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
MENTHOL (UNII: L7T10EIP3A)	
WATER (UNII: 059QF0KO0R)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	

Product Characteristics					
Color	white	Score			
Shape		Size			
Flavor		Imprint Code			
Contains					

Packaging						
# Item Code	Package Description	Marketing Start Date	Marketing End Date			
1 NDC:67172-393-01	1 in 1 BOX	02/15/2011				
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 final	part349	02/15/2011				

Labeler - Prestige Brands Holdings, Inc. (159655021)

Revised: 6/2023 Prestige Brands Holdings, Inc.