CLEAR EYES MAXIMUM REDNESS RELIEF- naphazoline hydrochloride and glycerin 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 Max 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 the 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 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 20°-25°C (68°-77°F)
- remove contact lenses before using

Inactive ingredients

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

Questions?

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

PRINCIPAL DISPLAY PANEL

CLEAR EYES®
MAXIMUM
REDNESS RELIEF
LUBRICANT/REDNESS RELIEVER EYE DROPS
STERILE 1 FL OZ (30 mL)

CLEAR EYES MAXIMUM REDNESS RELIEF

naphazoline hydrochloride and glycerin liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67172-696
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
NAPHAZOLINE HYDROCHLORIDE (UNII: MZ1131787D) (NAPHAZOLINE - UNII:H231GF11BV)	NAPHAZOLINE 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)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
WATER (UNII: 059QF0KO0R)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67172-696-01	1 in 1 CARTON	09/01/2010	
1		30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
2	NDC:67172-696-02	1 in 1 CARTON	09/01/2010	
2		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	09/01/2010	

Labeler - Prestige Brands Holdings, Inc. (159655021)

Revised: 6/2023 Prestige Brands Holdings, Inc.