

CLEAR EYES MAXIMUM REDNESS RELIEF- naphazoline hydrochloride and glycerin liquid

Prestige Brands Holdings, Inc.

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)

Drug Facts

KEEP CARTON FOR FUTURE REFERENCE

Active ingredients	Purposes
Glycerin 0.5%.....Lubricant	Naphazoline hydrochloride 0.02%.....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 to relieve 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 to any surface
- replace cap after using
- overuse may produce increased redness of the eye
- pupils may become temporarily enlarged.

Stop use and 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)

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Questions? 1-877-234-1787 Cleareyes.com

BOTTLE IMAGE IS ACTUAL SIZE

UP TO 12 HOURS OF SOOTHING COMFORT



— MAXIMUM —
REDNESS RELIEF

To open cap, push down and turn.



6 78112 25510 8

UP TO 12 HOURS OF SOOTHING COMFORT

MAXIMUM strength

TRANSFORM YOUR DAY™



CLEAR EYES

— MAXIMUM —
REDNESS RELIEF

LUBRICANT/REDNESS RELIEVER EYE DROPS

relieves redness, dryness, burning & irritation

STERILE 1 fl oz (30mL)

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 Drug	M018	09/01/2010	

Labeler - Prestige Brands Holdings, Inc. (159655021)

Revised: 2/2025

Prestige Brands Holdings, Inc.