CLEAR EYES REDNESS RELIEF HANDY POCKET PAL- naphazoline hydrochloride and glycerin liquid

CLEAR EYES 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 Redness Relief

Clear Eyes Redness Relief

Drug Facts

Active Ingredients

Glycerin 0.25%

Purpose

Lubricant

Active Ingredients

Naphazoline hydrochloride 0.012%

Purpose

Redness Reliever

Uses

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

- pupils may become enlarged temporarily
- overuse may produce increased redness of the eye
- remove contact lenses before using
- to avoid contamination, do not touch tip of container to any surface

replace cap after using

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
- 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 4 times daily.

Other Information

- store at room temperature

Inactive Ingredients

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

Questions?

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

Clear Eyes Redness Relief, Handy Pocket Pal

Drug Facts

Active Ingredients

Glycerin 0.25%

Purpose

Lubricant

Active Ingredients

Naphazoline hydrochloride 0.012%

Purpose

Redness Reliever

Uses

- Relieves redness of the eye due to minor eye irritations.
- For use as a protectant against further irritation or to relieve dryness of the eye.

• Temporarily relieves burning and irritation due to 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 cause increased redness of the eye.
- Pupils may become enlarged temporarily.

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
- 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 Between 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 REDNESS RELIEF LUBRICANT/REDNESS RELIEVER EYE DROPS Sterile 0.5 FL OZ (15 mL)



PRINCIPAL DISPLAY PANEL

UP TO 12 HOURS OF SOOTHING COMFORT Clear eyes REDNESS RELIEF LUBRICANT/REDNESS RELIEVER EYE DROPS Handy Pocket Pal[®] Sterile 0.2 FL OZ (6 mL)





CLEAR EYES REDNESS RELIEF HANDY POCKET PAL

naphazoline hydrochloride and glycerin liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:67172-796
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
NAPHAZOLINE HYDRO CHLO RIDE (UNII: MZ1131787D) (NAPHAZOLINE - UNII:H231GF11BV)	NAPHAZOLINE HYDROCHLORIDE	.00012 mg in 1 mL		
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	.0025 mg in 1 mL		

Inactive Ingredients		
	Ingredient Name	Strength

BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7)	
BORIC ACID (UNII: R57ZHV85D4)	
EDETATE DISO DIUM (UNII: 7FLD91C86K)	
WATER (UNII: 059QF0KO0R)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	

Product Characteristics					
Color WHITE Score					
Shape		Size			
Flavor		Imprint Code			
Contains					

1	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:67172-796-01	1 in 1 BOX	11/0 1/20 12		
1		6 mL in 1 BOTTLE; 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	11/0 1/20 12		

CLEAR EYES REDNESS RELIEF

naphazoline hydrochloride and glycerin liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:67172-797
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
NAPHAZOLINE HYDRO CHLO RIDE (UNII: MZ1131787D) (NAPHAZOLINE - UNII:H231GF11BV)	NAPHAZOLINE HYDROCHLORIDE	0.12 mg in 1 mL
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	2.5 mg in 1 mL

Inactive Ingredients				
Ingredient Name	Strength			
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)				
BORIC ACID (UNII: R57ZHV85D4)				
EDETATE DISO DIUM (UNII: 7FLD9 1C8 6 K)				
WATER (UNII: 059QF0KO0R)				

SODIUM BORATE (UNII: 91MBZ8H3QO)

Product Characteristics					
Color WHITE Score					
Shape		Size			
Flavor		Imprint Code			
Contains					

P	ackaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67172-797- 01	1 in 1 BOX	04/27/2011	
1		30 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:67172-797- 15	1 in 1 CARTON	04/27/2011	
2		15 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
3	NDC:67172-797- 06	1 in 1 CARTON	04/27/2011	
3		6 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	04/27/2011	

Labeler - Prestige Brands Holdings, Inc. (159655021)

Revised: 2/2019 Prestige Brands Holdings, Inc.