CLEAR EYES MAXIMUM ITCHY EYE RELIEF- glycerin and naphazoline hydrochloride and zinc sulfate 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 Maximum Itchy Eye Relief

Drug Facts

Active Ingredient

Glycerin 0.25%

Purpose

Lubricant

Active Ingredient

Naphazoline hydrochloride 0.012%

Purpose

Redness Reliever

Active Ingredient

Zinc Sulfate 0.25%

Purpose

Astringent

Uses

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

- 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 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 right away.

Directions

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

Other Information

Store at room temperature

Inactive Ingredients

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

Questions?

1-877-274-1787 Cleareyes.com

PRINCIPAL DISPLAY PANEL

Clear eyes®
MAXIMUM ITCHY EYE RELIEF
ASTRINGENT/LUBRICANT/REDNESS RELIEVER EYE DROPS

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CLEAR EYES MAXIMUM ITCHY EYE RELIEF

glycerin and naphazoline hydrochloride and zinc sulfate liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67172-999
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	2.5 mg in 1 mL
NAPHAZOLINE HYDROCHLORIDE (UNII: MZ1131787D) (NAPHAZOLINE - UNII:H231GF11BV)	NAPHAZOLINE HYDROCHLORIDE	.12 mg in 1 mL
ZINC SULFATE (UNII: 89DS0H96TB) (ZINC CATION - UNII:13S1S8SF37)	ZINC SULFATE	2.5 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 CHLORIDE (UNII: 451W47IQ8X)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	

Product Characteristics

Color	WHITE	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67172-999-01	1 in 1 BOX	03/15/2011	
1		15 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
2	NDC:67172-999-06	1 in 1 CARTON	09/04/2018	
2		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	03/15/2011	

Labeler - Prestige Brands Holdings, Inc. (159655021)

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