CLEAR EYES MAXIMUM ITCHY EYE RELIEF- glycerin and naphazoline hydrochloride and zinc sulfate liquid
Prestige Brands Holdings, Inc.

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

- to avoid contamination, do not touch tip to any surface
- replace cap after using
- overuse may produce increased redness of the eye
- overuse may produce increased redness of the eye

Stop useandask 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 20-25C (68-77F)
- remove contact lenses before using

Inactive Ingredients

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

Questions?

1-877-274-1787Cleareyes.com

PRINCIPAL DISPLAY PANEL

CLEAR
EYES ®
MAXIMUM
ITCHY EYE

RELIEF
ASTRINGENT LUBRICANT/
REDNESS RELIEVER EYE DROPS

PRINCIPAL DISPLAY PANEL

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

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	0.12 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	05/31/2020
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 Drug	M018	03/15/2011	

Labeler - Prestige Brands Holdings, Inc. (159655021)

Revised: 9/2026 Prestige Brands Holdings, Inc.