CLEAR EYES NATURAL TEARS- polyvinyl alcohol and povidone 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 Natural Tears

Drug Facts

Active ingredient

Polyvinyl alcohol 0.5%

Purpose

Lubricant

Active ingredient

Povidone 0.6%

Purpose

Lubricant

Uses

- 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

When using this product

- 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 lasts 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) as needed.

Other information

- store at room temperature
- remove contact lenses before using

Inactive ingredients

benzalkonium chloride, dextrose, edetate disodium, potassium chloride, purified water, sodium bicarbonate, sodium chloride, sodium citrate, sodium phosphate (mono- and dibasic)

Questions?

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

PRINCIPAL DISPLAY PANEL

Clear eyes® Natural Tears
Lubricating Eye Drops
STERILE 0.5 FL OZ (15 mL)

CLEAR EYES NATURAL TEARS

polyvinyl alcohol and povidone liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67172-557
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990) (POLYVINYL ALCOHOL, UNSPECIFIED - UNII:532B59J990)	POLYVINYL ALCOHOL, UNSPECIFIED	5.0 mg in 1 mL
POVIDONE (UNII: FZ989GH94E) (POVIDONE - UNII:FZ989GH94E)	POVIDONE	6.0 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
DEXTROSE (UNII: IY9XDZ35W2)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SODIUM PHOSPHATE (UNII: SE337SVY37)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67172-557-02	1 in 1 BOX	05/15/2013	
1		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	05/15/2013	

Labeler - Prestige Brands Holdings, Inc. (159655021)

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