ARTIFICIAL TEARS LUBRICANT EYE- polyvinyl alcohol and povidone liquid
Prestige Brands Holdings, Inc.

Artificial Tears Lubricant Eye Drops

Drug Facts

Active ingredients

Polyvinyl alcohol 0.5%

Povidone 0.6%

Purpose

Lubricant

Uses

- For the temporary relief of burning & irritations 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 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) 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

PRINCIPAL DISPLAY PANEL

Sterile
Artificial Tears
Lubricant Eye Drops
0.5 FL OZ (15 mL)

ARTIFICIAL TEARS LUBRICANT EYE

polyvinyl alcohol and povidone liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67172-181
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, DIBASIC (UNII: GR686LBA74)	
SODIUM PHOSPHATE, MONOBASIC (UNII: 3980JIH2SW)	

Product Characteristics

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67172-181-01	1 in 1 BOX	06/01/2012	
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 Drug	M018	06/01/2012	

Labeler - Prestige Brands Holdings, Inc. (159655021)

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