MURINE TEARS FOR DRY EYES- 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.

Murine Tears for Dry Eyes

Drug Facts

Active ingredient

Polyvinyl alcohol 0.5%

Purpose

Lubricant

Active ingredient

Povidone 0.6%

Purpose

Lubricant

Uses

- For the temporary relief of burning & irritation due to dryness of the eye.
- For use as a protectant against further irritation or 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.
- recap 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 right away.

Directions

Instill 1 to 2 drops in the affected eye(s) as needed.

Other information

- store at 20°- 25°C (68°-77°F)
- remove contact lenses before using
- **Tamper Evident:** Do not use if neckband on bottle is broken or missing.

Inactive ingredients

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

Questions?

1-877-854-0853 www.murine.com

PRINCIPAL DISPLAY PANEL

Murine
Tears® For DRY EYES
Lubricant Eye Drops
Original Natural Tears Formula
● Moisture Enriched
● Eye Doctor Recommended
0.5 fl oz. (15 mL)

MURINE TEARS FOR DRY EYES

polyvinyl alcohol and povidone liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67172-574
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 mg in 1 mL
POVIDONE (UNII: FZ989GH94E) (POVIDONE - UNII:FZ989GH94E)	POVIDONE	6 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:67172-574-01	1 in 1 BOX	10/01/2012	
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 FINAL	part349	10/01/2012	

Labeler - Prestige Brands Holdings, Inc. (159655021)

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