PREMIER VALUE EYE DROPS MT - polyvinyl alcohol and povidone solution
HANLIM PHARM. CO., LTD.

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.

Drug Facts

Active ingredients	Purpose
Polyvinyl Alcohol 0.5%	Lubricant
Povidone 0.6%	Lubricant

Uses
- relieves dryness of the eye(s)
- temporary relieves burning and irritations due to dryness of the eye
- protects against further irritation

For external use only.

Do not use: If solution changes color or becomes cloudy or if you are sensitive to any ingredient in this product.

When using this product
- remove contact lenses before using
- do not touch tip of container to any surface to avoid contamination
- replace cap after each use

Stop use and ask a doctor if
- you feel eye pain
- changes in vision occur
- redness of irritation of the eye gets worse or lasts more than 72 hours

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of the reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions
- Instill 1 or 2 drops in the affected eye(s) as needed.
- Store at 15º-25ºC (59º-77ºF).
- Children under 6 years of age: Ask a doctor

Inactive Ingredients: Benzalkonium Chloride, Dextrose, Disodium Edetate, Potassium Chloride, Purified Water, Sodium Bicarbonate, Sodium Chloride, Sodium Citrate,

Sodium Phosphate (Mono- and Dibasic)

Distributed By:

Chain Drug Consortium, LLC.

3301 NW Boca Raton Blvd. Suite 101

Boca Raton, FL33431

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PREMIER VALUE EYE DROPS MT

polyvinyl alcohol and povidone solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11716-0224
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

POLYVINYL ALCOHOL (UNII: 532B59J990) (POLYVINYL ALCOHOL - UNII:532B59J990)	POLYVINYL ALCOHOL	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:11716-0224-2	1 in 1 CARTON		
1		15 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part349	10/19/2010	

Labeler - HANLIM PHARM. CO., LTD. (687986034)

Registrant - UNITED EXCHANGE CORP. (840130579)

Establishment

Name	Address	ID/FEI	Business Operations
HANLIM PHARM. CO., LTD.		687986034	manufacture

Revised: 10/2010 HANLIM PHARM. CO., LTD.