GOODSENSE LUBRICATING EYE DROPS - carboxymethylcellulose sodium 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

Carboxymethylcellulose Sodium 0.5%----------Lubricant

Glycerin 0.9%----------Lubricant

Uses

- For the temporary relief of burning, irritation, and discomfort due to dryness of the eye or from irritation from wind or sun.
- May be sued to protect against further irritation.

Warnings

- For external use only.

When using this product

- To avoid contamination, do not touch tip of container to any surface. Replace cap after using.
- Do not use if solution changes color or gets cloudy.

Stop use and ask a doctor if

You feel eye pain, changes in vision, continued redness, or irritation of the eye, or if the condition worsens or persists for more than 72 hours.

Keep out of reach of children.

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

Directions

Put 1 or 2 drops in the affected eye/s as needed.

Inactive ingredients: Boric Acid, Calcium Chloride Dihydrate, Chlorhexidine Gluconate, Erythritol Hexahydrate, Levocarnitine, Magnesium Chloride, Potassium Chloride, Purified Water, Sodium Borate Decahydrate, Sodium Citrate Dihydrate

Process Yellow | PMS 871 | PMS 274 | PMS Black

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GOODSENSE LUBRICATING EYE DROPS
carboxymethylcellulose sodium solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11716-0041
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311) (CARBOXYMETHYLCELLULOSE - UNII:05JZI7B19X)	CARBOXYMETHYLCELLULOSE SODIUM	5 mg in 1 mL
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	9 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
BORIC ACID (UNII: R57ZHV85D4)	
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	
CHLORHEXIDINE GLUCONATE (UNII: MOR84MUD8E)	
ERYTHRITOL (UNII: RA96B954X6)	
LEVOCARNITINE (UNII: 0G389FZZ9M)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:11716-0041-8	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	07/30/2010	

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

Revised: 7/2010 HANLIM PHARM. CO., LTD.