LEADER LUBRICATING EYE- carboxymethylcellulose sodium, and glycerin solution/ drops
CARDINAL HEALTH

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.

Active ingredients Purpose

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

Glycerin 0.9%---------- Lubricnat

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 used 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

benzalkonium chloride, boric acid, calcium chloride dihydrate, erythritol, levocarnitine, magnesium chloride, potassium chloride, purified water, sodium borate, sodium citrate dihydrate

DISTRIBUTED BY CARDINAL HEALTH

DUBLIN, OHIO 43017

CIN 4290763

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LEADER LUBRICATING EYE

carboxymethylcellulose sodium, and glycerin solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:37205-635
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
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BORIC ACID (UNII: R57ZHV85D4)	
CALCIUM CHLORATE DIHYDRATE (UNII: BBB2RG413E)	
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:37205-635-05	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	01/04/2012	

Labeler - CARDINAL HEALTH (097537435)

Revised: 12/2014 CARDINAL HEALTH