QUALITY CHOICE LUBRICATING EYE DROPS - carboxymethylcellulose sodium, glycerin solution
CHAIN DRUG MARKETING ASSOCIATION 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.

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.

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.

Other information

- some users may experience a brief tingling sensation
- Store at room temperature.

Inactive ingredients:

Boric Acid, Sodium Borate, Levocarnitin, Potassium Chloride, Calcium Chloride Dihydrate, Magnesium Chloride, Erythritol, Sodium Citrate Dihydrate, Potassium Sorbate, Purified Water

Distributed by C.D.M.A., Inc.

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Made in Korea

QUALITY CHOICE LUBRICATING EYE DROPS

carboxymethylcellulose sodium, glycerin solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63868-969
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)	

SODIUM BORATE (UNII: 91MBZ8H3QO)	
LEVOCARNITINE (UNII: 0G389FZZ9M)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
ERYTHRITOL (UNII: RA96B954X6)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-969-15	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	09/30/2012	

Labeler - CHAIN DRUG MARKETING ASSOCIATION INC (011920774)

Revised: 9/2012 CHAIN DRUG MARKETING ASSOCIATION INC