LUBRICANT EYE PLUS- carboxymethylcellulose sodium solution/ drops
Rite Aid Corporation

Rite Aid Corporation Lubricant Eye Drops Plus Drug Facts

Active ingredient (in each single-use container)

Carboxymethylcellulose sodium 0.5%

Purpose

Eye lubricant

Uses

- for the temporary relief of burning, irritation, and discomfort due to dryness of the eye or exposure to wind or sun
- may be used as a protectant against further irritation

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
- do not reuse
- once opened, discard
- do not touch unit-dose tip to eye

Stop use and ask a doctor if

- you experience eye pain
- changes in vision occur
- redness or irritation of the eye continues
- redness or irritation of the eye 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 right away (1-800-222-1222).

Directions

- to open, twist and pull tab to remove
- instill 1 or 2 drops in the affected eye(s) as needed and discard container

Other information

- store at 20-25°C (68-77°F)
- RETAIN THIS CARTON FOR FUTURE REFERENCE.

Inactive ingredients

calcium chloride dihydrate, magnesium chloride hexahydrate, potassium chloride, sodium chloride, sodium lactate solution, water for injection. May also contain hydrochloric acid and/or sodium hydroxide to adjust pH.

Questions or comments?

1-800-719-9260

Package/Label Principal Display Panel

FREE FROM | PRESERVATIVE FREE

Compare to the active ingredient of Refresh Plus® Lubricant Eye Drops

MOISTURIZING RELIEF

ACTUAL SIZE

LUBRICANT EYE DROPS PLUS

CARBOXYMETHYLCELLULOSE SODIUM 0.5%

Instantly moisturizes & relieves dry, irritated eyes

For sensitive eyes

30 SINGLE-USE CONTAINERS

0.01 FL OZ (0.4 mL) each Sterile

LUBRICANT EYE PLUS

carboxymethylcellulose sodium solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11822-0323
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM (UNII: K679OBS311) (CARBOXYMETHYLCELLULOSE - UNII:05JZI7B19X)	CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM	0.5 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	

MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM LACTATE (UNII: TU7HW0W0QT)	
WATER (UNII: 059QF0KO0R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822-0323-1	6 in 1 CARTON	09/16/2013	
1		5 in 1 POUCH		
1		0.4 mL in 1 VIAL, SINGLE-USE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M018	09/16/2013	

Labeler - Rite Aid Corporation (014578892)

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