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

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.

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

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

LUBRICANT EYE DROPS PLUS

LUBRICANT EYE DROPS

carboxymethylcellulose sodium 0.5%

ACTUAL SIZE

preservative-free

immediate, soothing relief for dry eyes

30 STERILE Single-Use Containers

0.01 FL OZ (0.4 mL) EACH



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 (UNII: K679OBS311) (CARBOXYMETHYLCELLULOSE - UNII:05JZI7B19X)	CARBOXYMETHYLCELLULOSE SODIUM	0.5 g in 100 mL			

Inactive Ingredients		
Ingredient Name	Strength	
CALCIUM CHLORIDE (UNII: M4I0 D6 VV5M)		

MAGNESIUM CHLO RIDE (UNII: 02F3473H9O)	
POTASSIUM CHLORIDE (UNII: 660 YQ98 I10)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM LACTATE (UNII: TU7HW0W0QT)	
WATER (UNII: 059QF0KO0R)	
SO DIUM HYDRO XIDE (UNII: 55X0 4QC32I)	
HYDRO CHLO RIC ACID (UNII: QTT17582CB)	

P	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 final	part349	09/16/2013		

Labeler - Rite Aid Corporation (014578892)

Revised: 7/2019 Rite Aid Corporation