LUBRICANT EYE DROPS PRESERVATIVE FREE 30 SINGLE USE CONTAINER-carboxymethylcellulose sodium liquid Nexus Optical 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.

Lubricant Eye Drops Preservative-free, 30 Single Use Container

Drug Facts

Active ingredients

Carboxymethylcellulose sodium 0.5%

Purposes

Lubricant

Use

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

Warnings

For external use only

Do not use

solution changes color or becomes cloudy

When using this product

- do not reuse
- once opened, discard
- to avoid contamination, do not touch tip of container to any surface
- 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 accidentally swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) immediately.

Directions

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

• if used for post-operative (e.g., LASIK) dryness and discomfort, follow your eye doctor's instructions

Other information

- store at 15°-25°C (59°-77°F)
- use only if single-use container is intact
- use before expiration date marked on container
- RETAIN THIS CARTON FOR FUTURE REFERENCE

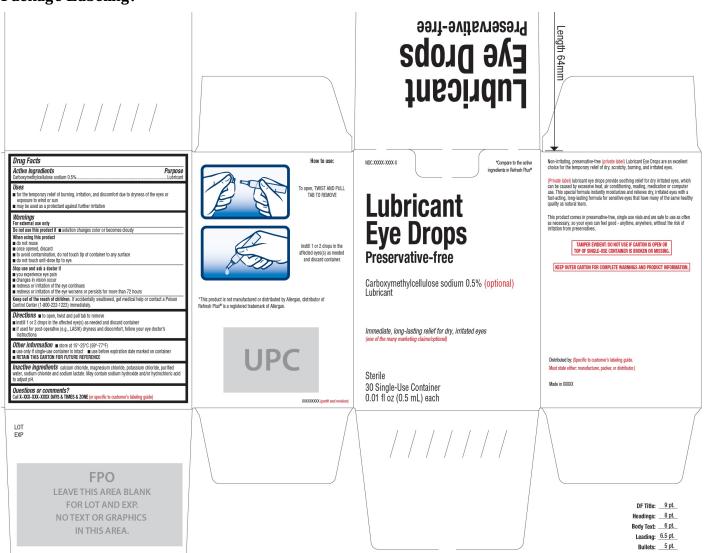
Inactive ingredients

calcium chloride, magnesium chloride, potassium chloride, purified water, sodium chloride and sodium lactate. May contain sodium hydroxide and/or hydrochloric acid to adjust pH.

Questions or comments?

Call X-XXX-XXX-XXXX DAYS & TIMES & ZONE

Package Labeling:



carboxymethylcellulose sodium liquid

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73450-002		
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		

Inactive Ingredients		
Ingredient Name	Strength	
CALCIUM CHLO RIDE (UNII: M4I0 D6 VV5M)		
MAGNESIUM CHLO RIDE (UNII: 02F3473H9O)		
POTASSIUM CHLORIDE (UNII: 660 YQ 98 I10)		
WATER (UNII: 059QF0KO0R)		
SODIUM CHLORIDE (UNII: 451W47IQ8X)		
SODIUM LACTATE (UNII: TU7HW0W0QT)		

l	Packaging				
	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
	1 NDC:73450-002- 05	30 in 1 BOX	04/20/2020		
	1	0.5 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	04/20/2020		

Labeler - Nexus Optical Inc (107289657)

Revised: 2/2020 Nexus Optical Inc