LUBRICANT EYE DROPS- carboxymethylcellulose sodium solution/ drops
Oliver Landon Intl 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

Drug Facts

Active ingredient

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.
- protects against further irritation.

Warnings

For external use only

Do not use if solution changes color or becomes cloudy

When using this product

- do not touch tip of container to any surface to avoid contamination.
- do not reuse. Once opened, discard.

Stop use and ask a doctor if

- you experience eye pain, changes in vision, continued redness or irritation of the eye
- 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 right away.

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 dryness and discomfort, follow your eye doctor's instructions.

Other information

- use before expiration date marked on container.
- store at 59°-86°F (15°-33°C)

RETAIN THIS CARD FOR FUTURE REFERENCE.

Use only if single-use container is intact.

Inactive ingredients Calcium chloride, magnesium chloride, potassium chloride, purified water, sodium chloride and sodium lactate.

May also contain hydrochloric acid and/or sodium hydroxide to adjust pH.

Questions & Comments? **1 (800) 839-5929**

Manufactured for: Oliver Landon Intl. Inc.

The Grove, 21 Pine Road

Belleville, St. Michael, BARBADOS BB11113

Lubricant Eye Drops
0.4 ml (0.014 fl oz) per vial

FRONT

BACK

LUBRICANT EYE DROPS

carboxymethylcellulose sodium solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:59276-905
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 CHLORIDE (UNII: M4I0D6VV5M)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	

SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM LACTATE (UNII: TU7HW0W0QT)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59276-905-05	5 in 1 CELLO PACK	02/25/2019	
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	02/25/2019	

Labeler - Oliver Landon Intl Inc. (815240195)

Revised: 11/2020 Oliver Landon Intl Inc.