# 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.

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### **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.

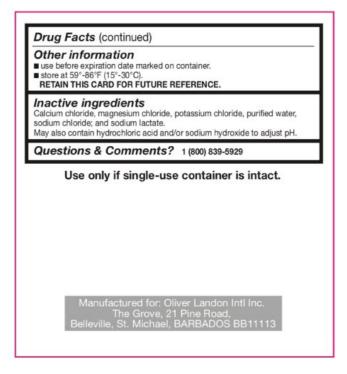
The Grove, 21 Pine Road

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	
CARBO XYMETHYLCELLULOSE SODIUM (UNII: K679 OBS 311) (CARBO XYMETHYLCELLULOSE - UNII: 05 JZI7B 19 X)	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 YQ98 I10)		
WATER (UNII: 059QF0KO0R)		

SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM LACTATE (UNII: TU7HW0W0QT)	
HYDRO CHLORIC ACID (UNII: QTT17582CB)	
SO DIUM HYDRO XIDE (UNII: 55X04QC32I)	

F	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.