

REAL- 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.

Real Lubricant Eye Drops

Drug Facts

Active ingredient

Carboxymethylcellulose sodium 0.5%

Purpose

Eye Lubricant

Use

- For 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
- 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
- If solution changes color or becomes cloudy, do not use

Stop use and ask a doctor if you experience eye pain, changes in vision, continued redness or irritation of the eye, or if the condition worsens or persists for more than 72 hours.

If pregnant or breast-feeding, ask a doctor before use.

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 only if single-use container is intact
- Use before expiration date marked on container
- Store at 59°-86°F (15°-33°C)
- RETAIN THIS CARTON FOR FUTURE REFERENCE

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 or Comments?

info@oliverlandon.biz

NDC 59276-403-05

Real Lubricant Eye Drops

- Moisturizing relief
- Immediate, soothing relief for dry eyes.

Preservative-free

Directions

To open, TWIST AND PULL TAB TO REMOVE

Instill 1 or 2 drops in the affected eye(s) as needed and discard container.

5 Single-Use Containers

0.01 fl oz (0.4 mL) each Sterile



REAL

carboxymethylcellulose sodium solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59276-403
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:59276-403-05	5 in 1 CARTON	11/14/2018	
1		0.4 mL in 1 VIAL, SINGLE-USE; Type 0: Not a Combination Product		
2	NDC:59276-403-30	30 in 1 CARTON	11/14/2018	
2		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	11/14/2018	

Labeler - Oliver Landon Intl Inc. (815240195)