FAMILY CARE LUBRICANT SINGLE USE EYE- carboxymethylcellulose sodium solution/ drops
UNITED EXCHANGE CORP

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.

Active ingredient Purpose

Carboxymethylcellulose Sodium 0.5%.....Eye Lubricant

Uses

- For the temporary relief of burning 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 of eye
- if solution changes color or becomes cloudy, do not use

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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

Keep out of the 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 (e.g., LASIK) 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° to 86°F (15° to 30°C)
- RETAIN THIS CARTON FOR FUTURE REFERENCE.

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.

Distributed by:

UNITED EXCHANGE CORP.

17211 Valley View Ave.

Cerritos, CA 90703 USA

MADE IN KOREA

FAMILY CARE LUBRICANT SINGLE USE EYE

carboxymethylcellulose sodium solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:65923-555
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:65923-555-02	5 in 1 CARTON	03/28/2013	
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	02/28/2013	

Labeler - UNITED EXCHANGE CORP (840130579)

Revised: 11/2021 UNITED EXCHANGE CORP