FAMILY CARE 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, 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.

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 (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.

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.

Distributed by:

United Exchange Corp.

17211 Valley View Ave.

Cerritos, CA 90703 U.S.A.

www.ueccorp.com

Toll Free: 1800 814 8028

Made in Korea





Drug Facts Active ingredient

Warnings

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 bouch unit-dose tip to eye.

**If solution changes color or becomes cloudy, do not use.

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

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.
- 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.

How to Use:

To open. TWIST AND PULL TAB TO REMOVE. Instill 1 or 2 drops in the affected eye(s) as needed and discard





Compare to the active ingredient in REFRESH PLUS® Lubricant Eye Drops*

FAMILY FAMILY **CARE**

SINGLE-USE EYE DROPS

- ures:

 For Temporary Relief of
 Burning & Irritation Due to
 Dryness of the Eye
 Small Enough for Your
 Purse or Pocket
 - **SENSITIVE**

CARBOXYMETHYLCELLULOSE SODIUM 0.5% (Eye Lubricant)

> 30 Single-Use Containers 0.01 fl oz (0.4 ml) each Sterile



instantly moisturize and relieve dry irritated eyes. This fast-acting, long-lasting formula has many of the same healthy

qualities as your own natural tears making it perfect for sensitive eyes. Improves dry eyes when used after LASIK surgery.

Family Care™ Single-Use Eye Drops come in preservative-free single-use vials and are safe to use as often as needed, so your eyes can feel good anytime, anywhere

FAMILY CARE SINGLE-USE EYE

carboxymethylcellulose sodium solution/ drops

Product Information

HUMAN OTC DRUG Product Type Item Code (Source) NDC:65923-583

Route of Administration OPHTHALMIC

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CARBO XYMETHYLCELLULO SE SODIUM (UNII: K679 OBS 311) (CARBO XYMETHYLCELLULO SE - UNII: 0 5 J Z I 7 B 19 X)	CARBOXYMETHYLCELLULOSE SODIUM	5 mg in 1 mL

Inactive Ingredients	
Ingredient Name	Strength

CALCIUM CHLORIDE (UNII: M4I0 D6 VV5M)	
MAGNESIUM CHLO RIDE (UNII: 02F3473H9O)	
POTASSIUM CHLORIDE (UNII: 660 YQ98110)	
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM LACTATE (UNII: TU7HW0W0QT)	

F	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:65923-583- 30	30 in 1 BOX	03/28/2016		
1		.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	03/28/2016			

Labeler - United Exchange Corp. (840130579)

Revised: 8/2017 United Exchange Corp.