**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.*

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**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.

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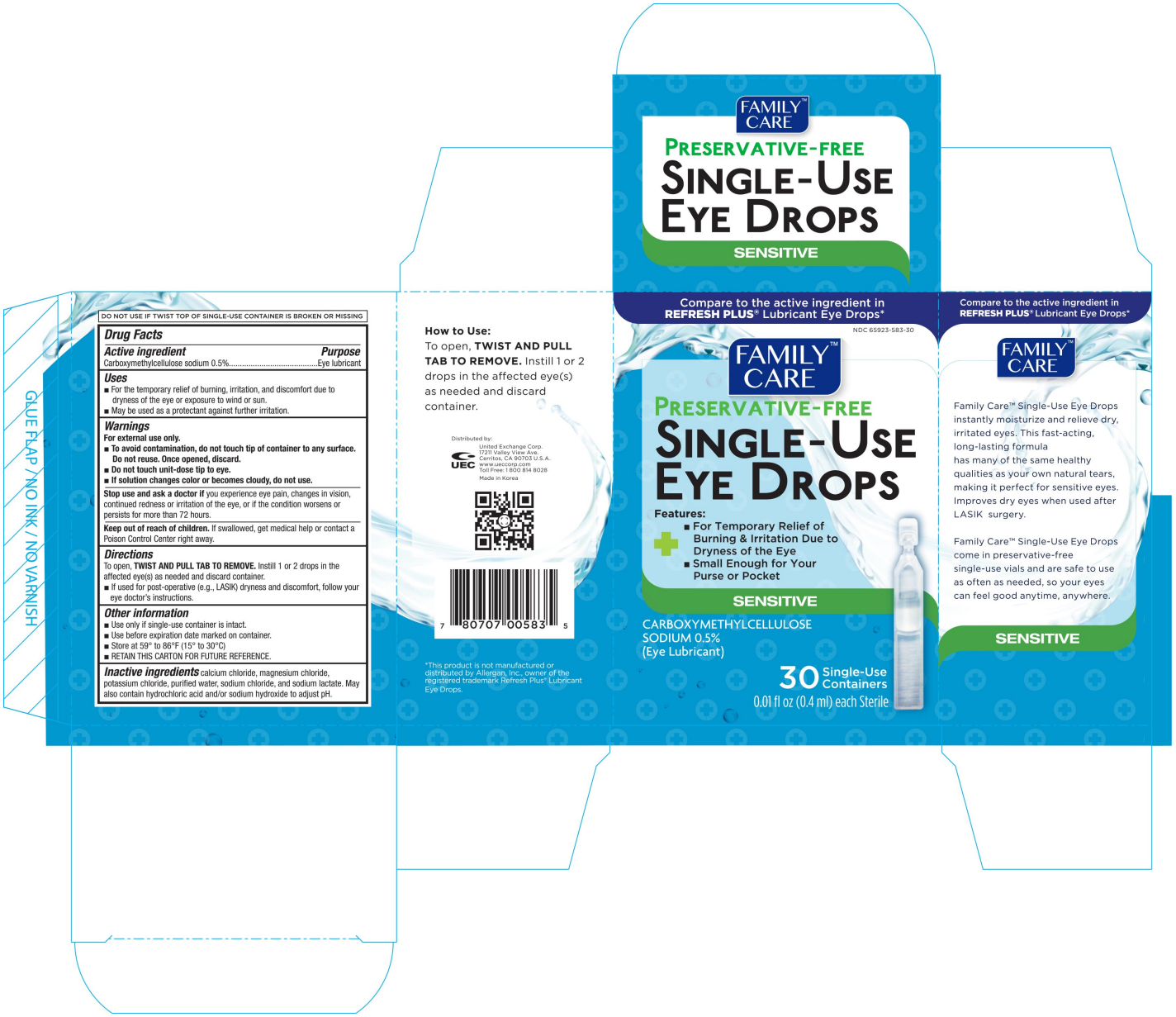
Cerritos, CA 90703 U.S.A.

www.ueccorp.com

Toll Free: 1 800 814 8028

Made in Korea




# FAMILY CARE SINGLE-USE EYE

carboxymethylcellulose sodium solution/ drops

## Product Information

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

## 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.