

**FAMILY CARE SINGLE USE LUBRICANT EYE DROP 0.4ML 30CT- carboxymethyl cellulose 0.5% solution/ drops**  
**United Exchange Corp**

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**Family Care 685 Single Use Lubricant Eye Drop 0.4mL 30ct**

Active ingredient Purpose

Carboxymethylcellulose sodium (0.5%).....Eye lubricant

Uses

- For the temporary relief of burning and irritation due to dryness of the eye.
- For the temporary relief of discomfort due to minor irritations of the eye or from exposure to wind or sun.
- For use as a lubricant to prevent further irritation or to relieve dryness of the eye.

Warnings

For use in the eyes 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.
- 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

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

Other information

- Use only if tamper seal on top and bottom flaps are intact.
- Store between 15-30°C (59-86°F).  Keep carton for complete product information.

Inactive ingredients

Calcium chloride, magnesium chloride, potassium chloride, purified water, sodium chloride, sodium lactate. May also contain hydrochloric acid and/or sodium hydroxide to adjust pH.

Questions or comments?

Call 1-888-645-8204 Monday-Friday 9AM-5PM (PST)

Distributed by: UNITED EXCHANGE CORP.

Cypress, CA 90630 USA | 1-888-645-8204

Made in U.S.A.



## FAMILY CARE SINGLE USE LUBRICANT EYE DROP 0.4ML 30CT

carboxymethyl cellulose 0.5% solution/ drops

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:65923-576
Route of Administration	OPHTHALMIC		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM</b> (UNII: K679OBS311) (CARBOXYMETHYLCELLULOSE - UNII:05JZ17B19X)	CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM	5 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KOOR)	
<b>SODIUM LACTATE</b> (UNII: TU7HW0W0QT)	
<b>CALCIUM CHLORIDE</b> (UNII: M4I0D6VV5M)	
<b>MAGNESIUM CHLORIDE</b> (UNII: 02F3473H9O)	
<b>HYDROCHLORIC ACID</b> (UNII: QTT17582CB)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	

**POTASSIUM CHLORIDE** (UNII: 660YQ98I10)

**SODIUM HYDROXIDE** (UNII: 55X04QC32I)

### Product Characteristics

Color	Score
Shape	Size
Flavor	Imprint Code
Contains	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65923-576-30	30 in 1 BOX	01/16/2026	
1		5 mL in 1 VIAL, SINGLE-DOSE; Type 0: Not a Combination Product		



## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M018	01/16/2026	

**Labeler** - United Exchange Corp (840130579)

Revised: 1/2026

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