**REFRESH CELLUVISC- carboxymethylcellulose sodium gel**
**Allergan, Inc.**

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**REFRESH® CELLUVISC®**
***Drug Facts***

***Active ingredient***

Carboxymethylcellulose sodium 1%

***Purpose***

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.

***Other information***

- Use only if single-use container is intact.
- REFRESH® CELLUVISC® may cause temporary blurring due to its viscosity.
- Store at 59°-77°F (15°-25°C).
- Use before expiration date marked on container.

- RETAIN THIS CARTON FOR FUTURE REFERENCE.

### *Inactive ingredients*

Calcium chloride dihydrate; potassium chloride; purified water; sodium chloride; and sodium lactate.

### *Questions or comments?*

**1.800.678.1605**

**refreshbrand.com**

v1.0DFL4554

### PRINCIPAL DISPLAY PANEL

NDC 0023-4554-30
Preservative-free
Refresh®
Celluvisc®
Lubricant Eye Gel
EXTRA-STRENGTH GEL
Soothes & Comforts
Dry, Irritated Eyes30 Single-Use Containers
0.01 fl oz (0.4 mL) each Sterile


## REFRESH CELLUVISC

carboxymethylcellulose sodium gel

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:0023-4554 |

| Route of Administration | OPHTHALMIC |
|---|---|

## Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|---|---|
| **CARBOXYMETHYLCELLULOSE SODIUM** (UNII: K679OBS311) (CARBOXYMETHYLCELLULOSE - UNII:05JZI7B19X) | CARBOXYMETHYLCELLULOSE SODIUM | 10 mg in 1 mL |

## Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **CALCIUM CHLORIDE** (UNII: M4I0D6VV5M) | |
| **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:0023-4554-05 | 5 in 1 CARTON | 10/04/1989 | |
| **1** | | 0.4 mL in 1 VIAL, SINGLE-USE; Type 0: Not a Combination Product | | |
| **2** | NDC:0023-4554-30 | 30 in 1 CARTON | 10/04/1989 | |
| **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 Drug | M018 | 10/04/1989 | |

## Labeler - Allergan, Inc. (144796497)

Revised: 8/2022 Allergan, Inc.