REFRESH CELLUVISC- carboxymethylcellulose sodium gel
Allergan, Inc.

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.

REFRESH® CELLUVISC®
Lubricant Eye Gel
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.
- REFRESH® CELLUVISC® may cause temporary blurring due to its viscosity.
- Store at 59°-86°F (15°-30°C).
- Use before expiration date marked on container.
- RETAIN THIS CARTON FOR FUTURE REFERENCE.

Inactive ingredients

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

Questions or comments?

☎ 1.800.433.8871

refreshbrand.com

PRINCIPAL DISPLAY PANEL

NDC 0023-4554-30

Preservative-free

Refresh®
Celluvisc®
Lubricant Eye Gel

Soothing Gel

Long-lasting relief
for dry eyes in a
soothing gel formula

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

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How To Use:
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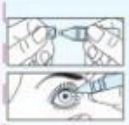

refreshbrand.com

Allergan, Inc.
2525 Dupont Drive
Irvine, CA 92612, U.S.A.

Made in the U.S.A.

Drug Facts

Active ingredient	Purpose
Carboxymethylcellulose sodium 1%	Eye lubricant

Uses
- For the temporary relief of burning, irritation, and discomfort due to dryness of the eye or exposure to wind or sun.
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- **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°-86°F (15°-30°C).
- Use before expiration date marked on container.
- RETAIN THIS CARTON FOR FUTURE REFERENCE.

Inactive ingredients Calcium chloride; potassium chloride; purified water; sodium chloride; and sodium lactate.

Questions or comments?
☎ 1.800.433.8671
refreshbrand.com

Refresh®

Make your dry eyes feel better with the drops doctors recommend.

REFRESH® CELLUVISC® Lubricant Eye Gel provides extra strength moisturizing relief plus protection for dry, irritated eyes with our thickest gel formula for sensitive eyes.

REFRESH® CELLUVISC® 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.

refreshbrand.com

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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 FINAL	part349	10/04/1989	

Labeler - Allergan, Inc. (144796497)

Revised: 2/2018 Allergan, Inc.