REFRESH CELLUVISC- carboxymethylcellulose sodium gel
AvPAK

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Carboxymethylcellulose Sodium Ophthalmic Gel !% PF
(preservative free)

Lubricant Eye Gel

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.
- Store at room temperature 15°-30°C (59°-86°F).
- RETAIN THIS CARTON FOR FUTURE REFERENCE.

Inactive ingredients

Calcium chloride, hydrochloric acid, magnesium chloride, potassium chloride, puried water, sodium chloride, sodium hydroxide and sodium lactate.

Questions or comments?

1-855-361-3993

PRINCIPAL DISPLAY PANEL

REFRESH CELLUVISC

carboxymethylcellulose sodium gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50268-065
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**
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM LACTATE (UNII: TU7HW0W0QT)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:50268-065-30	30 in 1 CARTON	11/18/2020	
1		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
unapproved drug other		11/18/2020	

Labeler - AvPAK (832926666)

Revised: 11/2020 AvPAK