REFRESH RELIEVA PF- carboxymethylcellulose sodium and glycerin solution/ drops
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® RELIEVA™ PF (Unit Dose)
Lubricant Eye Drops
Drug Facts

Active ingredients

Carboxymethylcellulose sodium 0.5%

Glycerin 1.0%

Purpose

Eye lubricant

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, 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
- *Follow your eye doctor’s instructions if you are using this product after an eye surgery (e.g., LASIK) to relieve eye dryness and discomfort.

Other information

- Use only if single-use container is intact.
- Use before expiration date marked on container.
- Store at 59°-77°F (15°-25°C).
- Protect from sunlight.
- RETAIN THIS CARTON FOR FUTURE REFERENCE.

Inactive ingredients

Calcium chloride dihydrate; erythritol; levocarnitine; magnesium chloride hexahydrate; potassium chloride; purified water; sodium lactate; and sodium hyaluronate. May contain hydrochloric acid and/or sodium hydroxide (to adjust pH).

Questions or comments?

☎ 1.800.678.1605
refreshbrand.com

PRINCIPAL DISPLAY PANEL

Refresh®
RELIEVA™ PF
Lubricates and Protects
Recommended for

Sensitive Eyes
PRESERVATIVE-FREE
30 Vials 0.01 fl oz (0.4 mL) Sterile

PRINCIPAL DISPLAY PANEL

Refresh®
RELIEVA™ PF
Lubricates and Protects
Recommended for Sensitive Eyes
5 Vials 0.01 fl oz (0.4 mL) Sterile

REFRESH RELIEVA PF

carboxymethylcellulose sodium and glycerin solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0023-4515
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
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	10 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	
ERYTHRITOL (UNII: RA96B954X6)	
LEVOCARNITINE (UNII: 0G389FZZ9M)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM LACTATE (UNII: TU7HW0W0QT)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	

SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0023-4515-30	30 in 1 CARTON	09/25/2020	
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
OTC monograph final	part349	09/25/2020	

REFRESH RELIEVA PF

carboxymethylcellulose sodium and glycerin solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0023-9537
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
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	10 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	
ERYTHRITOL (UNII: RA96B954X6)	
LEVOCARNITINE (UNII: 0G389FZZ9M)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM LACTATE (UNII: TU7HW0W0QT)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0023-9537-05	5 in 1 CARTON	09/25/2020	
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
OTC monograph final	part349	09/25/2020	

Labeler - Allergan, Inc. (144796497)

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