REFRESH DIGITAL PF- carboxymethylcellulose sodium, glycerin and polysorbate 80 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® DIGITAL PRESERVATIVE-FREE
Drug Facts

Active ingredients

Carboxymethylcellulose sodium 0.5%

Glycerin 1%

Polysorbate 80 0.5%

Purpose

Eye lubricant

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.

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).
- RETAIN THIS CARTON FOR FUTURE REFERENCE.

Inactive ingredients

Boric acid; carbomer copolymer type A; castor oil; erythritol; levocarnitine; purified water; and sodium hydroxide (to adjust pH).

Questions or comments?

☎ **1.800.678.1605**

refreshbrand.com

v1.0DFL6954

Principal Display Panel

NDC 0023-6954-30

Refresh®
DIGITAL PF
Lubricant Eye Drops
For Eye Dryness
with HydroCell™
Fast-acting Relief
That Soothes Your Eyes
30 Vials (0.01 fl oz (0.4 mL) each Sterile

REFRESH DIGITAL PF

carboxymethylcellulose sodium, glycerin and polysorbate 80 solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0023-6954
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
POLYSORBATE 80 (UNII: 6OZP39ZG8H) (POLYSORBATE 80 - UNII:6OZP39ZG8H)	POLYSORBATE 80	5 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BORIC ACID (UNII: R57ZHV85D4)	
CARBOMER COPOLYMER TYPE A (UNII: 71DD5V995L)	
CASTOR OIL (UNII: D5340Y2I9G)	
ERYTHRITOL (UNII: RA96B954X6)	
LEVOCARNITINE (UNII: 0G389FZZ9M)	
WATER (UNII: 059QF0KO0R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0023-6954-30	30 in 1 CARTON	08/01/2020	
1		0.4 mL in 1 VIAL, PLASTIC; Type 0: Not a Combination Product		
2	NDC:0023-6954-05	1 in 1 CARTON	08/01/2020	
2		.40 mL in 1 VIAL, PLASTIC; 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	08/01/2020	

Labeler - Allergan, Inc. (144796497)

Revised: 11/2022 Allergan, Inc.