

REFRESH OPTIVE- carboxymethylcellulose sodium, glycerin solution/ drops
Allergan, Inc.

REFRESH OPTIVE®
Drug Facts

Active ingredients

Carboxymethylcellulose sodium 0.5%

Glycerin 0.9%

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 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. *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).
- RETAIN THIS CARTON FOR FUTURE REFERENCE.

Inactive ingredients

Boric acid; calcium chloride dihydrate; erythritol; levocarnitine; magnesium chloride hexahydrate; potassium chloride; purified water; sodium borate decahydrate; and sodium citrate dihydrate.

Questions or comments?

1.800.678.1605

refresheyedrops.com

v2.0DFL3416

PRINCIPAL DISPLAY PANEL

NDC 0023-3416-30

Preservative-free

Refresh

Optive®

Lubricant Eye Drops

Dual-Action Relief

For Dry, Sensitive Eyes

30 Vials 0.01 fl oz (0.4 mL) each Sterile



REFRESH OPTIVE

carboxymethylcellulose sodium, glycerin solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0023-3416
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Carboxymethylcellulose Sodium (UNII: K679OBS311) (Carboxymethylcellulose - UNII:05JZ17B19X)	Carboxymethylcellulose Sodium	5 mg in 1 mL
Glycerin (UNII: PDC6A3C00X) (Glycerin - UNII:PDC6A3C00X)	Glycerin	9 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
Boric Acid (UNII: R57ZHV85D4)	
Calcium Chloride (UNII: M4I0D6VV5M)	
Erythritol (UNII: RA96B954X6)	
Levocarnitine (UNII: 0G389FZZ9M)	

Magnesium Chloride (UNII: 02F3473H9O)	
potassium chloride (UNII: 660YQ98I10)	
water (UNII: 059QF0KO0R)	
sodium borate (UNII: 91MBZ8H3QO)	
trisodium citrate dihydrate (UNII: B22547B95K)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0023-3416-05	5 in 1 CARTON	08/22/2008	
1		0.4 mL in 1 VIAL, SINGLE-USE; Type 0: Not a Combination Product		
2	NDC:0023-3416-30	30 in 1 CARTON	08/22/2008	
2		0.4 mL in 1 VIAL, SINGLE-USE; Type 0: Not a Combination Product		
3	NDC:0023-3416-60	60 in 1 CARTON	08/22/2008	
3		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	08/22/2008	

Labeler - Allergan, Inc. (144796497)