

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

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**REFRESH OPTIVE® GEL DROPS**  
***Drug Facts***

***Active ingredients***

Carboxymethylcellulose sodium 1%

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. Replace cap after using.**
- **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***

Instill 1 or 2 drops in the affected eye(s) as needed.

***Other information***

- Use only if tape seals on top and bottom flaps are intact.
- Use before expiration date marked on container.

- Discard 90 days after opening.
- 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; PURITE® (stabilized oxychloro complex); sodium borate decahydrate; and sodium citrate dihydrate.

### ***Questions or comments?***

**1.800.678.1605**

**refreshbrand.com**

v1.0DFL5459

### **PRINCIPAL DISPLAY PANEL**

**Refresh**

**Optive®**

**GEL DROPS**

***Lubricant Eye Gel***

**Extended Relief**

**0.33 fl oz (10 mL) Sterile**



## REFRESH OPTIVE DROPS

carboxymethylcellulose sodium and glycerin solution/ drops

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:0023-5459
<b>Route of Administration</b>	OPHTHALMIC		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>carboxymethylcellulose sodium</b> (UNII: K679OBS311) (carboxymethylcellulose - UNII:05JZ17B19X)	carboxymethylcellulose sodium	10 mg in 1 mL
<b>glycerin</b> (UNII: PDC6A3C0OX) (glycerin - UNII:PDC6A3C0OX)	glycerin	9 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>boric acid</b> (UNII: R57ZHV85D4)	
<b>calcium chloride</b> (UNII: M410D6VW5M)	
<b>erythritol</b> (UNII: RA96B954X6)	
<b>levocarnitine</b> (UNII: 0G389FZ Z9M)	
<b>magnesium chloride</b> (UNII: 02F3473H9O)	
<b>potassium chloride</b> (UNII: 660YQ98110)	

<b>water</b> (UNII: 059QF0KO0R)	
<b>sodium borate</b> (UNII: 91MBZ8H3QO)	
<b>sodium chlorite</b> (UNII: G538EBV4VF)	
<b>TRISODIUM CITRATE DIHYDRATE</b> (UNII: B22547B95K)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0023-5459-10	1 in 1 CARTON	09/30/2015	
1		10 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
2	NDC:0023-5459-02	1 in 1 CARTON	09/30/2015	
2		2 mL in 1 BOTTLE, DROPPER; 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	09/30/2015	

**Labeler** - Allergan, Inc. (144796497)

Revised: 7/2022

Allergan, Inc.