

**REFRESH LIQUIGEL- carboxymethylcellulose sodium gel**  
**Allergan, Inc.**

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**REFRESH LIQUIGEL®**  
**Drug Facts**

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

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; magnesium chloride hexahydrate; potassium chloride; purified water; PURITE® (stabilized oxychloro complex); sodium borate decahydrate; and sodium chloride. May contain hydrochloric acid or sodium hydroxide (to adjust pH).

## Questions or comments?



1.800.687.1605

refreshbrand.com

v1.0DFL9205

## PRINCIPAL DISPLAY PANEL

NDC 0023-9205-15

**Refresh**

**Liquigel®**

Lubricant Eye Gel

Soothes & Comforts

Dry, Irritated Eyes

0.5 fl oz (15 mL) Sterile

The image shows the principal display panel for Refresh Liquigel Lubricant Eye Gel. The panel is divided into several sections:

- Left Side:** Contains 'Drug Facts (continued)', 'Inactive ingredients', and 'Questions or comments?' with the phone number 1.800.687.1605 and website refreshbrand.com. A barcode is located at the bottom left.
- Center:** Features the product name 'Refresh Liquigel' in large green and purple font, with 'Lubricant Eye Gel' below it. The tagline 'Soothes & Comforts Dry, Irritated Eyes' is in a blue oval. At the bottom, it says '0.5 fl oz (15 mL) Sterile'. The Allergan logo is at the top.
- Right Side:** Contains 'Drug Facts' with 'Active ingredient' (Carboxymethylcellulose sodium 1%) and 'Purpose' (Eye lubricant). It includes 'Uses', 'Warnings', 'Directions', and 'Other information'. Below this is the slogan 'TRUST YOUR EYES TO REFRESH!' and the website refreshbrand.com.
- Bottom:** Includes the text 'The brand with over 30 years of experience.' and '© 2022 Allergan. All rights reserved. All trademarks are the property of their respective owners.'

Lot: 016152 Exp.: 20074634

**REFRESH LIQUIGEL**

carboxymethylcellulose sodium gel

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:0023-9205
<b>Route of Administration</b>	OPHTHALMIC		

## Active Ingredient/Active Moiety

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED</b> (UNII: K679OBS311) (Carboxymethylcellulose - UNII:05JZI7B19X)	CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED	10 mg in 1 mL

## Inactive Ingredients

<b>Ingredient Name</b>	<b>Strength</b>
<b>BORIC ACID</b> (UNII: R57ZHV85D4)	
<b>CALCIUM CHLORIDE</b> (UNII: M4I0D6VV5M)	
<b>MAGNESIUM CHLORIDE</b> (UNII: 02F3473H9O)	
<b>POTASSIUM CHLORIDE</b> (UNII: 660YQ98I10)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM BORATE</b> (UNII: 91MBZ8H3QO)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>SODIUM CHLORITE</b> (UNII: G538EBV4VF)	
<b>HYDROCHLORIC ACID</b> (UNII: QTT17582CB)	
<b>SODIUM HYDROXIDE</b> (UNII: 55X04QC32I)	

## Packaging

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:0023-9205-03	1 in 1 CARTON	10/04/2001	04/12/2020
1		3 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
2	NDC:0023-9205-15	1 in 1 CARTON	10/04/2001	
2		15 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
3	NDC:0023-9205-02	2 in 1 CARTON	10/04/2001	04/12/2020
3		15 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		

## Marketing Information

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC Monograph Drug	M018	10/04/2001	

