REFRESH LIQUIGEL- carboxymethylcellulose sodium gel
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 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

SOOTHING GEL

Long-lasting relief
for dry eyes in a
soothing gel formula

0.5 fl oz (15 mL) Sterile

REFRESH LIQUIGEL

carboxymethylcellulose sodium gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0023-9205
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Carboxymethylcellulose sodium (UNII: K679OBS311) (Carboxymethylcellulose - UNII:05JZI7B19X)	Carboxymethylcellulose sodium	10 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
Boric acid (UNII: R57ZHV85D4)	
calcium chloride (UNII: M4I0D6VV5M)	
magnesium chloride (UNII: 02F3473H9O)	
potassium chloride (UNII: 660YQ98I10)	
water (UNII: 059QF0KO0R)	
sodium borate (UNII: 91MBZ8H3QO)	
sodium chloride (UNII: 451W47IQ8X)	
sodium chlorite (UNII: G538EBV4VF)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
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

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part349	10/04/2001	

Labeler - Allergan, Inc. (144796497)

Revised: 11/2022 Allergan, Inc.