REFRESH LIQUIGEL- carboxymethylcellulose sodium gel
AvPAK

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Carboxymethylcellulose Sodium Ophthalmic Gel 1%

Lubricant Eye Gel

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 use in the eyes 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 this and all drugs out of the reach of children. If swallowed, get medical help or contact a Poison Control Center right away (1-800-222-1222).

Directions

Shake well before use.

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

Other information

- Do Not Use if imprinted seal on cap is torn, broken or missing.
- Discard 90 days after opening.
- Store at room temperature 15°-30°C (59°-86°F).
- Retain outer carton for full product drug information.

Inactive ingredients

Boric acid, calcium chloride, magnesium chloride, potassium chloride, purified water, sodium borate,

sodium chloride. Stabilized Oxychloro Complex 2.5%. May contain hydrochloric acid and/or sodium hydroxide to adjust pH.

Questions or comments?

1-855-361-3993

Distributed by:
AvKARE

Pulaski, TN 38478

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REFRESH LIQUIGEL

carboxymethylcellulose sodium gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50268-066
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)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50268-066-15	1 in 1 CARTON	11/18/2020	
1		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
unapproved drug other		11/18/2020	

Labeler - AvPAK (832926666)

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