REFRESH TEARS LUBRICANT- carboxymethylcellulose sodium solution/ drops
Navajo Manufacturing Company Inc.

Refresh Tears Lubricant Eye Drops

Active ingredient

Carboxymethylcellulose sodium 0.5%

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 the imprinted seal around the bottle neck is intact and not damaged.**
- **Use before expiration date marked on container.**
- **Store at 59° - 86° F (15° - 30 C)**
- **RETAIN THIS PACKAGING FOR FUTURE REFERENCE.**

Inactive ingredients

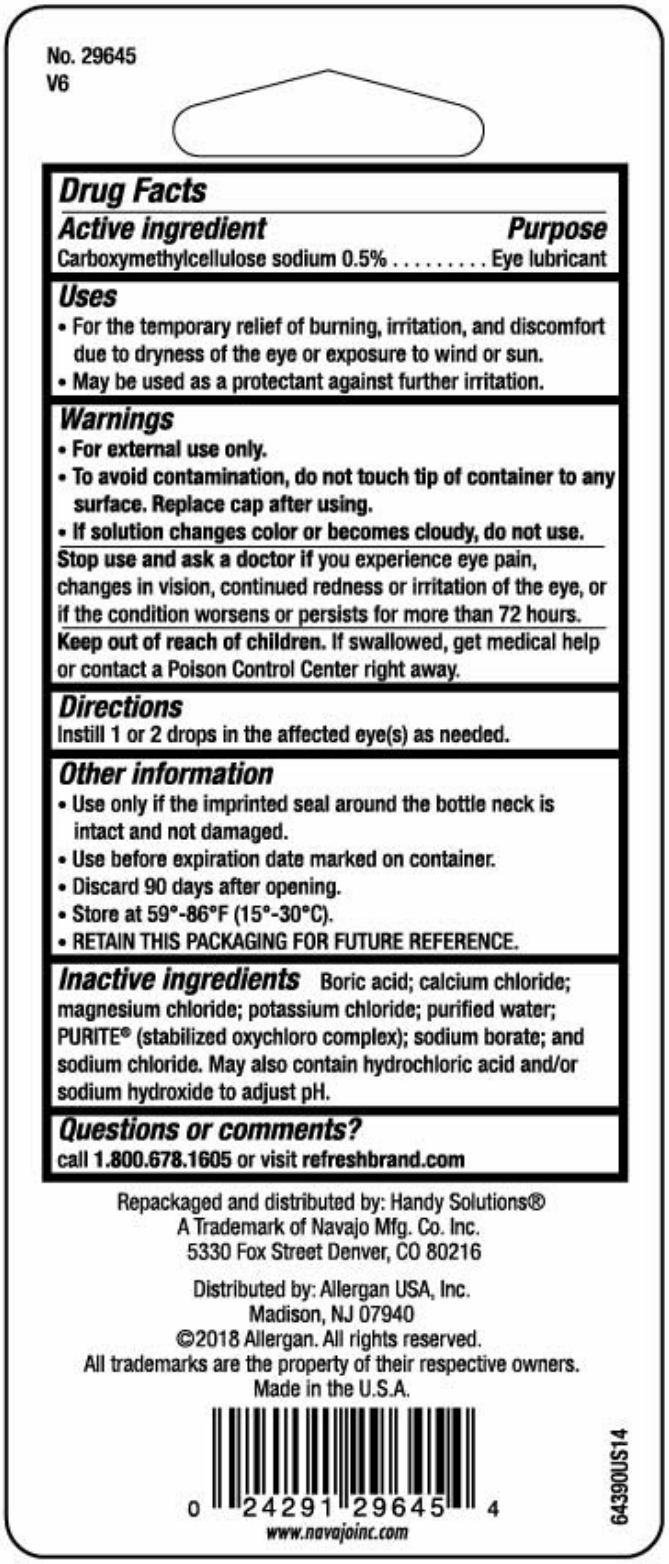
Boric acid; calcium chloride; magnesium chloride; potassium chloride; purified water; PURITE ® (stablilized oxychloro complex); sodium borate; and sodium chloride. May also contain hydrochloric acid and/or sodium hydroxide to adjust pH.

Questions or comments?

1.800.433.8871, M-F 6 AM - 4:30 PM Pacific Time

refreshbrand.com

Principal Display Panel



REFRESH TEARS LUBRICANT

carboxymethylcellulose sodium solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67751-130(NDC:0023-0798)
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311) (CARBOXYMETHYLCELLULOSE - UNII:05JZI7B19X)	CARBOXYMETHYLCELLULOSE SODIUM	5 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 CHLORITE (UNII: G538EBV4VF)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67751-130-01	3 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	12/22/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M018	09/01/2004	

Labeler - Navajo Manufacturing Company Inc. (091917799)

Establishment

Name	Address	ID/FEI	Business Operations
Navajo Manufacturing Company Inc.		136941411	relabel(67751-130) , repack(67751-130)

Establishment

Name	Address	ID/FEI	Business Operations
Allergan, Inc.		362898611	manufacture(67751-130)

Revised: 10/2024 Navajo Manufacturing Company Inc.