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 ® (stabilized 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

<table>
<thead>
<tr>
<th>Product Type</th>
<th>HUMAN OTC DRUG</th>
<th>Item Code (Source)</th>
<th>NDC:67751-130(NDC:0023-0798)</th>
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<tbody>
<tr>
<td>Route of Administration</td>
<td>OPHTHALMIC</td>
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Drug Facts

Active ingredient  Purpose
Carboxymethylcellulose sodium 0.5%  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.
- Discard 90 days after opening.
- 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® (stabilized oxychloro complex); sodium borate; and sodium chloride. May also contain hydrochloric acid and/or sodium hydroxide to adjust pH.

Questions or comments?
call 1.800.678.1665 or visit refreshbrand.com

Repackaged and distributed by: Handy Solutions®
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5330 Fox Street Denver, CO 80219

Distributed by: Allergan USA, Inc.
Maddison, NJ 07940
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Made in the U.S.A.
### Active Ingredient/Active Moiety

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<tr>
<th>Ingredient Name</th>
<th>Basis of Strength</th>
<th>Strength</th>
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<tr>
<td>CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OB5311) (CARBOXYMETHYLCELLULOSE - UNII:05JZ17B19X)</td>
<td>CARBOXYMETHYLCELLULOSE SODIUM</td>
<td>5 mg in 1 mL</td>
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### Inactive Ingredients

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<tr>
<th>Ingredient Name</th>
<th>Strength</th>
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<tr>
<td>BORIC ACID (UNII: R57ZH85D4)</td>
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<tr>
<td>CALCIUM CHLORIDE (UNII: M410D6VV5M)</td>
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<tr>
<td>MAGNESIUM CHLORIDE (UNII: 02F3473H9O)</td>
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<td>POTASSIUM CHLORIDE (UNII: 660YQ98I10)</td>
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<tr>
<td>WATER (UNII: 059QF0KO0R)</td>
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<tr>
<td>SODIUM CHLORITE (UNII: G538EBV4VF)</td>
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<td>SODIUM BORATE (UNII: 91MBZ8H3Q0)</td>
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<tr>
<td>SODIUM CHLORIDE (UNII: 451W47I08X)</td>
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<td>HYDROCHLORIC ACID (UNII: QTT17582CB)</td>
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<td>SODIUM HYDROXIDE (UNII: 55X04QC32I)</td>
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### Packaging

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

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<th>Application Number or Monograph Citation</th>
<th>Marketing Start Date</th>
<th>Marketing End Date</th>
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<tr>
<td>OTC monograph final</td>
<td>part349</td>
<td>09/01/2004</td>
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### Labeler
- Navajo Manufacturing Company Inc. (091917799)

### Establishment

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

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

Revised: 3/2023