LUBRICATING PLUS- carboxymethylcellulose sodium solution/drops
Major Pharmaceuticals

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.

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Major Pharmaceuticals Lubricating Plus Drug Facts

Active ingredient (in each single-use container)
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

Do not use
if solution changes color or becomes cloudy

When using this product
to avoid contamination

• do not touch tip of container to any surface
• do not reuse
• once opened, discard
• do not touch unit-dose tip to eye

Stop use and ask a doctor if

• you experience eye pain
• changes in vision occur
• redness or irritation of the eye continues
• redness or irritation of the eye 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 (1-800-222-1222).

Directions
• to open, twist and pull tab to remove
• instill 1 or 2 drops in the affected eye(s) as needed and discard container

Other information
• store at 20-25°C (68-77°F)
• RETAIN THIS CARTON FOR FUTURE REFERENCE.

Inactive ingredients
calcium chloride dihydrate, magnesium chloride hexahydrate, potassium chloride, sodium chloride, sodium lactate solution, water for injection. May also contain hydrochloric acid and/or sodium hydroxide to adjust pH.

Questions or comments?
1-800-616-2471

Package/Label Principal Display Panel
COMPARE TO the active ingredient of REFRESH PLUS®
Lubricating PLUS
Lubricant Eye Drops
Immediate, soothing relief for dry eyes
Moisturizing Relief
Carboxymethylcellulose Sodium 0.5%
30 Sterile Single-Use Containers
0.01 FL OZ (0.4 mL) each
Actual Size
LUBRICATING PLUS
carboxymethylcellulose sodium solution/ drops

Product Information

Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:0904-6329
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Route of Administration | OPHTHALMIC |  | |

Active Ingredient/Active Moiety

<table>
<thead>
<tr>
<th>Ingredient Name</th>
<th>Basis of Strength</th>
<th>Strength</th>
</tr>
</thead>
<tbody>
<tr>
<td>CARBOXYMETHYLCELLULOSE SODIUM (UNII: K6790BS311) (CARBOXYMETHYLCELLULOSE - UNII:05JZI7B19X)</td>
<td>CARBOXYMETHYLCELLULOSE SODIUM</td>
<td>0.5 g in 100 mL</td>
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Inactive Ingredients

<table>
<thead>
<tr>
<th>Ingredient Name</th>
<th>Strength</th>
</tr>
</thead>
<tbody>
<tr>
<td>CALCIUM CHLORIDE (UNII: M4I0D6VV5M)</td>
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</tbody>
</table>
MAGNESIUM CHLORIDE (UNII: 02F3473H0O)
POTASSIUM CHLORIDE (UNII: 660YQ98I10)
SODIUM CHLORIDE (UNII: 451W47IQ8X)
SODIUM LACTATE (UNII: TU7HW0W0QT)
WATER (UNII: 059QF0KO0R)
SODIUM HYDROXIDE (UNII: 55X04QC32I)
HYDROCHLORIC ACID (UNII: QTT17582CB)

### Packaging

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<thead>
<tr>
<th>#</th>
<th>Item Code</th>
<th>Package Description</th>
<th>Marketing Start Date</th>
<th>Marketing End Date</th>
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<tbody>
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<td>1</td>
<td>NDC:0904-6329-46</td>
<td>6 in 1 CARTON</td>
<td>05/09/2013</td>
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<tr>
<td>1</td>
<td></td>
<td>5 in 1 POUCH</td>
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<tr>
<td>1</td>
<td></td>
<td>0.4 mL in 1 VIAL, SINGLE-USE; Type 0: Not a Combination Product</td>
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<td>05/31/2013</td>
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### Marketing Information

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<th>Marketing Start Date</th>
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<tr>
<td>OTC monograph final</td>
<td>part349</td>
<td>05/09/2013</td>
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**Labeler** - Major Pharmaceuticals (191427277)