RAIN RENEWAL EYE DROPS- carboxymethylcellulose sodium solution/ drops Rain Eye Drops, LLC

Rain Eye Drops - 70 ct Pack/Label - US

Carboxymethylcellulose sodium 0.5%

Carboxymethylcellulose sodium.....Lubricant

- 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 the product

- do not reuse
- once opened, discard
- to avoid contamination do not touch tip of container to any surface
- 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 accidentally swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) immediately.

Directions

- To open, twist and pull tab to remove
- Instill 1 or 2 drops in the affected eye(s) as needed and discard container
- If used for post-operative (e.g., LASIK) dryness and discomfort, follow your eye doctor's instructions.

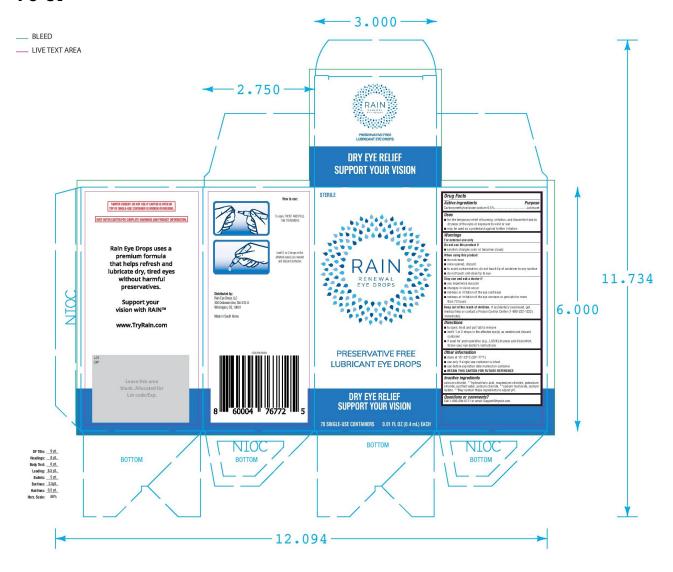
Other information

- Store at 15°-25°C (59°-77°F)
- Use only if single-use container is intact
- Use before expiration date marked on container

RETAIN THIS CARTON FOR FUTURE REFERENCE

calcium chloride, **hydrochloric acid, magnesium chloride, potassium chloride, purified water, sodium chloride, **sodium hydroxide, sodium lactate. **May contain these ingredients to adjust pH.

70 ct



RAIN RENEWAL EYE DROPS

carboxymethylcellulose sodium solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79662-002
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM (UNII: K6790BS311) (CARBOXYMETHYLCELLULOSE - UNII:05JZ17B19X)	CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM	0.5 g in 100 mL	

Inactive Ingredients		
Ingredient Name	Strength	
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)		

MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
WATER (UNII: 059QF0KO0R)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
SODIUM LACTATE (UNII: TU7HW0W0QT)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

F	Packaging			
#	tem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79662- 002-70	70 in 1 BOX	09/18/2020	11/30/2023
1	1.	0.4 mL in 1 VIAL, DISPENSING; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M018	09/18/2020	

Labeler - Rain Eye Drops, LLC (117579363)

Revised: 11/2023 Rain Eye Drops, LLC