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

Rain Renewal Eye Drops - 30 ct

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.

- to avoid contamination do not touch tip of container to any surface
- do not touch unit-dose tip to eye.
- do not reuse
- once opened, discard
- do not use if solution changes color or becomes cloudy.
- remove contact lenses before using

Stop Use and ASK Doctor if

- you feel eye pain
- changes in vision occurs
- continued redness or irritations of the eye last
- conditions worsens or last more than 72 hours

Stop use and ask a doctor if

- you experience eye pain
- changes in vision occur
- redness or irritation of the eye lasts
- conditions worsens or lasts 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
- Children under 6 years of age; ask a doctor

Other information

- Store at 15°-30°C (59°-86°F)
- Protect from light

RETAIN THIS CARTON FOR FUTURE REFERENCE

calcium chloride, hydrochloric acid, magnesium chloride, potassium chloride, purified water, sodium chloride, sodium hydroxide, sodium lactate.

Rain Renewal Eye Drops - 30 Ct - Updated package

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: K679OBS311) (CARBOXYMETHYLCELLULOSE - UNII:05JZI7B19X)	CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM	0.5 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
WATER (UNII: 059QF0KO0R)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
SODIUM LACTATE (UNII: TU7HW0W0QT)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79662-002-30	30 in 1 BOX	02/15/2021	
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	02/15/2021	

Labeler - Rain Eye Drops, LLC (117579363)

Revised: 10/2024 Rain Eye Drops, LLC