

GREEN GUARD LUBRICANT EYE DROPS- carboxymethylcellulose sodium solution
UniFirst First Aid Corporation

Green Guard Lubricant Eye Drops

Drug Facts

Active ingredient

Carboxymethylcellulose sodium 0.5%

Purpose

Eye lubricant

Uses

- for the temporary relief of burning, irritation, and discomfort due to dryness of the eyes or exposure to wind or sun
- for use a protectant against further irritation

Warnings

For external use only

Do not use

- if solution changes color or becomes cloudy

When using this 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 od 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 59°-77°F (15°-25°C)

- use only if single-use container is intact
- use before expiration date marked on container
- **RETAIN THIS CARTON FOR FUTURE REFERENCE**

Inactive ingredients

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

** May contain these ingredients to adjust pH

Questions or comments? 1-800-869-6970

Preservative-free

Lubricant eye drops

NDC 47682-194-84

*Compare to the active ingredients in Refresh Plus®

Carboxymethylcellulose Sodium 0.5% Eye Lubricant

Immediate, long-lasting relief for dry, irritated eyes

Green Guard

Sterile

30 Single-Use Containers

0.01 fl oz (0.4 mL) each



GREEN GUARD LUBRICANT EYE DROPS

carboxymethylcellulose sodium solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:47682-194
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311) (CARBOXYMETHYLCELLULOSE - UNII:05JZ17B19X)	CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED	0.5 g in 100 mL

Inactive Ingredients

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47682-194-83	30 in 1 BOX	09/01/2021	
1		0.4 mL in 1 AMPULE; 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	09/01/2021	

Labeler - UniFirst First Aid Corporation (832947092)

Revised: 1/2024

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