EYE DROPS- carboxymethylcellulose sodium liquid
Foshan Sugar Max Cosmetics CO.,Ltd

EYE DROPS

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Carboxymethylcellulose sodium 0.25%

Eye lubricant

Uses For the temporary relief of burning, irritation, anddiscomfort due to dryness of the eye or exposure towind or sun.
May be used as a protectant against further irritation.

For external use only.
To avoid contamination, do not touch tip of container toany surface. Replace cap after using.
lf solution changes color or becomes cloudy, do not use.

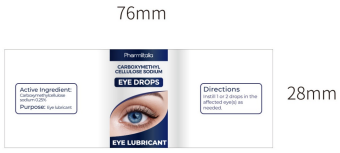
Stop use and ask a doctor if you experience eye pain,changes in vision, continued redness or irritation of theeye, or if the condition worsens or persists for more than72 hours.

Keep out of reach of children. lf swallowed, get medicalhelp or contact a Poison Control Center right away.

Directions Instilllor 2 drops in the affected eye(s) asneeded

Use only if tape seals on top and bottom flaps areintact.
Use before expiration date marked on container.
Discard 90 days after opening.

calciumchloride
Dequest
magnesium chloride
potassium chloride
purified water
sodium bicarbonate
sodium chloride
sodium perborate
sodium phosphate



EYE DROPS

carboxymethylcellulose sodium liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:84938-001
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311) (CARBOXYMETHYLCELLULOSE - UNII:05JZI7B19X)	CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED	0.25 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
CALCIUM CHLORIDE ANHYDROUS (UNII: OFM21057LP)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM PERBORATE (UNII: Y52BK1W96C)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
SODIUM PHOSPHATE (UNII: SE337SVY37)	
AMINOTRIS(METHYLENEPHOSPHONIC ACID) (UNII: 1Y702GD0FG)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84938-001-01	2 in 1 BOX	11/14/2024	
1		15 mL in 1 BOTTLE; Type 0: Not a Combination		

1		Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M018	11/14/2024	

Labeler - Foshan Sugar Max Cosmetics CO.,Ltd (700689935)

Establishment

Name	Address	ID/FEI	Business Operations
Foshan Sugar Max Cosmetics CO.,Ltd		700689935	manufacture(84938-001)

Revised: 11/2024 Foshan Sugar Max Cosmetics CO.,Ltd