

PERFECTEARS- carboxymethylcellulose sodium, carboxymethylcellulose liquid
Cetel Scientific LLC

Active ingredient

Carboxymethylcellulose sodium 0.5%

Purpose

Eye Lubricant

Uses

- For temporary relief of burning, irritation and discomfort due to dryness of the eye or exposure to wind or sun
- 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 touch tip of container to any surface to avoid contamination
- do not reuse. Once opened, discard.

Stop use and ask a doctor if

- you experience eye pain, changes in vision, continued redness or irritation of the eye
- condition 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.

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

- use before expiration date marked on container.
- store at 59°-86°F (15°-30°C).

Inactive ingredients

Calcium chloride, magnesium chloride, potassium chloride, purified water, gelatin,

sodium chloride; and sodium lactate. May also contain hydrochloric acid and/or sodium hydroxide to adjust pH.

Product label



PERFECTEARS

carboxymethylcellulose sodium, carboxymethylcellulose liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84535-001
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311) (CARBOXYMETHYLCELLULOSE - UNII:05JZI7B19X)	CARBOXYMETHYLCELLULOSE SODIUM	5 mg in 1 mL
CARBOXYMETHYLCELLULOSE (UNII: 05JZI7B19X) (CARBOXYMETHYLCELLULOSE - UNII:05JZI7B19X)	CARBOXYMETHYLCELLULOSE	5 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM LACTATE (UNII: TU7HW0W0QT)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
GELATIN (UNII: 2G86QN327L)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84535-001-01	0.4 mL in 1 CONTAINER; Type 0: Not a Combination Product	07/24/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M018	07/24/2024	

Labeler - Cetel Scientific LLC (079453529)

Revised: 7/2024

Cetel Scientific LLC