

EYS-ED- carboxymethylcellulose sodium 0.5% solution/ drops

Eyes-Plus LLC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Eys Lubricant Eye Drops Single Use

Active ingredient Purpose

Carboxymethylcellulose sodium 0.5%.....Eye lubricant

Uses

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 reuse. Once opened, discard.
- Do not touch unit does tip to eye.
- If 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 the eye, or if the 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.

- If used for post-operative (e.g., SILAS) dryness and discomfort, follow your eye doctor's instructions.

Other Information

- Use only if single-use container is intact.
- Use before expiration date marked on container.
- Store at 59° to 86°F (15° to 30°C)
- RETAIN THIS CARTON FOR FUTURE REFERENCE.

Inactive ingredients

calcium chloride, magnesium chloride, potassium chloride, purified water, sodium chloride, and sodium lactate. May also contain hydrochloric acid and/or sodium hydroxide to adjust pH.

Distributed by:

Eys Plus LLC

Lomita, CA 90717'

Made in Korea



EYS-ED

carboxymethylcellulose sodium 0.5% solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:57349-397
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

Inactive Ingredients

Ingredient Name	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM LACTATE (UNII: TU7HW0W0QT)	

POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:57349-397-04	5 in 1 BOX	06/13/2018	
1		0.4 mL in 1 VIAL; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part349	06/13/2018	

Labeler - Eyes-Plus LLC (078751656)

Registrant - Eyes-Plus LLC (078751656)

Revised: 6/2018

Eyes-Plus LLC