

EQUATE RESTORE PLUS LUBRICANT EYE- carboxymethylcellulose sodium solution/ drops
Wal-Mart Stores Inc.

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.

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-dose 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., LASIK) 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°-86°F (15°-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:

Wal-Mart Stores Inc.

Bentonville, AR 72716

Made in Korea



EQUATE RESTORE PLUS LUBRICANT EYE

carboxymethylcellulose sodium solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49035-657	
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	
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)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49035-657-70	70 in 1 CARTON	02/13/2017	
1		0.4 mL in 1 VIAL, SINGLE-USE; Type 1: Convenience Kit of Co-Package		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part349	09/05/2013		

Labeler - Wal-Mart Stores Inc. (051957769)

Revised: 10/2021

Wal-Mart Stores Inc.