RESTORE PLUS SINGLE VIAL- carboxmethylcellulose sodium solution/ drops
United Exchange Corp.

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.

Refresh Plus Single Vial Eye Drops 0.4 ml Bulk 6577ZZ (2018)

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 eyes.
- 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 in the condtion 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 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

RESTORE PLUS SINGLE VIAL

carboxmethylcellulose sodium solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:65923-657
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**
SODIUM LACTATE (UNII: TU7HW0W0QT)	
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65923-657-04	0.4 mL in 1 VIAL; Type 0: Not a Combination Product	01/01/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part349	01/01/2016	

Labeler - United Exchange Corp. (840130579)

Registrant - United Exchange Corp. (840130579)

Revised: 11/2021 United Exchange Corp.