UP AND UP LUBRICANT- carboxymethylcellulose sodium solution/ drops
Target Corporation

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.

Up & Up Lubricant Single Use Eye Drops 70 ct. 675

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

When using this product

- do not touch tip of container to any surface to avoid contamination
- do not reuse
- once opened, discard
- do not use if this solution changes color or becomes cloudy

Stop use and ask a doctor if

- you feel eye pain
- changes in vision occur
- continued redness or irritation of the eye lasts
- condition worsens or lasts more than 72 hours

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach children. If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) 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
- children under 6 years of age: ask a doctor

Other information

- store between 15-30°C (59-86°F)
- protect from light

Inactive ingredients

calcium chloride, hydrochloric acid, magnesium chloride, potassium chloride. purified water, sodium chloride, sodium hydroxide, sodium lactate

Distributed by Target Corporation

Made in South Korea

UP AND UP LUBRICANT

carboxymethylcellulose sodium solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-126

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)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
WATER (UNII: 059QF0KO0R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SODIUM LACTATE (UNII: TU7HW0W0QT)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-126-70	70 in 1 CARTON	01/26/2021	
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	01/26/2021	

Labeler - Target Corporation (006961700)

Revised: 12/2022 Target Corporation