UP AND UP LUBRICANT REFRESHING- 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 Preservative Free Eye Drops 70 ct. NBE Refresh Plus 636 (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

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
- remove contact lenses before using

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 of 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
- children under 6 years of age: ask a doctor

Other information

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

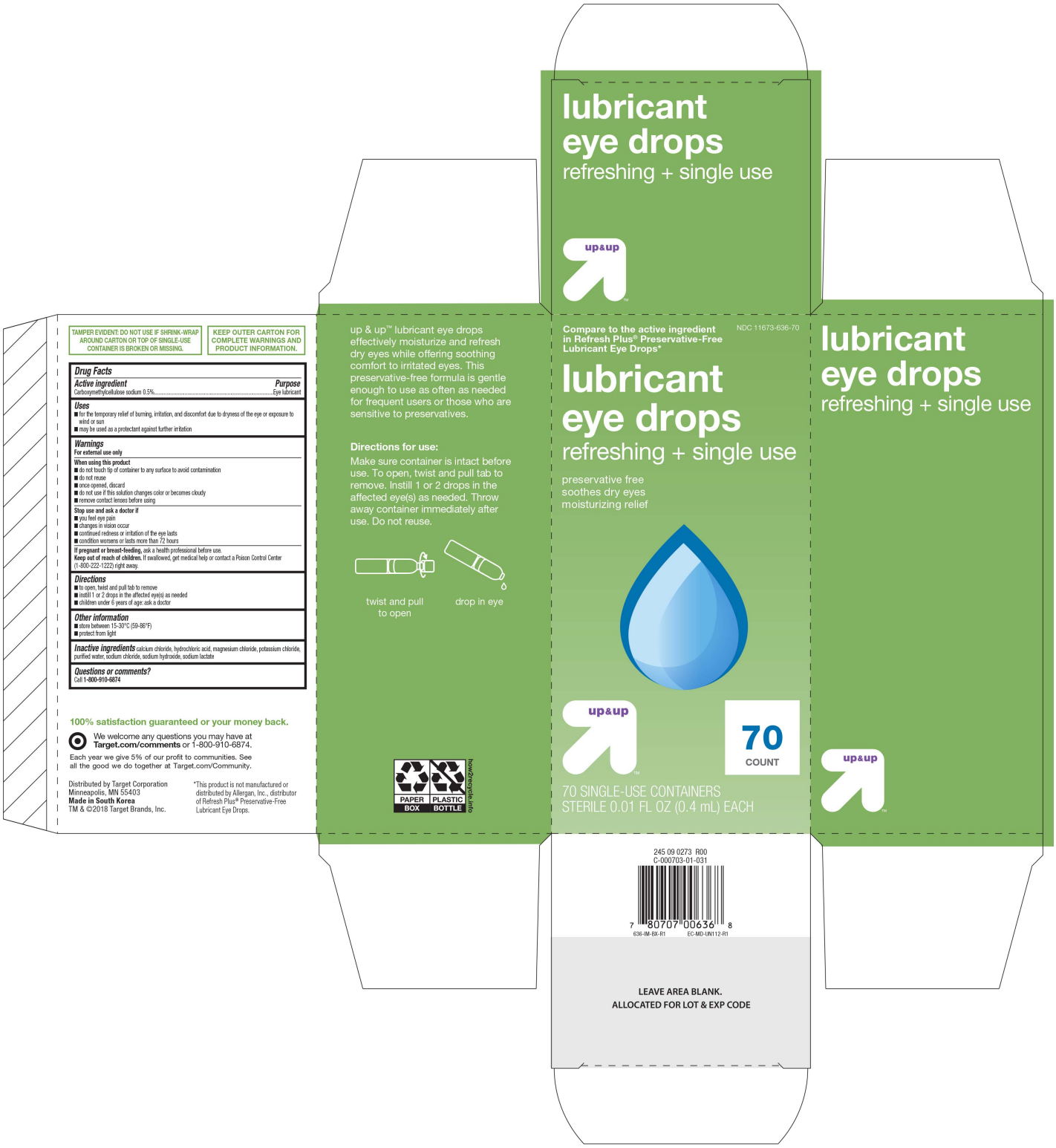
Inactive ingredients calcium chloride, hydrochloric acid, magnesium chloride, potassium chloride, purified water, sodium chloride, sodium hydroxide, sodium lactate

Distributed by:

Target Corporation

Minneapolis, MN 55403

Made in South Korea



UP AND UP LUBRICANT REFRESHING

carboxymethylcellulose sodium solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-636
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)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SODIUM LACTATE (UNII: TU7HW0W0QT)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:11673-636-70	70 in 1 CARTON	04/19/2018	
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	04/19/2018	

Labeler - Target Corporation (006961700)

Revised: 4/2018 Target Corporation