LUBRICANT EYE DROPS PRESERVATIVE FREE- carboxymethylcellulose sodium solution/ drops
Target Corporation

Target Lubricant Eye Drops Preservative Free 70 ct (PLD)

Active ingredient
Carboxymethylcellulose sodium 0.5%

Purpose
Carboxymethylcellulose sodium.....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.

Do not use if solution changes color or becomes cloudy.

When using the product
- do not reuse
- once opened, discard
- to avoid contamination do not touch tip of container to any surface
- do not touch unit-dose tip to eye.

Stop use and ask a doctor if
- you experience eye pain
- changes in vision occur
- redness or irritation of the eye continues
- redness or irritation of the eye worsens or persists for more than 72 hours

Keep out of reach of children. If accidentally swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) immediately.

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
- Store at 15°-25°C (59°-77°F)
- Use only if single-use container is intact
- Use before expiration date marked on container
- **RETAIN THIS CARTON FOR FUTURE REFERENCE**

Inactive ingredients
Calcium chloride, magnesium chloride, potassium chloride, purified water, sodium

chloride, and sodium lactate. May contain sodium hydroxide and/or hydrochloric acid to adjust pH.

LUBRICANT EYE DROPS PRESERVATIVE FREE

carboxymethylcellulose sodium solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-150
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311) (CARBOXYMETHYLCELLULOSE - UNII:05JZI7B19X)	CARBOXYMETHYLCELLULOSE SODIUM	0.5 g in 100 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)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-150-01	70 in 1 BOX	02/12/2019	
1		0.4 mL in 1 VIAL, DISPENSING; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M018	02/12/2019	

Labeler - Target Corporation (006961700)

Registrant - Unimed (689852052)

Establishment

Name	Address	ID/FEI	Business Operations
KC Pharmaceuticals, Inc.		174450460	pack(11673-150) , label(11673-150)

Establishment

Name	Address	ID/FEI	Business Operations
Unimed		689852052	manufacture(11673-150)

Revised: 12/2023 Target Corporation