LUBRICANT EYE DROPS- carboxymethylcellulose sodium solution/ drops
KC Pharmaceuticals, Inc.

KC Pharmaceuticals (Packager & labeler) NDC for Cintas Lubricant Eye Drops 5 ct (PLD)

Carboxymethylcellulose sodium 0.5%

Eye Lubricant

- 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

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, pull and twist 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

Calcium Chloride, **Hydrochloric acid, Magnesium Chloride, Potassium Chloride, Purified water, Sodium Chloride, **Sodium Hydroxide, Sodium Lactate. **May contain these ingredients to adjust pH.

Warnings

For external use only

Do not use this product if:

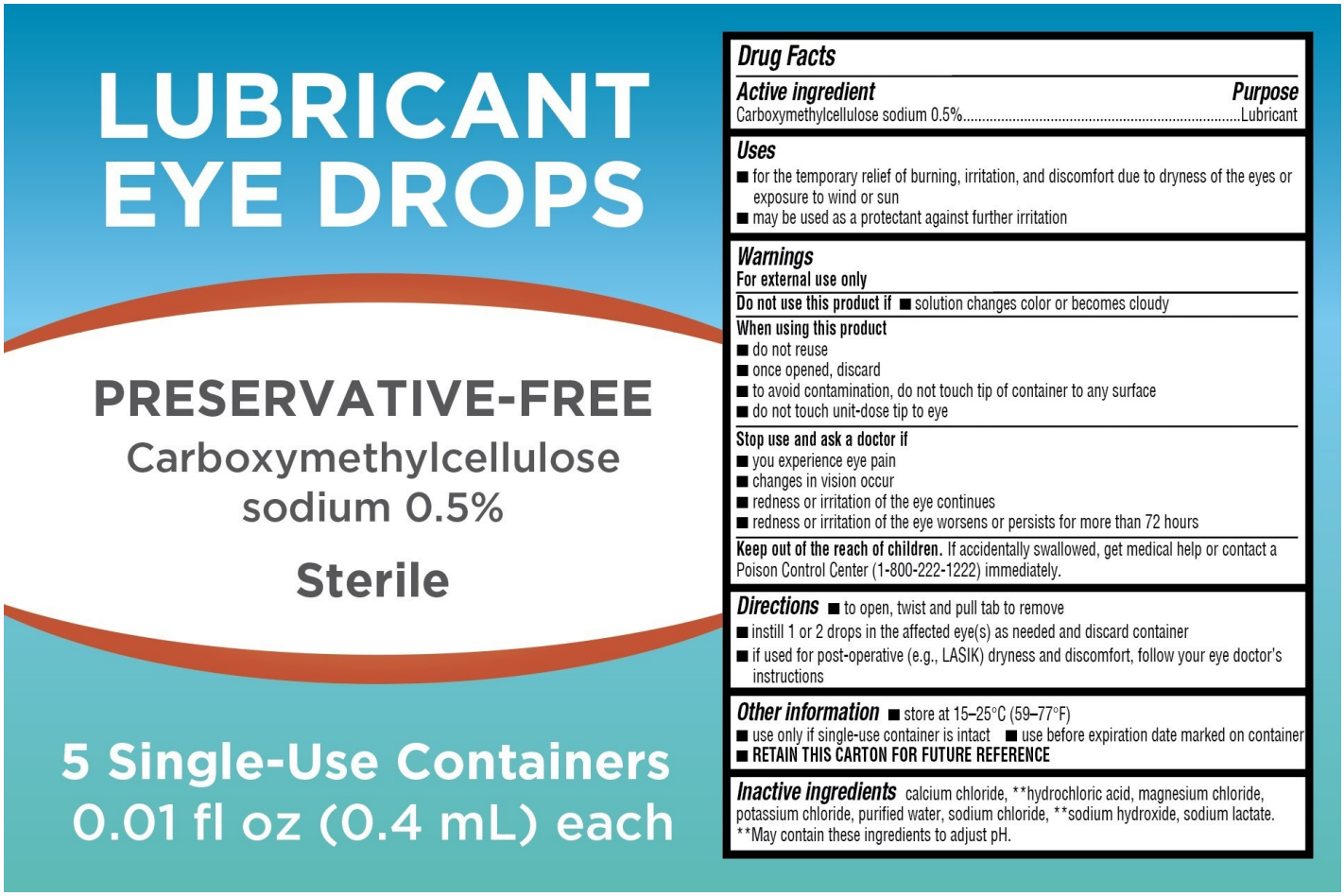
- solution changes color or becomes cloudy

When using this product

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

Stop use and ask a doctor if you experience any of the following

- you experience eye pain
- changes in vision occur
- redness or irritation of the eye continues
- redness or irritation of the eye gets worse or lasts more than 72 hours



LUBRICANT EYE DROPS

carboxymethylcellulose sodium solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55651-400
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**
WATER (UNII: 059QF0KO0R)	
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SODIUM LACTATE (UNII: TU7HW0W0QT)	

HYDROCHLORIC ACID (UNII: QTT17582CB)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55651-400-05	5 in 1 BOX	03/15/2026	
1		0.4 mL in 1 BOTTLE, DROPPER; 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	03/15/2026	

Labeler - KC Pharmaceuticals, Inc. (174450460)

Revised: 3/2026 KC Pharmaceuticals, Inc.