HEB LUBRICANT EYE DROPS- carboxymethylcellulose sodium solution/ drops
HEB

HEB Lubricant Eye Drops 30 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 this product 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.

HEB LUBRICANT EYE DROPS

carboxymethylcellulose sodium solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:37808-100
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)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37808-100-01	30 in 1 CARTON	01/11/2019	
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	01/11/2019	

Labeler - HEB (007924756)

Registrant - Unimed Pharmaceuticals, Inc. (689852052)

Establishment

Name	Address	ID/FEI	Business Operations
Unimed Pharmaceuticals, Inc.		689852052	label(37808-100) , manufacture(37808-100) , pack(37808-100)

Revised: 12/2023 HEB