WALGREENS LUBRICANT EYE DROPS PRESERVATIVE FREE-
carboxymethylcellulose sodium solution/ drops
Walgreen Company

Walgreens Lubricant Eye Drops Preservative-Free (PLD)

Active Ingredients

Carboxymethylcellulose sodium 0.5%

Purpose

Lubricant

Uses

- for the temporary relief of burning, irritation, and discomfort due to dryness of the eyes 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 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 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 the 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°-25ºC (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, **hydrochloric acid, magnesium chloride, potassium chloride, purified water, sodium chloride, **sodium hydroxide, sodium lactate,

**May contain these ingredients to adjust pH.

Questions or comments?

Call 1-888-527-4276

Walgreens Lubricant Eye Drops Preservative Free 30ct

Walgreens Lubricant Eye Drops Preservative Free 70ct

WALGREENS LUBRICANT EYE DROPS PRESERVATIVE FREE

carboxymethylcellulose sodium solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0363-8800
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
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
SODIUM LACTATE (UNII: TU7HW0W0QT)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-8800-01	30 in 1 BOX	12/01/2019	
1		0.4 mL in 1 VIAL, SINGLE-USE; Type 0: Not a Combination Product		
2	NDC:0363-8800-02	70 in 1 BOX	12/01/2019	
2		0.4 mL in 1 VIAL, SINGLE-USE; 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	12/01/2019	

Labeler - Walgreen Company (008965063)

Registrant - KC Pharmaceuticals, Inc. (174450460)

Establishment

Name	Address	ID/FEI	Business Operations
KC Pharmaceuticals, Inc.		174450460	pack(0363-8800) , label(0363-8800)

Establishment

Name	Address	ID/FEI	Business Operations
Unimed		689852052	manufacture(0363-8800) , pack(0363-8800) , label(0363-8800)

Revised: 12/2023 Walgreen Company