

EQUATE LUBRICANT EYE DROPS PRESERVATIVE-FREE- carboxymethylcellulose sodium solution/ drops
Wal-Mart Stores, Inc.

Equate 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°-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, **hydrochloric acid, magnesium chloride, potassium chloride, purified water, sodium chloride, **sodium hydroxide, sodium lactate.

**May contain these ingredients to adjust pH.

Questions or comments?

1-888-287-1915

Equate Preservative-Free Lubricant Eye Drops 25ct



EQUATE LUBRICANT EYE DROPS PRESERVATIVE-FREE

carboxymethylcellulose sodium solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49035-245
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
HYDROCHLORIC ACID (UNII: QTT17582CB)	

CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SODIUM LACTATE (UNII: TU7HW0W0QT)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

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Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49035-245-01	25 in 1 BOX	11/01/2019	
1		0.4 mL in 1 VIAL, SINGLE-USE; Type 0: Not a Combination Product		

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Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M018	11/01/2019	

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Labeler - Wal-Mart Stores, Inc. (051957769)

Registrant - KC Pharmaceuticals, Inc. (174450460)

Establishment			
Name	Address	ID/FEI	Business Operations
KC Pharmaceuticals, Inc.		174450460	pack(49035-245) , label(49035-245)

Establishment			
Name	Address	ID/FEI	Business Operations
Unimed		689852052	manufacture(49035-245)

Revised: 12/2023

Wal-Mart Stores, Inc.