

QUALITY CHOICE LUBRICANT EYE DROPS- carboxymethylcellulose sodium solution/ drops
Chain Drug Marketing Assoc., Inc.

Quality Choice Lubricant Eye Drops 30 ct (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-operation (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.

Drug Facts

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How to use:

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*This product is not manufactured or distributed by Allergan, distributor of Refresh Plus®.

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Non-irritating, preservative-free Quality Choice Lubricant Eye Drops are an excellent choice for the temporary relief of dry, scratchy, burning, and irritated eyes.

Quality Choice Lubricant Eye Drops provide soothing relief for dry irritated eyes, which can be caused by excessive heat, air conditioning, reading, medication or computer use. This special formula instantly moisturizes and relieves dry, irritated eyes with a fast-acting, long-lasting formula for sensitive eyes that has many of the same healthy qualities as natural tears.

This product comes in preservative-free, single use vials and are safe to use as often as necessary, so your eyes can feel good - anytime, anywhere, without the risk of irritation from preservatives.

TAMPER EVIDENT: DO NOT USE IF CARTON IS OPEN OR TOP OF SINGLE-USE CONTAINER IS BROKEN OR MISSING.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

SATISFACTION 100% GUARANTEED

Distributed by C.D.M.A., Inc.®
43157 W 9 Mile Rd
Novi, MI 48375
www.qualitychoice.com
Questions: 800-935-2362

QUALITY CHOICE LUBRICANT EYE DROPS

carboxymethylcellulose sodium solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63868-618
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: M410D6VV5M)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
WATER (UNII: 059QF0KO0R)	

SODIUM HYDROXIDE (UNII: 55X04QC32I)

SODIUM LACTATE (UNII: TU7HW0W0QT)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-618-30	30 in 1 CARTON	03/13/2020	
1		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	03/13/2020	

Labeler - Chain Drug Marketing Assoc., Inc. (011920774)

Registrant - K.C. Pharmaceuticals, Inc. (174450460)

Establishment

Name	Address	ID/FEI	Business Operations
Unimed Pharmaceuticals, Inc.		689852052	manufacture(63868-618) , pack(63868-618) , label(63868-618)

Revised: 12/2023

Chain Drug Marketing Assoc., Inc.