

**MEDICS CHOICE FRESH AND SOOTHE LUBRICANT EYE DROPS- polyethylene glycol 400, propylene glycol solution/ drops
KC Pharmaceuticals, Inc.**

Medic's Choice Fresh & Soothe Lubricant Eye Drops 10mL

Active ingredients

Polyethylene glycol 400 0.4%

Propylene glycol 0.3%

Purposes

Lubricant

Lubricant

Uses

- for use as a protectant against further irritation or to relieve dryness of the eye
- for the temporary relief of discomfort due to minor irritations of the eye or to exposure to wind or sun

Warnings

For external use only

Do not use

- if this product changes color or becomes cloudy
- if you are sensitive to any ingredient in this product

When using this product

- do not touch tip of container to any surface to avoid contamination
- replace cap after each use

Stop use and ask a doctor if

you experience any of the following:

- eye pain
- changes in vision
- continued redness or irritation of the eye
- condition worsens or persists for more than 72 hours

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

put 1 or 2 drops in the affected eye(s) as needed

Other information

Store at 15-25°C (59-77°F)

Inactive ingredients

boric acid, hypromellose, potassium chloride, purified water, sodium hyaluronate, sodium hydroxide

Questions or comments?

Call 1-888-527-4276

Medic's Choice Fresh & Soothe Lubricant Eye Drops 10mL



MEDICS CHOICE FRESH AND SOOTHE LUBRICANT EYE DROPS

polyethylene glycol 400, propylene glycol solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55651-999
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PROPYLENE GLYCOL (UNII: 6DC9Q167V3) (PROPYLENE GLYCOL - UNII:6DC9Q167V3)	PROPYLENE GLYCOL	0.3 g in 100 mL
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ) (POLYETHYLENE GLYCOL 400 - UNII:B697894SGQ)	POLYETHYLENE GLYCOL 400	0.4 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
BORIC ACID (UNII: R57ZHV85D4)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55651-999-01	1 in 1 BOX	09/01/2024	
1		10 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	09/01/2024	

Labeler - KC Pharmaceuticals, Inc. (174450460)

Establishment

Name	Address	ID/FEI	Business Operations
Unimed Pharmaceuticals, Inc.		689852052	pack(55651-999) , manufacture(55651-999) , label(55651-999)

Revised: 12/2025

KC Pharmaceuticals, Inc.