QUALITY CHOICE LUBRICANT EYE DROPS LONG LASTING- polyethylene glycol 400, propylene glycol solution/ drops
Chain Drug Marketing Assoc., Inc.

Quality Choice Lubricant Eye Drops Long Lasting 15mL (PLD)

Active ingredients

Polyethylene glycol 400 0.4%

Propylene glycol 0.3%

Purposes

Lubricant

Lubricant

Use

- for the temporary relief of burning and irritation due to dryness of the eye

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 feel eye pain
- changes in vision occur
- redness or irritation of the eye(s) gets worse or lasts more than 72 hours

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

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

Directions

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

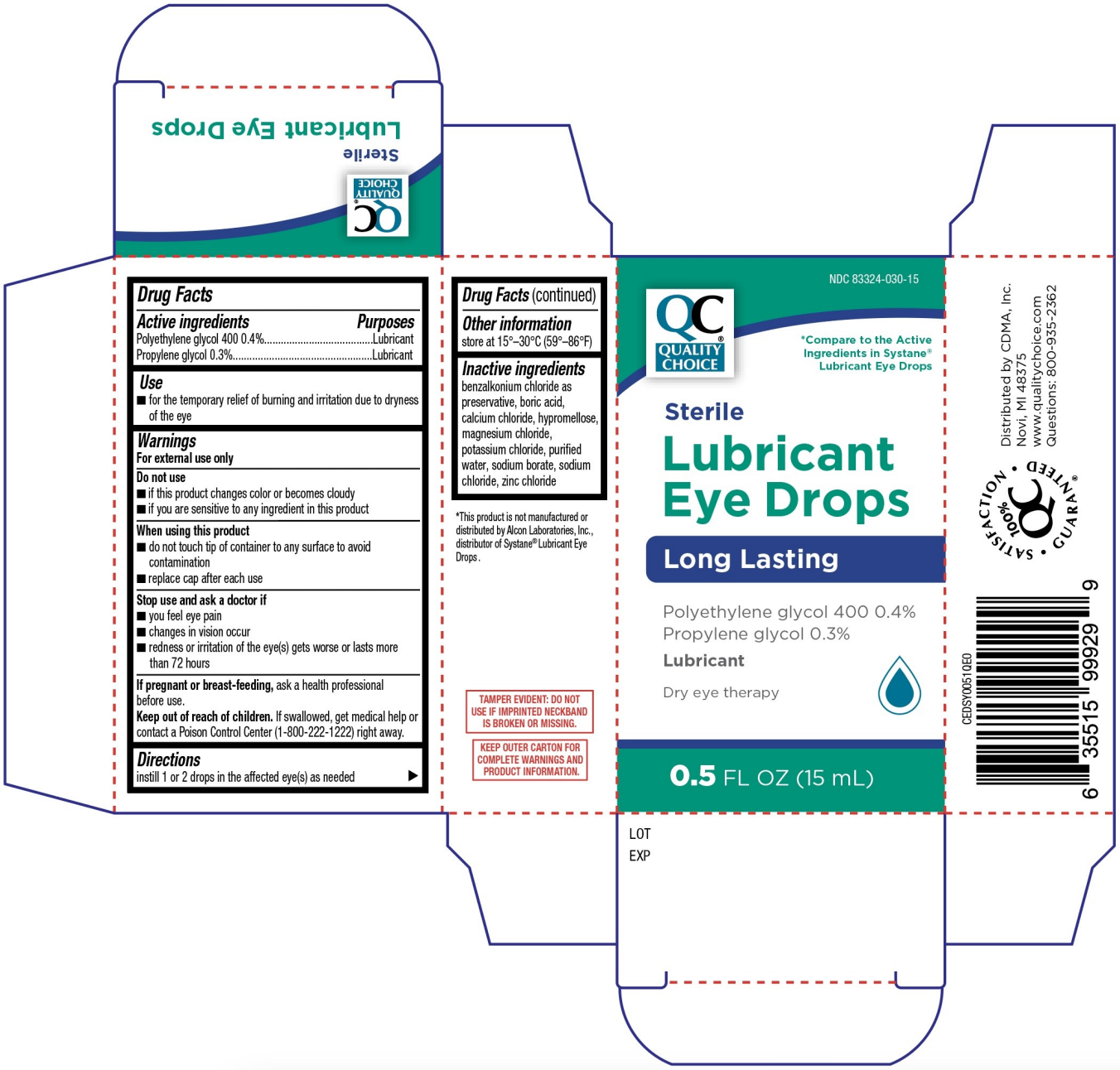
Other information

store at 15º-30ºC (59º-86°F)

Inactive ingredients

benzalkonium chloride as preservative, boric acid, calcium chloride, hypromellose, magnesium chloride, potassium chloride, purified water, sodium borate, sodium chloride, zinc chloride

Quality Choice Lubricant Eye Drops Long Lasting 15mL



QUALITY CHOICE LUBRICANT EYE DROPS LONG LASTING

polyethylene glycol 400, propylene glycol solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83324-030
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
BORIC ACID (UNII: R57ZHV85D4)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
WATER (UNII: 059QF0KO0R)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
ZINC CHLORIDE (UNII: 86Q357L16B)	
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83324-030-15	1 in 1 BOX	12/19/2023	
1		15 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	12/19/2023	

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

Registrant - KC Pharmaceuticals, Inc. (174450460)

Establishment

Name	Address	ID/FEI	Business Operations
KC Pharmaceuticals, Inc.		174450460	pack(83324-030) , manufacture(83324-030) , label(83324-030)

Revised: 12/2023 Chain Drug Marketing Assoc., Inc.