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

Quality Choice Lubricant Eye Drops High Performance 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

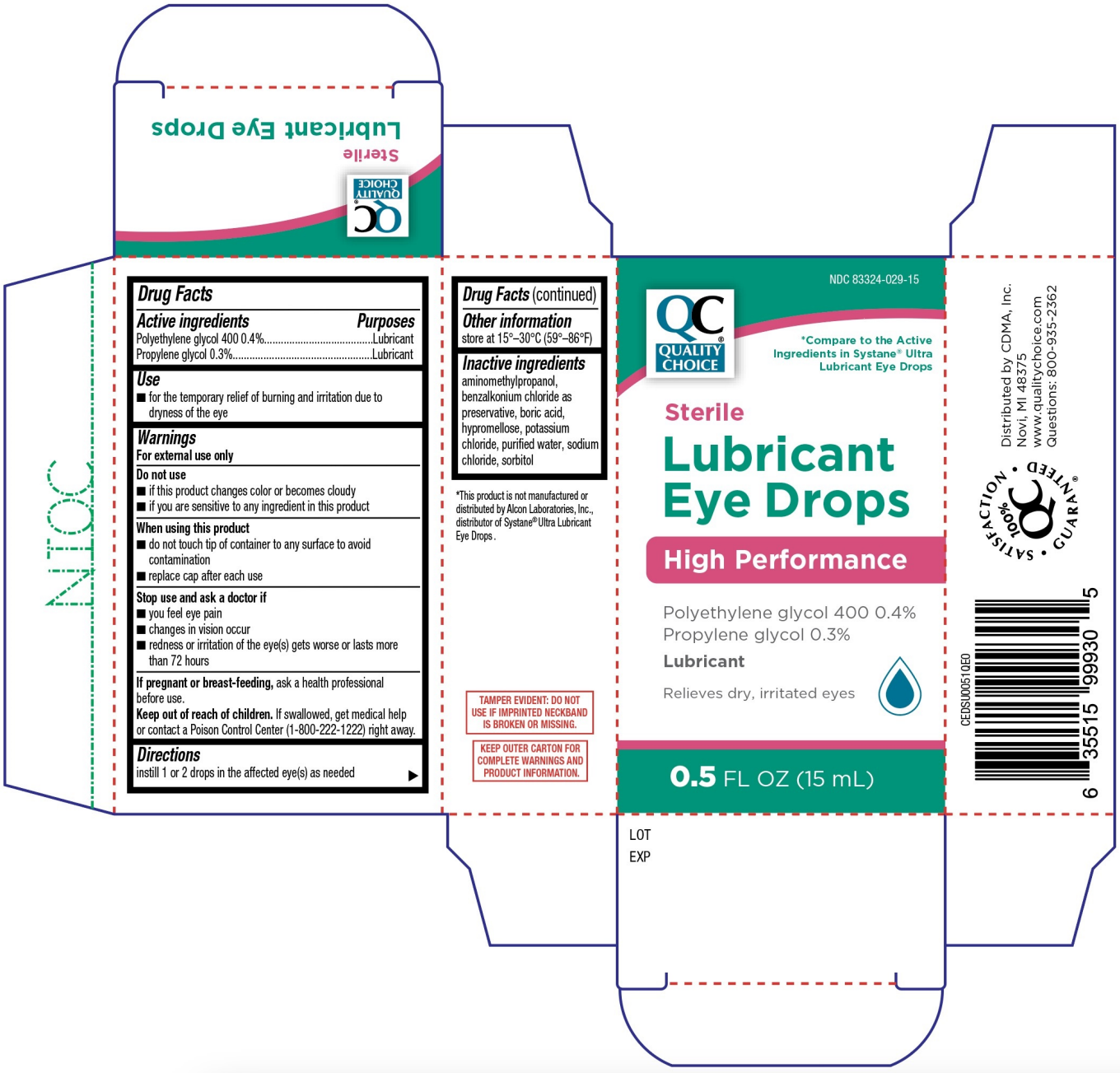
Other information

store at 15º-30ºC (59º-86ºF)

Inactive ingredients

aminomethylpropanol, benzalkonium chloride as preservative, boric acid, hypromellose, potassium chloride, purified water, sodium chloride, sorbitol;

Quality Choice Lubricant Eye Drops High Performance15mL



QUALITY CHOICE LUBRICANT EYE DROPS HIGH PERFORMANCE

polyethylene glycol 400, propylene glycol solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83324-029
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**
SORBITOL (UNII: 506T60A25R)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
BORIC ACID (UNII: R57ZHV85D4)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:83324-029-15	1 in 1 BOX	12/18/2023	
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	12/18/2023	

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

Registrant - KC Pharmaceuticals, Inc. (174450460)

Establishment

Name	Address	ID/FEI	Business Operations
KC Pharmaceuticals, Inc.		174450460	manufacture(83324-029) , pack(83324-029) , label(83324-029)

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