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

Quality Choice Lubricant Eye Drops Restorative Performance 15mL (PLD)

Active ingredient

Propylene glycol 0.6%

Purpose

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
- 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 eye pain
- changes in vision occur
- redness or irritation of the eye(s) gets worse, persists or lasts 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

- shake well before using
- instill 1 or 2 drops in the affected eye(s) as needed

Other information

- store at room temperature

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Inactive ingredients

benzalkonium chloride, boric acid, castor oil, disodium edetate hydrate, **hydrochloric acid, polyoxyethylene sorbitan monooleate, potassium chloride, purified water, sodium borate, sodium chloride, **sodium hydroxide

**May contain these ingredients to adjust pH

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propylene glycol solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83324-028
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
PROPYLENE GLYCOL (UNII: 6DC9Q167V3) (PROPYLENE GLYCOL - UNII:6DC9Q167V3)	PROPYLENE GLYCOL	0.6 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
BORIC ACID (UNII: R57ZHV85D4)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
CASTOR OIL (UNII: D5340Y2I9G)	
PEG-6 SORBITAN OLEATE (UNII: 58O7V09UCI)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
WATER (UNII: 059QF0KO0R)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:83324-028-15	1 in 1 BOX	12/12/2023	
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	12/12/2023	

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

Registrant - KC Pharmaceuticals, Inc. (174450460)

Establishment

Name	Address	ID/FEI	Business Operations
Daewoo Pharmaceuticals, Co., Ltd.		689046329	manufacture(83324-028) , pack(83324-028) , label(83324-028)

Revised: 12/2025 Chain Drug Marketing Assoc., Inc.