

**HEB LUBRICANT EYE DROPS DRY EYE THERAPY- propylene glycol solution/
drops
HEB**

HEB Lubricant Eye Drops Dry Eye Therapy 15mL and 15mL twin pack (PLD)

Active ingredient

Propylene glycol 0.6%

Purpose

Lubricant

Use

- for 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

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.

HEB Lubricant Eye Drops Dry Eye Therapy 15mL



HEB Lubricant Eye Drops Dry Eye Therapy 15mL twin pack



HEB LUBRICANT EYE DROPS DRY EYE THERAPY

propylene glycol solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:37808-650
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of	Strength
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Ingredient Name		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)				
CASTOR OIL (UNII: D5340Y2I9G)				
PEG-6 SORBITAN OLEATE (UNII: 58O7V09UCI)				
POTASSIUM CHLORIDE (UNII: 660YQ98I10)				
WATER (UNII: 059QF0KO0R)				
SODIUM BORATE (UNII: 91MBZ8H3QO)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)				
EDETATE DISODIUM (UNII: 7FLD91C86K)				
SODIUM HYDROXIDE (UNII: 55X04QC32I)				
HYDROCHLORIC ACID (UNII: QTT17582CB)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37808-650-01	1 in 1 BOX	09/12/2020	
1		15 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
2	NDC:37808-650-02	2 in 1 BOX	09/12/2020	
2		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	09/12/2020		

Labeler - HEB (007924756)

Registrant - KC Pharmaceuticals, Inc. (174450460)

Establishment

Name	Address	ID/FEI	Business Operations
Daewoo Pharmaceutical Co., Ltd.		689046329	manufacture(37808-650) , pack(37808-650) , label(37808-650)