EXCHANGE SELECT HIGH PERFORMANCE LUBRICANT EYE DROPS- polyethylene glycol 400, propylene glycol solution/ drops
Army and Air Force Exchange Service

Exchange Select High Performance Lubricant Eye Drops 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 sensative to any ingredients 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 15°-30°C (59°-86°F)

Inactive ingredients

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

Questions or Comments?

Call **1-888-527-4276**

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polyethylene glycol 400, propylene glycol solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55301-347
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, UNSPECIFIED - UNII:3WQ0SDW1A)	POLYETHYLENE GLYCOL 400	0.4 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
BORIC ACID (UNII: R57ZHV85D4)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SORBITOL (UNII: 506T60A25R)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55301-347-01	1 in 1 BOX	01/01/2024	
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	01/01/2024	

Labeler - Army and Air Force Exchange Service (001695568)

Registrant - KC Pharmaceuticals, Inc. (174450460)

Establishment

Name	Address	ID/FEI	Business Operations
KC Pharmaceuticals, Inc.		174450460	manufacture(55301-347) , pack(55301-347) , label(55301-347)

Revised: 1/2024 Army and Air Force Exchange Service