

TARGET LUBRICANT EYE DROPS HIGH PERFORMANCE- polyethylene glycol 400, propylene glycol solution/ drops
Target Corporation

Target Lubricant Eye Drops High Performance Twin (PLD)

Active ingredients

Polyethylene glycol 400 0.4%

Propylene glycol 0.3%

Purpose

Polyethylene glycol 400..... Lubricant

Propylene glycol..... Lubricant

Uses

- 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 the 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

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Instill 1 or 2 drops in the affected eye(s) as needed
- Children under 6 years of age: ask a doctor

Other information

- RETAIN THIS CARTON FOR FUTURE REFERENCE
- Store at room temperature

□Inactive ingredients

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



TARGET LUBRICANT EYE DROPS HIGH PERFORMANCE

polyethylene glycol 400, propylene glycol solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-110
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	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)				
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)				
BORIC ACID (UNII: R57ZHV85D4)				
HYPROMELLOSES (UNII: 3NXW29V3WO)				
POTASSIUM CHLORIDE (UNII: 660YQ98I10)				
WATER (UNII: 059QF0KO0R)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
SORBITOL (UNII: 506T60A25R)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-110-01	2 in 1 BOX	02/12/2019	
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	02/12/2019	

Labeler - Target Corporation (006961700)

Registrant - KC Pharmaceuticals, Inc. (174450460)

Establishment			
Name	Address	ID/FEI	Business Operations
KC Pharmaceuticals, Inc.		174450460	manufacture(11673-110) , pack(11673-110) , label(11673-110)

Revised: 12/2023

Target Corporation