FAMILY CARE ULTRA LUBRICANT- polyethylene glycol 400, and propylene glycol solution/ drops
United Exchange Corp.

Family Care Ultra Lubricant Eye Drops ZDP

Drug Facts

Active ingredients Purpose

Polyethylene Glycol 400 0.4%..... Eye lubricant

Propylene Glycol 0.3%.....Eye lubricant

Uses
- for the temporary relief of burning and irritation due to:
- dryness of the eyes
- exposure to wind or sun

Warnings

For external use only

Do not use
- if this product changes color or becomes cloudy
- if you are sensitive to any ingredient is this product

When using this product
- remove contact lenses before use
- replace cap after each use
- do not touch tip of bottle to any surface to avoid contamination

Stop use and ask a doctor if
- you experience eye pain and/or vision change
- you continue to have eye redness and/or irritation
- irritation, burning persists or worsens for 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
- shake well before using
- instill 1 or 2 drops in the affected eye(s) as needed
- children under 6 years of age: ask a doctor

Inactive ingredients

benzalkonium chloride, boric acid, hypromellose, hydrochloric acid, potassium chloride, purified water, sodium chloride, sodium hydroxide

Distributed by:

United Exchange Corp,

Cypress, CA 90630 USA

1-888-645-8204

Made in South Korea

FAMILY CARE ULTRA LUBRICANT

polyethylene glycol 400, and propylene glycol solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:65923-588
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	4 mg in 1 mL
PROPYLENE GLYCOL (UNII: 6DC9Q167V3) (PROPYLENE GLYCOL - UNII:6DC9Q167V3)	PROPYLENE GLYCOL	3 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
WATER (UNII: 059QF0KO0R)	
BORIC ACID (UNII: R57ZHV85D4)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65923-588-15	1 in 1 CARTON	07/22/2024	
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	07/22/2024	

Labeler - United Exchange Corp. (840130579)

Revised: 7/2024 United Exchange Corp.