FAMILY CARE LUBRICANT EYE- polyethylene glycol 400, and propylene glycol solution/drops

United Exchange Corp.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Active ingredients

Purpose

 Polyethylene glycol 400 0.4%.....

 Eye lubricant

Propylene glycol 0.3%.....Eye lubricant

Use

• Ifor the temporary relief of burning and irritation due to dryness of the eye

• Warnings

For external use only

Do not use

- Iif this solution changes color or becomes cloudy
- if you are sensitive to any ingredient in this product

When using this product

- Ido not touch tip of container to any surface to avoid contamination
- replace cap after each use
- remove contact lense before using

Stop use and ask a doctor if

- Iyou feel eye pain
- changes in vision occur
- redness or irritation of the eye(s) lasts
- condition worsens or lasts more than 72 hours

IIf pregnant or breast-feeding, Task 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

- Ishake well before using
- instill 1 or 2 drops in the affected eye(s) as needed
- children under 6 years of age: ask a doctor

Other information

• Istore at room temperature 20-25°C (68-77°F)

Inactive ingredients

boric acid, calcium chloride, chlorhexidine gluconate, hydrochloric acid, hypromellose, magnesium chloride, potassium chloride, purified water,

sodium chloride, sodium hydroxide, zinc chloride []

Distributed by:

United Exchange Corp.

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www.ueccorp.com

Made in Korea



FAMILY CARE LUBRICANT EYE

polyethylene glycol 400, and propylene glycol solution/ drops

Product Information

| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:65923-580 |
|-------------------------|----------------|--------------------|---------------|
| 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 | |
| BORIC ACID (UNII: R57ZHV85D4) | | |
| CALCIUM CHLO RIDE (UNII: M4I0 D6 VV5M) | | |
| CHLORHEXIDINE GLUCONATE (UNII: MOR84MUD8E) | | |
| HYDRO CHLO RIC ACID (UNII: QTT17582CB) | | |
| HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO) | | |
| MAGNESIUM CHLO RIDE (UNII: 02F3473H9O) | | |
| POTASSIUM CHLORIDE (UNII: 660 YQ98 I10) | | |
| WATER (UNII: 059QF0KO0R) | | |
| SODIUM CHLORIDE (UNII: 451W47IQ8X) | | |
| SODIUM HYDRO XIDE (UNII: 55X04QC32I) | | |
| ZINC CHLORIDE (UNII: 86Q357L16B) | | |

| F | Packaging | | | | |
|---|----------------------|---|-------------------------|-----------------------|--|
| # | Item Code | Package Description | Marketing Start Date | Marketing End Date | |
| 1 | NDC:65923-580- 15 | 1 in 1 CARTON | 05/23/2017 | | |
| 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 final | part349 | 05/23/2017 | | |
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Labeler - United Exchange Corp. (840130579)

Revised: 5/2017 United Exchange Corp.