

**WALGREENS MULTI-SYMPTOM RELIEF EYE DROPS- glycerin, hypromellose, polyethylene glycol 400, tetrahydrozoline hcl, zinc sulfate solution/ drops Walgreen Co.**

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**Walgreens Multi-Symptom Relief Eye Drops (PLD)**

***Active ingredients***

Glycerin ....0.2%

Hypromellose.....0.36%

Polyethylene glycol 400.....1%

Tetrahydrozoline HCl.....0.05%

Zinc sulfate 0.25%

***Purposes***

Glycerin ....Lubricant

Hypromellose.....Lubricant

Polyethylene glycol 400.....Lubricant

Tetrahydrozoline HCl.....Redness reliever

Zinc sulfate.....Astringent

***Uses***

- for temporary relief of discomfort and redness of the eye due to minor eye irritations
- relieves dryness of the eye
- for temporary relief of burning and irritation due to exposure to wind or sun
- for protection against further irritation

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***Warnings***

**For external use only**

**Ask a doctor before use if you have narrow angle glaucoma**

**When using this product**

- pupils may become enlarged temporarily
- overuse may cause more eye redness
- remove contact lenses before using
- do not touch tip of container to any surface to avoid contamination
- replace cap after using
- do not use if this solution changes color or becomes cloudy

**Stop use and ask a doctor if**

- you feel eye pain
- changes in vision occur
- redness or irritation of the eye lasts
- condition worsens 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 to 2 drops in the affected eye(s) up to 4 times daily
- children under 6 years of age: ask a doctor

***Other information***

- some users may experience a brief tingling sensation
- store at 20°-25°C (68°-77°F)

***Inactive ingredients***

- benzalkonium chloride, boric acid, edetate disodium, purified water, sodium chloride, sodium citrate

***Questions or comments?***

1-888-527-4276



## WALGREENS MULTI-SYMPATOM RELIEF EYE DROPS

glycerin, hypromellose, polyethylene glycol 400, tetrahydrozoline hcl, zinc sulfate solution/ drops

### Product Information

|                                |                |                           |               |
|--------------------------------|----------------|---------------------------|---------------|
| <b>Product Type</b>            | HUMAN OTC DRUG | <b>Item Code (Source)</b> | NDC:0363-1462 |
| <b>Route of Administration</b> | OPHTHALMIC     |                           |               |

### Active Ingredient/Active Moiety

| Ingredient Name   | Basis of Strength              | Strength         |
|---|--------------------------------|------------------|
| <b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO) (HYPROMELLOSE, UNSPECIFIED - UNII:3NXW29V3WO) | HYPROMELLOSE, UNSPECIFIED      | 0.36 g in 100 mL |
| <b>POLYETHYLENE GLYCOL 400</b> (UNII: B697894SGQ) (POLYETHYLENE GLYCOL 400 - UNII:B697894SGQ)     | POLYETHYLENE GLYCOL 400        | 1 g in 100 mL    |
| <b>ZINC SULFATE</b> (UNII: 89DS0H96TB) (ZINC CATION - UNII:13S1S8SF37)                            | ZINC SULFATE                   | 0.25 g in 100 mL |
| <b>GLYCERIN</b> (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)                                   | GLYCERIN                       | 0.2 g in 100 mL  |
| <b>TETRAHYDROZOLINE HYDROCHLORIDE</b> (UNII: 0YZT43HS7D) (TETRAHYDROZOLINE - UNII:S9U025Y077)     | TETRAHYDROZOLINE HYDROCHLORIDE | 0.05 g in 100 mL |

## Inactive Ingredients

| Ingredient Name                                 | Strength |
|---|----------|
| <b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7) |          |
| <b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)       |          |
| <b>SODIUM CITRATE</b> (UNII: 1Q73Q2JULR)        |          |
| <b>BORIC ACID</b> (UNII: R57ZHV85D4)            |          |
| <b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)      |          |
| <b>WATER</b> (UNII: 059QF0KO0R)                 |          |

## Packaging

| # | Item Code        | Package Description   | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:0363-1462-01 | 1 in 1 CARTON   | 03/06/2020           |                    |
| 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                                     | 03/06/2020           |                    |

**Labeler** - Walgreen Co. (008965063)

**Registrant** - K.C. Pharmaceuticals, Inc. (174450460)

## Establishment

| Name                       | Address | ID/FEI    | Business Operations   |
|----------------------------|---------|-----------|---|
| K.C. Pharmaceuticals, Inc. |         | 174450460 | manufacture(0363-1462) , pack(0363-1462) , label(0363-1462) |

Revised: 12/2023

Walgreen Co.