**TOPCARE HEALTH ULTRA ITCH RELIEF ASTRINGENT EYE DROPS-**
**tetrahydrozoline hcl, zinc sulfate solution**
**Topco Associates LLC**

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**Topcare Ultra Itch Relief Astringent (PLD)**

***Active ingredients***

Tetrahydrozoline HCl 0.05%

Zinc sulfate 0.25%

***Purposes***

Tetrahydrozoline HCl.....Redness reliever

Zinc sulfate.....Astringent

***Use***

- for temporary relief of discomfort and redness of the eye due to minor eye irritations

***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
- to avoid contamination, do not touch tip of container to any surface. Replace cap after using.
- if solution changes color or becomes cloudy, do not use
- overuse may produce increased redness of the eye
- remove contact lens before using

**Stop use and ask a doctor if** you experience

- eye pain
- changes in vision
- continued redness or irritation of the eye, or if the condition worsens or persists 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***

Instill 1 to 2 drops in the affected eye(s) up to 4 times daily

***Other information***

store at 15º-30ºC (59º-86ºF)

***Inactive ingredients***

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

sodium citrate


# TOPCARE HEALTH ULTRA ITCH RELIEF ASTRINGENT EYE DROPS

tetrahydrozoline hcl, zinc sulfate solution

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:36800-855 |
| **Route of Administration** | OPHTHALMIC | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **TETRAHYDROZOLINE HYDROCHLORIDE** (UNII: 0YZT43HS7D) (TETRAHYDROZOLINE - UNII:S9U025Y077) | TETRAHYDROZOLINE HYDROCHLORIDE | 0.05 g in 100 mL |
| **ZINC SULFATE** (UNII: 89DS0H96TB) (ZINC CATION - UNII:13S1S8SF37) | ZINC SULFATE | 0.25 g in 100 mL |

| Inactive Ingredients |
|---|

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

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:36800-855-01 | 1 in 1 CARTON | 02/01/2020 | |
| **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 | 02/01/2020 | |

## Labeler - Topco Associates LLC (006935977)

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

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| K.C. Pharmaceuticals, Inc. | | 174450460 | manufacture(36800-855) , pack(36800-855) , label(36800-855) |

Revised: 12/2023 Topco Associates LLC