

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

-----  
**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 15°-30°C (59°-86°F)

***Inactive ingredients***

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

sodium citrate

**TopCare health™**

**Eye Drops**  
Ultra Itch Relief Astringent

**Drug Facts**

Active ingredients	Purposes
Tetrahydrozoline HCl 0.05%	Redness reliever
Zinc sulfate 0.25%	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.

**Drug Facts (continued)**

**Directions**  
instill 1 to 2 drops in the affected eye(s) up to 4 times daily

**Other information**  
store at 15°-30°C (59°-86°F)

**Inactive ingredients**  
benzalkonium chloride, boric acid, edetate disodium, purified water, sodium chloride, sodium citrate

\*This product is not manufactured or distributed by Johnson & Johnson, Healthcare Products, distributor of Visine® A.C.\*

TAMPER EVIDENT: DO NOT USE IF IMPRINTED NECKBAND IS BROKEN OR MISSING.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

Scan here for more information

**TopCare health™**

COMPARE TO VISINE® A.C.\*  
ACTIVE INGREDIENTS\*

**Ultra Itch Relief Astringent Eye Drops**

Relieves Itchy Eyes Due to Pollen, Dust & Ragweed

**Sterile 0.5 FL OZ (15 mL)**

DISTRIBUTED BY  
TOPCO ASSOCIATES LLC  
ELK GROVE VILLAGE, IL 60007  
© TOPCO KCPA1118  
QUESTIONS? 1-888-423-0139  
topcare@topco.com  
www.topcarebrand.com  
PRODUCT OF USA

QUALITY GUARANTEED

This TopCare® product is laboratory tested to guarantee its highest quality. Your total satisfaction is guaranteed.

0 36800 03640 6

LOT  
EXP

## TOPCARE HEALTH ULTRA ITCH RELIEF ASTRINGENT EYE DROPS

tetrahydrozoline hcl, zinc sulfate solution

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:36800-855
<b>Route of Administration</b>	OPHTHALMIC		

### Active Ingredient/Active Moiety

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

### Inactive Ingredients

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