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
EYE DROPS AC
tetrahydrozoline hydrochloride, zinc sulfate solution/ drops

Product Information
Product Type: HUMAN OTC DRUG
Route of Administration: OPHTHALMIC
Item Code (Source): NDC:55651-024

Active Ingredient/Active Moiety

<table>
<thead>
<tr>
<th>Ingredient Name</th>
<th>Basis of Strength</th>
<th>Strength</th>
</tr>
</thead>
<tbody>
<tr>
<td>ZINC SULFATE (UNII: 89DS0H56TB) (ZINC CATION - UNII:13S18SF37)</td>
<td>ZINC SULFATE</td>
<td>2.5 mg in 1 mL</td>
</tr>
<tr>
<td>TETRAHYDROZOLINE HYDROCHLORIDE (UNII: 0YZT43HS7D) (TETRAHYDROZOLINE - UNII:59U025Y077)</td>
<td>TETRAHYDROZOLINE HYDROCHLORIDE</td>
<td>0.5 mg in 1 mL</td>
</tr>
</tbody>
</table>

Inactive Ingredients

<table>
<thead>
<tr>
<th>Ingredient Name</th>
<th>Strength</th>
</tr>
</thead>
<tbody>
<tr>
<td>SODIUM CHLORIDE (UNII: 451W47Q8X)</td>
<td></td>
</tr>
<tr>
<td>BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)</td>
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</tbody>
</table>
WATER (UNII: 059QF0KO0R)
BORIC ACID (UNII: R57ZHV85D4)
EDETATE DISODIUM (UNII: 7FLD91C86K)
SODIUM CITRATE (UNII: 1Q73Q2JULR)

<table>
<thead>
<tr>
<th>Packaging</th>
<th>Marketing Start Date</th>
<th>Marketing End Date</th>
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<tbody>
<tr>
<td>1 NDC:55651-024-01 1 in 1 CARTON</td>
<td>10/23/2003</td>
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<tr>
<td>1 15 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product</td>
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</table>

<table>
<thead>
<tr>
<th>Marketing Information</th>
<th>Marketing Start Date</th>
<th>Marketing End Date</th>
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<tbody>
<tr>
<td>OTC monograph final part349</td>
<td>10/23/2003</td>
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Labeler - KC Pharmaceuticals, Inc. (174450460)

Establishment

<table>
<thead>
<tr>
<th>Name</th>
<th>Address</th>
<th>ID/FEI</th>
<th>Business Operations</th>
</tr>
</thead>
<tbody>
<tr>
<td>KC Pharmaceuticals, Inc.</td>
<td>174450460</td>
<td>manufacture(55651-024) , pack(55651-024) , label(55651-024)</td>
<td></td>
</tr>
</tbody>
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Revised: 10/2019

KC Pharmaceuticals, Inc.