QUALITY CHOICE ARTIFICIAL TEARS- polyvinyl alcohol, povidone solution/ drops
Chain Drug Marketing Association, Inc.

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.

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Active ingredients
Polyvinyl alcohol 0.5%
Povidone 0.6%

Purpose
Polyvinyl alcohol: Eye Lubricant
Povidone: Eye Lubricant

Uses:
• for use as a protectant against further irritation or to relieve dryness of the eye
• for the temporary relief of discomfort due to minor irritations of the eye, or to exposure to wind or sun

Warnings
For external use only
• Do not use this product if solution changes color or becomes cloudy

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

When using this product
• to avoid contamination, do not touch tip of container to any surface
• replace cap after using. Keep container tightly closed
• remove contact lens before using

Keep out of the reach of children. If accidentally swallowed, get medical help or contact a Poison Control Center immediately.

Directions
Instill 1 or 2 drops in the affected eye(s) as needed.

Other information
• Tamper Evident. Do not use this product if imprinted neckband is missing or broken.
• RETAIN THIS CARTON FOR FUTURE REFERENCE
• Store at 15°-30°C (59°-86°F)

Inactive ingredients
benzalkonium chloride, dextrose, edetate disodium, potassium chloride, purified water, sodium bicarbonate, sodium chloride, sodium citrate, sodium phosphate dibasic, and sodium phosphate monobasic
## QUALITY CHOICE ARTIFICIAL TEARS
polyvinyl alcohol, povidone solution/ drops

### Product Information

<table>
<thead>
<tr>
<th>Product Type</th>
<th>HUMAN OTC DRUG</th>
<th>Item Code (Source)</th>
<th>NDC: 63868-223</th>
</tr>
</thead>
<tbody>
<tr>
<td>Route of Administration</td>
<td>OPHTHALMIC</td>
<td></td>
<td></td>
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</table>

### Active Ingredient/Active Moiety

<table>
<thead>
<tr>
<th>Ingredient Name</th>
<th>Basis of Strength</th>
<th>Strength</th>
</tr>
</thead>
<tbody>
<tr>
<td>POLYVINYL ALCOHOL (UNII: 532BS9J990)</td>
<td>POLYVINYL ALCOHOL</td>
<td>500 mg in 100 mL</td>
</tr>
<tr>
<td>Povidone (UNII: FZ989GH94E)</td>
<td>Povidone</td>
<td>600 mg in 100 mL</td>
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</table>

### Inactive Ingredients

<table>
<thead>
<tr>
<th>Ingredient Name</th>
<th>Strength</th>
</tr>
</thead>
<tbody>
<tr>
<td>BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)</td>
<td></td>
</tr>
</tbody>
</table>
DEXTROSE (UNII: IY9XDZ35W2)
EDETATE DISODIUM (UNII: 7FLD91C86K)
POTASSIUM CHLORIDE (UNII: 660YQ981I0)
WATER (UNII: 059QFO0K0R)
SODIUM BICARBONATE (UNII: 8MDF5V39QO)
SODIUM CHLORIDE (UNII: 451W47IQ8X)
SODIUM CITRATE (UNII: 1Q73Q2JULR)
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)
SODIUM PHOSPHATE, MONOBASIC (UNII: 3980J1H2SW)

Packaging

<table>
<thead>
<tr>
<th>#</th>
<th>Item Code</th>
<th>Package Description</th>
<th>Marketing Start Date</th>
<th>Marketing End Date</th>
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<tbody>
<tr>
<td>1</td>
<td>NDC:63868-223-15</td>
<td>1 in 1 BOX</td>
<td>07/26/2019</td>
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<tr>
<td>1</td>
<td></td>
<td>15 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product</td>
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Marketing Information

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<tr>
<th>Marketing Category</th>
<th>Application Number or Monograph Citation</th>
<th>Marketing Start Date</th>
<th>Marketing End Date</th>
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<tr>
<td>OTC monograph final</td>
<td>part349</td>
<td>07/26/2019</td>
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Labeler - Chain Drug Marketing Association, Inc. (011920774)

Registrant - KC Pharmaceuticals, Inc. (174450460)

Establishment

<table>
<thead>
<tr>
<th>Name</th>
<th>Address</th>
<th>ID/FEI</th>
<th>Business Operations</th>
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<tbody>
<tr>
<td>KC Pharmaceuticals, Inc.</td>
<td>174450460</td>
<td>manufacture(63868-223) , label(63868-223) , pack(63868-223)</td>
<td></td>
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</tbody>
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Revised: 7/2019