See Clean-One Eye Drops

Active ingredient
Boric acid (1.546mg/1ml)

Purpose
Lubricant

Uses
For the temporarily relief of
- burning and irritation due to dryness of the eye
- redness of the eye due to minor eye irritation

Warnings
For external use only
Do not use
- if this product changes color or becomes cloudy
- if you are sensitive to any ingredient in this product
When using this product
- do not touch tip of container to any surface to avoid contamination
- remove contact lenses before using
- replace cap after use
Stop use and ask a doctor if
- you experience eye pain, changes in vision, continued redness or irritation of the eye
- conditions worsen

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 1-3 drops in the affected eye(s) 3-6 times a day

Inactive ingredients
Sulfamethoxazole, dipotassium glycyrrhizinate, Chlorpheniramine Maleate, Aminocaproic Acid,
**SEE CLEAN-ONE EYE DROPS**
boric acid liquid

### Product Information

<table>
<thead>
<tr>
<th>Product Type</th>
<th>Item Code (Source)</th>
<th>NDC: 58354-105</th>
</tr>
</thead>
<tbody>
<tr>
<td>HUMAN OTC DRUG</td>
<td></td>
<td></td>
</tr>
<tr>
<td>OPHTHALMIC</td>
<td></td>
<td></td>
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</tbody>
</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>BORIC ACID</td>
<td>BORIC ACID</td>
<td>21.644 mg in 14 mL</td>
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</table>

### Inactive Ingredients

<table>
<thead>
<tr>
<th>Ingredient Name</th>
<th>Strength</th>
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<tbody>
<tr>
<td>SODIUM BORATE ANHYDROUS</td>
<td></td>
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</tbody>
</table>

**Product Type**
HUMAN OTC DRUG

**Route of Administration**
OPHTHALMIC
GLYCRRHIZINATE DIPO TASSIUM (UNII: CA2Y0FE3FX)
EDE rATE DISODIUM (UNII: 7FLD9 IC86K)
WATER (UNII: 059QF0KO0R)
CHLORPHENIRAMINE MALE rA TE (UNII: V1Q0O9QJ9Z)
AMINO CAPROIC ACID (UNII: U6F3787206)
SODIUM HYDROXIDE (UNII: 55X04QC32I)
SULFAMETHOXAZOLE (UNII: JE42381TNV)

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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</thead>
<tbody>
<tr>
<td>1</td>
<td>NDC:58354-105-02</td>
<td>1 in 1 BOX</td>
<td>12/29/2017</td>
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</tr>
<tr>
<td>1</td>
<td>NDC:58354-105-01</td>
<td>14 mL in 1 BOTTLE; Type 0: Not a Combination Product</td>
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Marketing Information

<table>
<thead>
<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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<tbody>
<tr>
<td>unapproved drug other</td>
<td></td>
<td>12/29/2017</td>
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Labeler - Cho-A Pharm.Co.,Ltd. (688056831)

Registrant - Cho-A Pharm.Co.,Ltd. (688056831)

Establishment

<table>
<thead>
<tr>
<th>Name</th>
<th>Address</th>
<th>ID/FEI</th>
<th>Business Operations</th>
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</thead>
<tbody>
<tr>
<td>Cho-A Pharm.Co.,Ltd.</td>
<td>688056831</td>
<td>manufacture(58354-105)</td>
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</table>

Revised: 2/2019