Drug Facts

Active ingredient
Ketotifen 0.025%
(equivalent to ketotifen fumarate 0.035%)

Purpose
Antihistamine

Uses
For the temporary relief of itchy eyes due to ragweed, pollen, grass, animal hair and dander.

Warnings
For external use only
Do not use

• if you are sensitive to any ingredient in this product
• if solution changes color or becomes cloudy
• to treat contact lens related irritation

When using this product
• remove contact lenses before use
• wait at least 10 minutes before re-inserting contact lenses after use
• do not touch tip of container to any surface to avoid contamination
• replace cap after each use

Stop use and ask a doctor if you experience any of the following:
• eye pain
• changes in vision
• redness of the eyes
• itching that worsens or lasts for more than 72 hours

Keep Out of Reach of Children
If swallowed, get medical help or contact a Poison Control Center right away.

Directions
Adults and children 3 years or older: put 1 drop in the affected eye(s) twice daily, every 8-12 hours, no more than twice per day.
Children under 3 years of age: consult a doctor
Other Information
Store at 4-25°C (39-77°F)

Inactive Ingredients
benzalkonium chloride 0.01%, glycerin, hydrochloric acid and/or sodium hydroxide, water for injection

Package/Label Principal Display Panel
NDC 49035-231-10
equate™

Compare to Zaditor® Active ingredient*
Eye Itch Relief
ketotifen fumarate
ophthalmic solution 0.035%
ANTIHISTAMINE EYE DROPS

• Works in minutes
• Original prescription strength
• For ages 3 years and older
• 60 day supply

UP TO 12 HOURS
2 x 10 mL BOTTLES
(STERILE 0.34 FL OZ EACH)
**EQUATE EYE ITCH RELIEF**
ketotifen fumarate solution

### Product Information

<table>
<thead>
<tr>
<th>Product Type</th>
<th>Item Code (Source)</th>
<th>NDC:49035-231</th>
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</thead>
<tbody>
<tr>
<td>HUMAN OTC DRUG</td>
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<tr>
<td>OPHTHALMIC</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>KETOTIFEN FUMARATE (UNII: HBD503WOR0) (KETOTIFEN - UNII:X49220T18G)</td>
<td>KETOTIFEN</td>
<td>0.25 mg in 1 mL</td>
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</tbody>
</table>

### Inactive Ingredients

<table>
<thead>
<tr>
<th>Ingredient Name</th>
<th>Strength</th>
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</thead>
<tbody>
<tr>
<td>BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)</td>
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### 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:49035-231-10</td>
<td>1 in 1 CARTON</td>
<td>01/01/2016</td>
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<tr>
<td>2</td>
<td>NDC:49035-231-11</td>
<td>10 mL in 1 BOTTLE; Type 0: Not a Combination Product</td>
<td>07/02/2018</td>
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<td>NDC:49035-231-10</td>
<td>10 mL in 1 BOTTLE; Type 0: Not a Combination Product</td>
<td>07/02/2018</td>
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<tr>
<td>2</td>
<td>NDC:49035-231-11</td>
<td>1 in 1 CARTON</td>
<td>07/02/2018</td>
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### Marketing Information

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<tr>
<th>Marketing Category</th>
<th>Application Number or Monograph Citation</th>
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<th>Marketing End Date</th>
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<tr>
<td>NDA</td>
<td>NDA021996</td>
<td>01/01/2016</td>
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### Labeler

- Wal-Mart Stores, Inc. (051957769)

### 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>Bausch &amp; Lomb Incorporated</td>
<td>079587625</td>
<td>MANUFACTURE(49035-231) , PACK(49035-231) , LABEL(49035-231)</td>
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</tbody>
</table>

Revised: 3/2020

Wal-Mart Stores, Inc.