trimethoprim 10 mg/mL Oral Suspension

<table>
<thead>
<tr>
<th>Ingredients</th>
<th>Mfr</th>
<th>Lot #</th>
<th>Expiry Date</th>
<th>Quantity</th>
<th>Measured</th>
<th>Checked</th>
</tr>
</thead>
<tbody>
<tr>
<td>trimethoprim 100mg Tablets</td>
<td>AA pharma</td>
<td></td>
<td></td>
<td>5</td>
<td></td>
<td></td>
</tr>
<tr>
<td>ORAL MIX</td>
<td>Medisca</td>
<td></td>
<td></td>
<td>q.s.50 mL</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**Additional Information:**

**Equipment:**
- mortar and pestle
- glass stirring rod
- graduated measure

**Procedure:**

Follow your Dept. procedures for risk assessment/training/PPE/equipment/facilities/NAPRA level

1. Crush tablets in the mortar to a fine powder with a pestle, or, soak tablets in a small amount of vehicle for at least 15 minutes.
2. Add a small amount of vehicle to powder and levigate to a smooth paste with a pestle. If soaked tablets, then levigate tablets into a smooth paste with a pestle. Continue to levigate as vehicle is added in small amounts until a liquid is formed.
3. Transfer liquid contents from mortar to graduate.
4. Use a small amount of vehicle to rinse mortar and add it to graduate.
5. Use vehicle to q.s. to the final volume. Stir well.
6. Transfer to amber bottle and label.

**Quality Control:**

<table>
<thead>
<tr>
<th>Expected Product Appearance</th>
<th>Additional Notes</th>
</tr>
</thead>
<tbody>
<tr>
<td>White suspension</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Storage:</th>
<th>Room temperature</th>
</tr>
</thead>
<tbody>
<tr>
<td>Packaging:</td>
<td>Amber glass/plastic PET bottles</td>
</tr>
<tr>
<td>BUD:</td>
<td>91 days</td>
</tr>
</tbody>
</table>

**Sample Label:**

trimethoprim 10 mg/mL Oral Suspension

Lot: 
BUD:

Room Temperature 
Shake Well

Date Made/Prepared By/Checked By: _________________________________

**Reference:**


**Formulation Reviewed:** October, 2020

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