hydroCHLOROthiazide 5 mg/mL Oral Suspension

Batch No: _________________________

<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>hydroCHLOROthiazide 50 mg tablets</td>
<td>TEVA/APO</td>
<td></td>
<td></td>
<td>10</td>
<td></td>
<td></td>
</tr>
<tr>
<td>ORA-Blend</td>
<td>Perrigo</td>
<td></td>
<td></td>
<td>q.s.100 mL</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Additional Information:

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

Procedure:

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

1. Soak tablets in small amount of vehicle in the mortar for a minimum of 15 minutes on the counter.
2. Use pestle to levigate into a smooth paste. 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:

Expected Product Appearance
White to light pink suspension

Storage: Room temperature
Packaging: Amber glass/plastic PET bottles
BUD: 60 days

Sample Label:

<p>| | |</p>
<table>
<thead>
<tr>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>hydroCHLOROthiazide 5 mg/mL Oral Suspension</td>
<td>Room Temperature</td>
</tr>
<tr>
<td>Lot:</td>
<td>BUD:</td>
</tr>
<tr>
<td>Shake Well</td>
<td></td>
</tr>
</tbody>
</table>

Date Made/Prepared By/Checked By: ________________________________

Reference:

Formulation Reviewed: October, 2020

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