cLONAZEpam 0.1 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>cLONAZEpam 2 mg Tablets</td>
<td>HLR/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.200 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. Soak tablets in a small amount of vehicle for at least 60 minutes.
2. 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:

Expected Product Appearance
- White to light pink suspension

Additional Notes

Storage:
- Room temperature

Packaging:
- Amber glass/plastic PET bottles

BUD:
- 60 days

Sample Label:

<table>
<thead>
<tr>
<th>SickKids</th>
<th>cloNAZEpam 0.1 mg/mL Oral Suspension</th>
</tr>
</thead>
<tbody>
<tr>
<td>Lot:</td>
<td>BUD:</td>
</tr>
<tr>
<td>Room Temperature</td>
<td>Shake Well</td>
</tr>
</tbody>
</table>

Date Made/Prepared By/Checked By: _____________________________________________________

Reference:


Formulation Reviewed: October, 2020

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