granisetron 0.05 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>granisetron 1mg tablets</td>
<td>AA Pharma/ Roche</td>
<td></td>
<td></td>
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<tr>
<td>ORA-Blend</td>
<td>Perrigo</td>
<td></td>
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</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 small amount of vehicle in the mortar for at least 30 minutes.
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

Additional Notes

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

Sample Label:

granisetron 0.05 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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