PRODUCT: HYDROCHLOROTHIAZIDE 5 mg/mL AND SPIRONOLACTONE 5 mg/mL SUSPENSION

Date Prepared: __________________________  FINAL PRODUCT CHECKED BY: __________________________

EXPIRY DATE: __________________________

<table>
<thead>
<tr>
<th>INGREDIENT</th>
<th>MANUFACTURER</th>
<th>LOT #</th>
<th>MAN. EXPIRY DATE</th>
<th>FORMULA QUANTITY</th>
<th>QUANTITY USED</th>
<th>MFG BY</th>
<th>CHK BY</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hydrochlorothiazide 25 mg Spironolactone 25 mg tablet</td>
<td>Ora-Blend®</td>
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<td>Paddock</td>
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<tr>
<td>Ora-Blend®</td>
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<td>qs to 100 mL</td>
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</table>

EQUIPMENT

- Mortar and pestle
- Graduated cylinder

MANUFACTURING DIRECTIONS

1. Crush tablets to make a fine powder in a mortar.
2. Levigate powder with a small amount of the vehicle to make a fine paste.
3. Continue to add vehicle until product liquid enough to transfer to a graduated cylinder.
4. Rinse mortar several times with vehicle and add to product in graduated cylinder.
5. QS to final volume with vehicle.
6. Dispense in amber plastic bottles. May be refrigerated or stored at room temperature.

SAMPLE LABEL

HYDROCHLOROTHIAZIDE 5 mg/mL and SPIRONOLACTONE 5 mg/mL Suspension
Shake well. Refrigerate. Take with Food.
Date Prepared: Date Expired:

STABILITY

60 days in fridge or at room temperature.

REFERENCE(S)

- Nahata MC, Hipple TF, Pediatric Drug Formulations, 4th ed. 2000, pp 115
- Allen LV, Erickson MA. AJHP 1996, 53: 2304-9

FINAL APPEARANCE

Opaque white Suspension

Master Sheet Revision Dates: 11 Dec 97; 14 Oct 05; 16 Feb 06; 10 Oct 06  Final Approval By: KS/CMB/JH

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