PRODUCT: **AZATHIOPRINE 50 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>Azathioprine 50 mg tablet</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>20</td>
<td></td>
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<tr>
<td>Ora-Blend®</td>
<td>Paddock</td>
<td></td>
<td></td>
<td></td>
<td>qs to 20 mL</td>
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</tbody>
</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.

**FINAL APPEARANCE**
Fine suspension of yellow particles.

**SAMPLE LABEL**
AZATHIOPRINE 50 mg/mL Suspension
Shake well. Room Temperature or Refrigerate.
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 17

*Master Sheet Revision Dates: 4 Feb 98; 7 July 03; 14 Oct 05; 16 Feb 06; 5 Oct 06  Final Approval By: KS/DI/CMB*

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