Children's Hospital Of Eastern Ontario

MASTER FORMULA SHEET – NON-STERILE MANUFACTURING

PRODUCT: ALLOPURINOL 20 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>Allopurinol 100 mg tablet</td>
<td>Paddock</td>
<td></td>
<td></td>
<td>20</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ora-Blend ®</td>
<td>Paddock</td>
<td></td>
<td></td>
<td>qs to 100 mL</td>
<td></td>
<td></td>
<td></td>
</tr>
</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.

FINAL APPEARANCE

White granular suspension

SAMPLE LABEL

ALLOPURINOL 20 mg/mL Suspension
Shake well. Refrigerate.
Date Prepared: _____________________  Date Expired: _____________________

STABILITY

60 days in fridge

REFERENCE(S)

- Nahata MC, Hipple TF, Pediatric Drug Formulations, 4th ed. 2000, pp 10

Master Sheet Revision Dates: 2 February 98; 14 May 01; 14 Oct 05; 29 Sept 06  Final Approval By: KS/DICMB

The non-sterile manufacturing formulas are provided for informational purposes for qualified health care professionals. The hospital will not assume responsibility for the continued currency of the information, any errors or omissions, and/or any consequences arising from the use of the information outside of CHEO.