PRODUCT: **CLONAZEPAM 0.1 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>Clonazepam 2 mg tablet</td>
<td>Ora-Blend®</td>
<td>1</td>
<td></td>
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<tr>
<td>Ora-Blend®</td>
<td>Paddock</td>
<td></td>
<td></td>
<td>qs to 20 mL</td>
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</tbody>
</table>

**EQUIPMENT**

- Mortar and pestle
- Graduated cylinder

***May use Ora-Blend sugar free to manufacture medication. No change in stability. Ref: Calgary Regional Health Authority

**MANUFACTURING DIRECTIONS**

1. Crush tablet 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.

**SAMPLE LABEL**

CLONAZEPAM 0.1 mg/mL Suspension
Shake well. Refrigerate. Protect from Light.

Date Prepared:   Date Expired:

**STABILITY**

60 days in fridge

**REFERENCE(S)**

- Nahata MC, Hipple TF, Pediatric Drug Formulations, 4th ed. 2000, pp 33
- Anon, Compounded Drug Formulas, Alberta Children’s Hospital, Calgary Regional Health Authority, May 2000, pp 40

*Master Sheet Revision Dates: 11 Dec 97; 19 Dec 01; 14 Oct 05; 16 Feb 06; 5 Oct 06  Final Approval By: KS/DI/CMB*

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