

Children's Hospital of Eastern Ontario

MASTER FORMULA SHEET – NON-STERILE MANUFACTURING

PRODUCT: AMLODIPINE 1 mg/mL SUSPENSION

Date Prepared: _____

FINAL PRODUCT CHECKED BY: _____

EXPIRY DATE: _____

INGREDIENTS	MANUFACTURER	LOT #	MAN. EXPIRY DATE	FORMULA QUANTITY	QUANTITY USED	MFG BY	CHK BY
Amlodipine 10 mg tablet				5			
Ora-Blend®				qs to 50 ml			

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 is 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. Refrigerate.

SAMPLE LABEL

AMLODIPINE 1 mg/mL SUSPENSION

Shake well. Refrigerate.

Date Prepared:

Date Expired:

STABILITY

91 days in fridge or 56 days at room temperature.

REFERENCE(S)

- Nahata MC, Morosco RS, Hipple TF. Stability of amlodipine besylate in two liquid dosage forms. J Am Pharm Assoc (Wash). 1999 May – Jun; 39 (3): 375-7

Master Sheet Revision Dates: 11 Nov 06

Final Approval By: CM/JH

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