Children's Hospital Of Eastern Ontario

MASTER FORMULA SHEET – NON-STERILE MANUFACTURING

PRODUCT: **TRIMETHOPRIM** 10 mg/mL SUSPENSION

Date Prepared: _____ **FINAL PRODUCT CHECKED BY: _____**

EXPIRY DATE: _____

INGREDIENT	MANUFACTURER	LOT #	MAN. EXPIRY DATE	FORMULA QUANTITY	QUANTITY USED	MFG BY	CHK BY
Trimethoprim 200 mg tablet				1			
Simple Syrup				qs to 20 mL			

EQUIPMENT

- Mortar and pestle
- Graduated cylinder

SAMPLE LABEL

TRIMETHOPRIM 10 mg/mL Suspension
Shake well. Refrigerate.
Date Prepared: Date Expired:

MANUFACTURING DIRECTIONS

1. Crush tablets to make a fine powder in a mortar.
2. Levigate powder with a small amount of 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.

STABILITY

91 days in fridge.

REFERENCE(S)

- Nahata MC, Hipple TF, Pediatric Drug Formulations, 4th ed. 2000, pp 124

Master Sheet Revision Dates: *21 Dec 01* ***Final Approval By:*** *BW/CMB*