Pharmacy Services

COMPOUND EVALUATION FORM			
Compound Name: Nifedipine	Container-closure system(s): Amber Plastic Bottle		
Strength: 4 mg/mL	Preservatives: Contained in SyrSpend SF PH4 Cherry		
Dosage Form: Oral Suspension	Beyond Use Date (compound type): 90 days		
Product Description: Opaque suspension	Storage: : Refrigerate (preferred) or room temperature		
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Auxiliary Labels: Refrigerate, Shake well and BUD

Quality control procedures (ex: pH test, etc.): Visual Inspection

Ingredients:

Nifedipine Bulk Powder

SyrSpend SF PH4 Cherry Flavored

Instructions for preparation:

- 1. Due to photosensitivity, nifedipine powder should not be exposed to light for prolonged periods. Please confirm pharmacist ingredient check is readily available and compound immediately after weighing powder do NOT allow ingredients to sit exposed to light.
- 2. Weigh the required quantity of powder using an electronic scale and triturate to fine powder using a mortar and pestle.
- 3. Wet powder with a minimal amount of vehicle and levigate to form a viscous but smooth and uniform paste
- 4. Geometrically continue adding vehicle and mixing well
- 5. Transfer to a conical graduate
- 6. Rinse mortar with vehicle and continue transferring to graduate until almost final volume
- 7. QS to final volume and stir well.

Calculations:

Ingredient	QS	Quantity	Units
Nifedipine		120	mg
SyrSpend SF	X	30	mL

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8. Package final product in bottle immediately after verification.	·		
Equipment/ Supplies:			
X Mortar and Pestle	X Stirring rod	☐ Ointment slab	
X Graduated cylinder(s)	☐ Spatula	X Balance	
☐ Weigh paper	X Amber (plastic/glass) bottle	☐ Ointment jar	
☐ Counting tray	☐ Syringe	□ Funnel	
References:			
1. Geiger CM, Sorenson B, Whaley P. Stability assessment of 10 active pharmaceutical ingredients			

- compounded in SyrSpend SF. *IJPC*. 2015;19(5):420-427.
- 2. Nifedipine 4mg/mL suspension. USP <51> Antimicrobial Effectiveness Testing; ARL Bio Pharma. Lot #240806-130, ARL #1068563. Date received: 8/16/24. Date tested: 11/6/24.

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