



NITROFURANTOIN 10 mg/mL ORAL SUSPENSION

Date Prepared: ____ / ____ / 20____

LOT: NS _____

Expiry Date: ____ / ____ / 20____

Final Checked By: _____

INGREDIENTS	MFR	LOT #	EXPIRY DATE	FORMULA QUANTITY	QUANTITY USED	PREPARED	CHECKED
Nitrofurantoin 50 mg tablets	AA pharma / novo			10 tablets			
Ora-blend				QS to 50 mL			

Risk Summary:

Always follow policies & procedures for risk assessment/NAPRA level as determined by your own pharmacy department.

Equipment and PPE:

- Mortar and pestle
- Graduated cylinder
- Glass stirring rod
- Standard CHEO Level B PPE (hair bonnet, mask, +/- beard cover, clean gown and nitrile gloves)

Procedure:

Always follow departments training protocols and policies and procedures that are currently in place.

1. Place tablets in mortar and pour a small amount of vehicle over the tablets before levigating.
2. Using the pestle, levigate tablets to create a smooth paste.
3. Continue to levigate, adding vehicle in small amounts until the product is liquid enough to transfer to a graduated cylinder.
4. Rinse mortar several times with vehicle and add product to the graduated cylinder.
5. QS to final volume with vehicle.
6. Stir well and transfer to amber bottle. Label bottle appropriately.
7. Before washing, rinse equipment into medication disposal bin, and not directly into sink.

Quality Control:

- Final Appearance: Opaque, yellow suspension
- Packaging: amber plastic bottles
- Storage & BUD: 91 days refrigerated or 60 days at room temperature

Sample Label:



Children's Hospital of Eastern Ontario
Pharmacy Department

Children's Hospital of Eastern Ontario

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SHAKE WELL. REFRIGERATE or ROOM TEMPERATURE.

Quantity:

LOT:

Expiry:

Reference(s):

- Ensom MHH, Decarie D. Stability of Nitrofurantoin in extemporaneously compounded suspensions. Can J Hosp Pharm 2006; 59: 29-33.

Formulation Review:

Master Sheet Revision Dates: nov 2016; jun 2018; jun 2021.