

Children's Hospital of Eastern Ontario

MASTER FORMULA SHEET – NON-STERILE MANUFACTURING

PRODUCT: DOMPERIDONE 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 |
|--------------------------|--------------|-------|------------------|------------------|---------------|--------|--------|
| Domperidone 10 mg tablet | | | | 1 | | | |
| Ora-Blend® | Paddock | | | qs to 10 mL | | | |

EQUIPMENT

- Mortar and pestle
- Graduated cylinder

SAMPLE LABEL

DOMPERIDONE 1 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 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.
6. Dispense in amber plastic bottles. Refrigerate.

STABILITY

91 days in fridge.

REFERENCE(S)

- Ensom MHH, Decarie, D, Hamilton DP, Stability of Domperidone in Extemporaneously Compounded Suspensions, Journal of Informed Pharmacotherapy. January – March 2002, Volume 8, pp 100-104

Master Sheet Revision Dates: 20 Dec 02; 3 Oct 03; 14 Oct 05; 16 Feb 06; 2 Oct 06 **Final Approval By:** CMB/EW/CMB

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