



New Zealand standardised oral formulation batch sheet

Domperidone suspension 1 mg/mL (100 mL)

Patient's name		Storage condition	In a refrigerator	At room temperature
NHI		Temperature	2-8°C	≤ 25°C
Date compounded		Shelf life	30 days	30 days
Batch number		Recommended storage	See note below	<input checked="" type="checkbox"/>

Ingredients required and formula:

Ingredient	Supplier	Batch number	Expiry date	Standard formula	Quantity dispensed	Dispensed by	Checked by
Domperidone 10 mg tablets				10 tablets			
Ora-Blend®	Perrigo			to 100 mL			
OR							
Ora-Plus®	Perrigo			50 mL			
Ora-Sweet®	Perrigo			50 mL			

Ora-Blend SF® or Ora-Sweet SF® (sugar free) can be substituted? Yes

Calculations performed by

Calculations checked by (pharmacist)

Area cleared for processing by

Instructions to compound suspension:

1. **If Ora-Blend® unavailable:** Pre-mix the Ora-Plus® and Ora-Sweet® to form the diluent.
2. Crush tablets in mortar to a fine powder.
3. Add diluent (Ora-Blend® or prepared diluent) to form a smooth paste.
4. Gradually add the diluent and transfer to the final measuring flask, rinsing the mortar well. Mix well.
5. Make up to the final volume and mix well.
6. Label and package product appropriately.

Labels

Attach product label and auxiliary labels:

Label checked by



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Final yield	_____ mL	Checked by	
Final appearance of product	Pale white to pink suspension.		
Container	Amber plastic mission bottle.		
Compounded by	<i>Name:</i>	<i>Signature:</i>	<i>Date:</i>
Final check and product release	<i>Name:</i>	<i>Signature:</i>	<i>Date:</i>
Special instructions: <ol style="list-style-type: none">1. Shake well before use.2. Store at room temperature.3. Solubility is only 50 microgram/mL so the domperidone will be in suspension. Storage at room temperature is preferred as if stored in a refrigerator the preparation will be more viscous and harder to re-suspend.			
References: <ol style="list-style-type: none">1. Allen LV, Jr. <i>Int J Pharm Compd</i>, 2006.2. Ensom MH, Decarie D, Hamilton D. <i>J Inform Pharmacother</i>, 2002; 8: 100-104.			
Prepared by	New Zealand Compounding Working Group		
Approved by	Medication Safety Expert Advisory Group of Health Quality and Safety Commission NZ		
Hosted by	Pharmaceutical Society of New Zealand Standard Formulations in the Practice Support section of the PSNZ website		
Version number	2.0	Version approval date	16/10/18
Document review due	01/11/20		

Disclaimer: this batch sheet has been designed to provide guidance and standardised formulations for New Zealand pharmacists. Please read and familiarise yourself with the general guidance for compounding oral liquids (available on the PSNZ web site) before use. Information was considered accurate at the time of publication.