



## New Zealand standardised oral formulation batch sheet

### Ursodeoxycholic acid suspension 50 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	<input checked="" type="checkbox"/>	May be stored at room temperature if this is more practical and temperatures greater than 25°C are avoided

#### Ingredients required and formula:

Ingredient	Supplier	Batch number	Expiry date	Standard formula	Quantity dispensed	Dispensed by	Checked by
Ursodeoxycholic acid 250 mg capsules				20 capsules			
Ora-Blend®	Perrigo			to 100 mL			
<b>OR</b>							
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. Empty the contents of the capsules in a mortar.
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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### Ursodeoxycholic acid suspension 50 mg/mL (100 mL)

Final yield	_____ mL	Checked by	
Final appearance of product	Pale white 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: 1. Shake well before use. 2. Store in a refrigerator.			
References: 1. Johnson CE, Streetman DD. <i>Am J Health Syst Pharm</i> , 2002; <b>59</b> (4): 361-3. 2. Allen LV, Jr. <i>Secundum Artem</i> , 2009; <b>14</b> (3).			
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 <a href="#">Standard Formulations in the Practice Support section of the PSNZ website</a>		
Version number	3.0	Version approval date	17/07/19
Document review due	01/08/21		

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.