

## New Zealand standardised oral formulation batch sheet

Terbinafine suspension 25 mg/mL (100 mL)											
Patient's name			Storage cond	Storage condition		In a refrigerator		At room temperature			
NHI	NHI		Temperature		2-8°C			≤ 25°C			
Date compounded		Shelf life		30 days			30 days				
Date compounded	Date compounded		Recommended storage				ı	May be stored at room temperature if this is more practical and temperatures greater than 25°C are avoided			
Ingredients required	and formula:						ı.				
Ingredient	Supplier	Batch number	Expiry date	Star	ndard nula			spensed	Checked by		
Terbinafine 250 mg tablets				10	tablets	ablets					
Ora-Blend®	Perrigo			to 2	100 mL						
OR		1	1								
Ora-Plus®	Perrigo				50 mL						
Ora-Sweet®	Perrigo				50 mL						
Ora-Blend SF® or Ora	-Sweet SF® (sug	gar free) can	be substituted	? '	Yes						
Calculations performed by											
Calculations checked by (pharmacist)											
Area cleared for proc	essing by										
Instructions to compo	ound suspensio	n:									
1. If Ora-Blend®	® <b>unavailable</b> : P	re-mix the O	ra-Plus® and C	)ra-Sv	veet® to	form the dilu	ıent.				
<ol><li>Crush tablets</li></ol>	in mortar to a f	fine powder.									
3. Add diluent (	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 page	ckage product a	ppropriately	•								
Labels								1 . 1 1	.111 1.		
Attach product label and auxiliary labels:								Label checked by			



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Final yield		mL		Checked by					
Final appearance of product		Pale white suspension.							
Container		Amber plastic mission bottle.							
Compounded by		Name:	Sigi	nature:	Date:				
Final check and product release		Name:	Sigi	nature:	Date:				
<ol> <li>Special instructions:</li> <li>Shake well before use.</li> <li>Storage in a refrigerator.</li> </ol>									
References:  1. Abdel-Rahman SM, Nahata MC. Am J Health Syst Pharm, 1999; <b>56</b> (3): 243-245.									
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	3.0		Version approval date 17/07/19		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.