

New Zealand standardised oral formulation batch sheet

Verapamil suspension 50 mg/mL (40 mL)											
Patient's name	Patient's name		Storage condition		In a refrigerator			At room temperature			
NHI	-11		Temperature		2-8°C			≤ 25°C			
Date compounded		Shelf life		30 days			30 days				
Batch number	Batch number		Recommended storage		$\overline{\checkmark}$			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		ndard nula	' '		spensed '	Checked by		
Verapamil 80 mg tablets				25	tablets	ablets					
Ora-Blend®	Perrigo			to	40 mL						
OR											
Ora-Plus®	Perrigo				20 mL						
Ora-Sweet®	Perrigo				20 mL						
Ora-Blend SF® or Ora-Sweet SF® (sugar free) can be substituted? Yes											
Calculations perform		•									
Calculations checked											
Area cleared for prod	essing by	sing by									
Instructions to comp	ound suspensio	n:									
1. If Ora-Blend	® unavailable : P	re-mix the O	ra-Plus® and C	ra-Sv	veet® to	form the dilu	ient				
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 pa	ckage product a	ppropriately.									
Labels											
Attach product label	and auxiliary la	bels:						Label	checked by		
Attach product laber and daxmary labers.									•		



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Final yield		mL		Checked by						
Final appearance of product		Pale white suspension.								
Container		Amber plastic mission bottle.								
Compounded by		Name:	Signatu	ire:	Date:					
Final check and product release		Name:	Signatu	ire:	Date:					
Special instructions:										
 Store in a refrigerator. Shake well before use. 										
References:										
1. Allen LV, Jr., Erickson MA, 3rd. <i>Am J Health Syst Pharm</i> , 1996; 53 (19): 2304-2309.										
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.