

New Zealand standardised oral formulation batch sheet

	Enal	april susp	ension 1	mg/n	nL (1	00 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		$\overline{\checkmark}$		May be stored at room temperature if this is more practical and temperatures greater than 25°C are avoided			
Ingredients require	d and formula:		<u> </u>		•						
Ingredient	Supplier	Batch number	Expiry date	Stand	, ,		Di by	spensed	Checked by		
Enalapril 20 mg tablets				5 ta	ablets						
Ora-Blend®	Perrigo			to 10	00 mL						
OR											
Ora-Plus®	Perrigo			į	50 mL						
Ora-Sweet®	Perrigo			. !	50 mL						
Ora-Blend SF® or O	ra-Sweet SF® (su	igar free) can	be substitute	d? Y	es						
Calculations perfor											
Calculations checked by (pharmacist)											
Area cleared for processing by											
Instructions to com	pound suspensi	on:									
1. If Ora-Blen	d® unavailable:	Pre-mix the O	ra-Plus® and	Ora-Swe	eet® to	form the dilu	ient				
2. Crush table	ts 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 p	ackage product	appropriately									
Labels											
Attach product labe	el and auxiliary l	abels:						Label	checked by		



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Enalapril suspension 1 mg/mL (100 mL)

Final yield		mL		Checked by					
Final appearance of product		Pink white suspension.							
Container		Amber plastic mission bottle/ amber glass.							
Compounded by		Name:	Sig	nature:	Date:				
Final check and product release		Name:	Sig	nature:	Date:				
Special instructions: 1. Shake well before use. 2. Store in a refrigerator.									
References: 1. Nahata MC, Morosco RS, Hipple TF. Am J Health Syst Pharm, 1998; 55 (11): 1155-7.									
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								
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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.