

## New Zealand standardised oral formulation batch sheet

Clonidine suspension 10 microgram/mL (90 mL)											
Patient's name	atient'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		<b>V</b>			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	Expiry	/ Star		ndard Quantity		pensed	Checked		
		number	date	forn	mula dispensed		by		by		
Clonidine 150 microgram tablets				6	tablets						
Ora-Blend®	Perrigo			to	90 mL						
OR	<del>-</del>										
Ora-Plus®	Perrigo				45 mL						
Ora-Sweet®	Perrigo				45 mL						
Ora-Blend SF® or Ora	be substituted	? ,	Yes								
Calculations performe											
Calculations checked by (pharmacist)											
Area cleared for proc											
Instructions to compound suspension:											
•	•		ra-Plus® and (	ارکاری اکتاح	weet® to	form the dil	uent.				
<ol> <li>If Ora-Blend® unavailable: Pre-mix the Ora-Plus® and Ora-Sweet® to form the diluent.</li> <li>Crush tablets in mortar to a fine powder.</li> </ol>											
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 lal	bels:						Label	checked by		



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Final yield		mL		Checked by					
Final appearance of product		Milky white suspension.							
Container		Amber plastic mission bottle/ amber glass.							
Compounded by		Name:	Sigi	nature:	Date:				
Final check and product release		Name:	Sigi	nature:	Date:				
Special instructions:  1. Shake well before use. 2. Store in a refrigerator.									
References:  1. Ma C, Decarie D, Ensom MHH. Am J Health Syst Pharm, 2014; 71(8): 657-61.									
Prepared by	New Zealand Compounding Working Group								
Approved by	National Medication Safety Advisory Group of the Health Quality & Safety Commission New Zealand								
Hosted by	Pharmaceutical Society of New Zealand Standard Formulations in the Practice Support section of the PSNZ website								
Version number	1.0		Version approval date 18/03/20		18/03/20				
Document review due	01/04/22								

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.