

New Zealand standardised oral formulation batch sheet

Metoprolol suspension 10 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		$\overline{\mathbf{V}}$		May be stored at room temperature if this is more practical and temperatures greater than 25°C are avoided			
Ingredients required	and formula:		Ü								
Ingredient	Supplier	Batch number	Expiry date	Star forn	•		Dis	spensed	Checked by		
Metoprolol 100 mg tablets (tartrate)				10	tablets						
Ora-Blend®	Perrigo			to	100 mL						
OR											
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 proc											
Instructions to compo	ound suspensio	n:									
1. If Ora-Blend®	unavailable:	Pre-mix the O	ra-Plus® and (اکاra-S	weet® to	form the dil	uent	t.			
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 page	ckage product a	ppropriately.									
Labels								1			
Attach product label						Label	checked by				



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Final yield		mL		Checked by					
Final appearance of product									
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. Allen LV, Jr., Erickson MA, 3rd. Am J Health Syst Pharm, 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.