

New Zealand standardised oral formulation batch sheet

Lorazepam suspension 1 mg/1 mL (60 mL)

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Patient's name			Storage condi	tion	In a	refrigerator		At room	temperature
NHI			Temperature		2-8°C			≤ 25°C	
Date compounded	Date compounded		Shelf life		30 days			30 days	
Batch number			Recommended storage					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	Quantity dispensed	Dis by	spensed	Checked by
Lorazepam 1 mg tablets				60	tablets				
Ora-Blend®	Perrigo			to	60 mL				
OR				ļ			1		
Ora-Plus®	Perrigo				30 mL				
Ora-Sweet®	Perrigo				30 mL				
Ora-Blend SF® or Ora		gar free) can l	be substituted	?	Yes				
Calculations perform	ed hy	<u> </u>							
Calculations perform	-								
Calculations checked by (pharmacist)									
Area cleared for processing by									
Instructions to comp	ound suspensio	n:							
1. If Ora-Blend	1. If Ora-Blend® unavailable: Pre-mix the Ora-Plus® and Ora-Sweet® to form the diluent.								
	2. Crush tablets in mortar to a fine powder.								
3. Gradually add the diluent (Ora-Blend® or prepared diluent) and transfer to the final measuring flask, rinsing the mortar well. Mix well.									
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Labels									
Attach product label	and auxiliary la	bels:						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:	Sig	nature:	Date:				
Final check and product release		Name:	Sig	nature:	Date:				
 Special instructions: Lorazepam easily dissolves in the mouth. If whole tablet doses are required there is generally no need to put the lorazepam into a suspension. A suspension will be required if using enteric tube administration. Shake well before use. Store in a refrigerator. 									
References: 1. Lee WME, Lugo RA 2. Nahata MC, Pai V.			•		Whitney Books.				
Prepared by	New Zealand Compounding Working Group								
Approved by	National Medication Safety Advisory Group of the Health Quality & Safety Commission New Zealand								
Hosted by		maceutical Society of New Zealand dard Formulations in the Practice Support section of the PSNZ website							
Version number	1.1		Version a	Version approval date 01/02/2023					
Document review due	01/02/2025)25							

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.