

New Zealand standardised oral formulation batch sheet

		Azathio	prine sus	spension !	50 n	ng/m	L (50 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				May be stored at room temperature if this is more practical and temperatures greater than 25°C are avoided		
Ingredients	required a	and formula:									
•		Supplier	Batch number	Expiry date	Stan forn	ndard nula	Quantity dispensed	Disp by	ensed	Checked by	
Azathioprin tablets	e 50 mg				50	tablets					
Ora-Blend®		Perrigo			to	50 mL					
OR	•		•	•	•			•		•	
Ora-Plus®		Perrigo				25 mL					
Ora-Sweet®		Perrigo				25 mL					
Ora-Blend SF® or Ora-Sweet SF® (sugar free) can be substituted? Yes											
Calculations performed by											
Calculations checked by (pharmacist)											
Area cleared for processing by											
Instructions	to compo	ound suspensic	n:								
-	toxic – Mu ective equ	ust use designa iipment.	ted cytotoxic	equipment ar	id pre	pare in	segregated ar	ea. ۱	Near pe	rsonal	
1. If C	1. If Ora-Blend® is unavailable: Pre-mix the Ora-Plus® and Ora-Sweet® to form the diluent.										
3. Gra	3. Gradually add the diluent (Ora-Blend® or prepared diluent) and transfer to the final measuring flask,										
	rinsing mortar well. Mix well.										
	4. Make up to final volume and mix well.5. Label and package product appropriately.										
Labels			· · · /								
Attach prod						Label	checked by				
,		,								,	



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Azathioprine suspension 50 mg/mL (50 mL)

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 re		Name:	Sig	nature:	Date:					
 Special instructions: Cytotoxic. Store in a refrigerator. Shake well before use. 										
References: 1. Allen LV, Jr., Erick 2. Allen LV, Jr., Erick		Am J Health Syst Ph Secundum Artem; 5		; 53 (16): 1944-9.						
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					
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.