

Acetazolamide suspension 25 mg/mL (100 mL)

Patient's name				tion	In a refrigerator		At room	At room temperature	
NHI				e	2-8°C		≤	≤ 25°C	
Date compounded					30 days		30 days		
Batch number				ed	V		May be stored at room temperature if this is more practical and temperatures greater than 25°C are avoided		
Ingredients required	l and formula:								
Ingredient	Supplier	Batch number	Expiry date	Stand formu		Quantity dispensed	Dispensed by	Checked by	
Acetazolamide 250 mg tablets				10 ta	ablets				
Ora-Blend [®]	Perrigo			to 10	00 mL				
OR									
Ora-Plus [®]	Perrigo			5	50 mL				
Ora-Sweet [®]	Perrigo			5	50 mL				
Ora-Blend SF [®] or Or	a-Sweet SF [®] (sug	ar free) can l	be substituted	? Ye	es				
Calculations performed by									
Calculations checked by (pharmacist)									
Area cleared for processing by									
Instructions to comp	ound suspension	n:							
1. If Ora-Blend	I [®] unavailable: F	Pre-mix the O	ra-Plus [®] and C	Dra-Swe	eet® to	form the dil	uent.		
2. Crush tablet	s in mortar to a f	fine powder.							
3. Add diluent	(Ora-Blend [®] or p	repared dilu	ent) to form a	smooth	h paste	2.			
4. Gradually ac	ld the diluent an	d transfer to	the final meas	uring fl	lask, ri	nsing the mo	rtar well. Mi	x well.	
5. Make up to	the final volume	and mix well							
6. Label and pa	ackage product a	ppropriately.							

Labels

Attach product label and auxiliary labels:

Label checked by



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Final yield		mL		Checked by					
Final appearance of product		Thick, white suspension. Cherry smell.							
Container		Amber plastic mission bottle/ amber glass.							
Compounded by		Name:	Sig	nature:	Date:				
Final check and product re		Name:	Sig	nature:	Date:				
 Special instructions: 1. Shake well before use. 2. Store in a refrigerator. 									
 References: 1. Allen LV, Jr., Erickson MA, 3rd. Secundum Artem; 5(4). 2. Allen LV, Jr., Erickson MA, 3rd. Am J Health Syst Pharm, 1996; 53(16): 1944-9. 3. Santoveña A, Suárez-González J, Martín-Rodríguez C, et al. Pharm Dev Tech, 2017; 22(2): 191-197. 									
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,		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.