

Phenobarbital sodium solution 10 mg/mL (40 mL)

Patient's name			Storage condit		on In a refrigerator		At roo	At room temperature	
NHI			Temperature 2-8°C		2-8°C	≤ 25°C			
Date compounded	ed		Shelf life		Precipitates			28 days	
Batch number		Recommended storage		There is a risk of precipitation if the solution is stored in a refrigerator			V		
Ingredients required	and formula:								
Ingredient	Supplier	Batch number	Expiry date	Star forn	ndard nula	Quantity dispensed	Dispense by	d Checked by	
Phenobarbital sodium powder				Z	400 mg				
Glycerol					4 mL				
Water [§]				to	40 mL				
Ora-Blend SF [®] or Ora	-Sweet SF [®] (sug	gar free) can l	be substituted	?	Not app	licable	1		
Calculations perform	ed by								
Calculations checked by (pharmacist)									
Area cleared for processing by									
 Wet the pow Gradually add Gradually add Make up to t Label and padd 	ound suspensio required amou der with glycero d water and trai he final volume ckage product a e phenobarbita	nt of phenob ol. nsfer to the fi and mix well ppropriately.	inal measuring	flask	x, rinsing				
Labels	,						Lak	el checked by	
Attach product label	una auxinary la	ueis:							



Phenobarbital sodium solution 10 mg/mL (40 mL)

Final yield	mL	Checked by			
Final appearance of product	Clear colourless solution.				
Container	Amber plastic mission bottle.				
Compounded by	Name: S	gnature:	Date:		
Final check and product release	Name: S	gnature:	Date:		
Special instructions:					

- 1. § Water used for compounding must be compliant with the Health and Disability Services Pharmacy Services Standard (NZS 8134.7.2010; Standards New Zealand, 2010), standard 5.19 *Starting materials shall be of a quality suitable for use in products intended for therapeutic use*. Potable water includes filtered tap water, water for irrigation or water for injection.
- 2. Store in a room temperature. There is a risk of precipitation if the solution is stored in a refrigerator.
- 3. Do not use if the colour of product changes to yellow or there are signs of precipitation.
- 4. Shake well before use.
- 5. If necessary, any unpleasant taste of the oral solution can be disguised by adding lemon or orange flavouring at the time of administration eg, ice blocks or cordial drink.

References:

1. Garg S, Svirskis D, Myftiu J, et al. J Pharm Prac Res, 2008; **38**(1): 28-31.

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					
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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.