

New Zealand standardised oral formulation batch sheet

Tramadol 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		$\overline{\checkmark}$		May be stored at room temperature if this is more practical and temperatures greater than 25°C are avoided			
Ingred	ients required	and formula:		<u> </u>		•						
Ingred		Supplier	Batch number	Expiry date	Stand	· ·		Disp by	pensed	Checked by		
Trama	dol 50 mg es				20 ca	capsules						
Ora-Bl	end SF®	Perrigo			to 2	100 mL						
OR		1	.	'			l			l		
Ora-Plus®		Perrigo				50 mL						
Ora-Sweet SF®		Perrigo				50 mL						
Ora-Bl	end® or Ora-Sv	weet® can be su		No)							
Calculations performed by					·							
Calculations checked by (pharmacist)												
Area cleared for processing by												
Instruc	ctions to comp	ound suspensio	n:									
Note:	this formula us	ses Ora-Blend <u>S</u>	<u>F</u> ® or Ora-Sw	eet <u>SF</u> ®								
1.	If Ora-Blend	SF® unavailable	e: Pre-mix the	e Ora-Plus® a	and Ora	-Sweet	SF® to form th	ne dil	uent.			
2.	2. Empty the contents of the capsules in a mortar.											
3.	3. Add diluent (Ora-Blend SF® or prepared diluent) to form a smooth paste.											
4.	4. Gradually add diluent and transfer to the final measuring flask, rinsing the mortar well. Mix well.											
5.	5. Make up to final volume and mix well.											
6. Label and package product appropriately.												
Labels												
	product lahel	and auxiliary la						Label	checked by			
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Document review due

01/08/21

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Final yield		m	L	Checked by						
Final appearance of produ	White suspension.									
Container	Amber plastic mission bottle/ amber glass.									
Compounded by		Name:	Signature:		Date:					
Final check and product re	elease	Name:	Sigi	nature:	Date:					
Special instructions: 1. Shake well before use. 2. Store in a refrigerator.										
 References: Polonini HC, Silva SL, Cunha CN, et al. <i>Pharmazie</i>, 2016; 71(4): 185-191. Wagner DS, Johnson CE, Cichon-Hensley BK, et al. <i>Am J Health-Syst Pharm</i>, 2003; 60: 1268-1270. 										
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 a	17/07/19						

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.