New Zealand standardised oral formulation batch sheet

Dipyridamole 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	See note below	☑

Ingredients required and formula:

Ingredient	Supplier	Batch number	Expiry date	Standard formula	Quantity dispensed	Dispensed by	Checked by
Dipyridamole 25 mg tablets				40 tablets			
Ora-Blend®	Perrigo			to 100 mL			
OR							
Ora-Plus®	Perrigo			50 mL			
Ora-Sweet®	Perrigo			50 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 compound suspension:

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. Add diluent (Ora-Blend® or prepared diluent) to form a smooth paste.
4. Gradually add the diluent and transfer to the final measuring flask, rinsing the mortar well. Mix well.
5. Make up to the final volume and mix well.
6. Label and package product appropriately.

Labels

Attach product label and auxiliary labels:	Label checked by

Dipyridamole suspension 10 mg/mL (100 mL)

Final yield	_____ mL	Checked by	
Final appearance of product	Orange suspension.		
Container	Amber plastic mission bottle.		
Compounded by	*Name:*	*Signature:*	*Date:*
Final check and product release	*Name:*	*Signature:*	*Date:*

Special instructions:

1. Shake well before use.
2. Store at room temperature.
3. Dipyridamole is only slightly soluble in water and most drug will be in suspension. Storage at room temperature is preferred as if stored in a refrigerator the preparation will be more viscous and harder to re-suspend.

References:

1. Allen LV, Jr., Erickson MA, 3rd. *Am J Health Syst Pharm*, 1996; **53**(18): 2179-84.
2. Allen LV, Jr., Erickson MA, 3rd. *Secundum Artem*; **6**(1).

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	2.0	Version approval date	16/10/18
Document review due	01/11/20		