



New Zealand standardised oral formulation batch sheet

Amiodarone 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	7 days	No data
Batch number		Recommended storage	<input checked="" type="checkbox"/>	-

Ingredients required and formula:

Ingredient	Supplier	Batch number	Expiry date	Standard formula	Quantity dispensed	Dispensed by	Checked by
Amiodarone 200 mg tablets				5 tablets			
Simple syrup				to 100 mL			

Ora-Blend SF[®] or Ora-Sweet SF[®] (sugar free) can be substituted? No

Calculations performed by

Calculations checked by (pharmacist)

Area cleared for processing by

Instructions to compound suspension:

Note: Visual physical stability has been established for Aratac and Cordarone-X brands of amiodarone.

1. Crush the tablets in a mortar to a fine powder and add the syrup to form a smooth paste.
2. Gradually add syrup and transfer to the final measuring flask, rinsing the mortar well. Mix well.
3. Make up to final volume with syrup and mix well.
4. Label and package product appropriately.

Labels

Attach product label and auxiliary labels:

Label checked by



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Final yield	_____ mL	Checked by	
Final appearance of product	Milky white suspension.		
Container	Amber plastic mission bottle/ amber glass.		
Compounded by	<i>Name:</i>	<i>Signature:</i>	<i>Date:</i>
Final check and product release	<i>Name:</i>	<i>Signature:</i>	<i>Date:</i>
Special instructions: 1. Shake well before use. 2. Store in a refrigerator.			
References: 1. Auckland City Hospital. Batch manufacturing order. Amiodarone 10mg in 1mL oral suspension. Auckland DHB. Original issue date 07/06/2004. Revised 20/01/2017. 2. Nahata MC. <i>Ann Pharmacother</i> , 1997; 31 (7-8): 851-852.			
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	05/11/19
Document review due	01/12/21		

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.