

TELEPHONE: 514-905-5096 FAX: 514-905-5097 technicalservices@medisca.net

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Suggested Formula	Amiodarone Hydrochloride 50 mg/mL Oral Liquid (Suspension, 50 mL)	FIN	F 003 363v4
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SUGGESTED FORMULATION

Ingredient Listing	Qty.	Unit	NDC#	Supplier	Lot Number	Expiry Date
Amiodarone Hydrochloride	2.500	g				
Glycerin, USP	2.0	mL				
Cherry Flavor	0.25	mL				
Methylcellulose (1500 CPS), USP	0.35	g				
Sodium Benzoate, NF	0.07	g				
Purified Water, USP	17.5	mL	(
Purified Water, USP	q.s. to 35.0	mL		·		
Syrup (Simple), NF	q.s. to 50.0	mL				
Citric Acid 20% Solution	As Required					
Monohydrate Potassium Citrate 20% Solution	As Required			7		

SPECIAL PREPARATORY CONSIDERATIONS

Ingredient-Specific Information		
Light Sensitive (protect from li	ight whenever possible):	Amiodarone Hydrochloride
Moisture Sensitive (protect fro	m humidity whenever possible):	Citric Acid
Hygroscopic (protect from moi	sture whenever possible):	Glycerin, Methylcellulose, Monohydrate Potassium Citrate
Suggested Preparatory Guidelines		
Non-Sterile Preparat	ion Sterile Preparation	
<u>Processing Error /</u> <u>Testing Considerations</u> :		and pH testing considerations during preparation, it is al 10 to 12% of the required quantities of ingredients.
Special Instruction:	Protective apparel, such as a lab of should always be worn.	coat, disposable gloves, eyewear and face-masks
		f very small quantities of ingredients. All calculations be verified before dispensing the final product.



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SUGGESTED PREPARATION (for 50 mL)

Weigh and / or measure the following ingredients when appropriate:

Ingredient Listing	Qty.	Unit	Multiplication factor (*):	Processing Error	Qty. to measure
Amiodarone Hydrochloride §	2.500	g			
Glycerin, USP §	2.0	mL			
Cherry Flavor	0.25	mL			
Methylcellulose (1500 CPS), USP §	0.35	g	8		
Sodium Benzoate, NF	0.07	g			
Purified Water, USP	17.5	mĹ			
Purified Water, USP	q.s. to 35.0	mL			
Syrup (Simple), NF	q.s. to 50.0	mL			
Citric Acid 20% Solution §	As Required				
Monohydrate Potassium Citrate 20% §	As Required				

- * Takes into account increased batch size conversions and density conversions, if required.
- § Weigh / measure just prior to use.



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Preparatory Instruction

1. **Powder-medium preparation:**

- A. Using direct heat, heat the Purified Water (17.5 mL plus processing error adjustments) between 90°C and 100°C.
- B. In the given order, sequentially add the following ingredients to the heated water:
 - -Sodium Benzoate
 - -Methylcellulose (1500 CPS)

Specifications: Continuously mix.

Maintain temperature between 90°C and 100°C.

End result: Homogeneous liquid-like dispersion.

Note: Add the next ingredient, once the previous one has been completely added and dissolved/dispersed.

C. Remove the mixture from the heat, and add ICE COLD Purified Water to the mixture (Step 1B) to fill to the required batch size (35.0 mL *plus* processing error adjustments).

Specifications: Continuously mix until all solid particles have completely dissolved.

End result: Homogeneous liquid-like solution.

2. **Powder-liquid preparation:**

- A. Triturate the Amiodarone Hydrochloride to form a fine, homogeneous powder.
- B. Levigate the fine, homogeneous powder (Step 2A) with the Glycerin.

End result: Homogeneous liquid-like dispersion.

3. **Powder-medium integration:**

- A. In the given order, sequentially add the following ingredients to the homogeneous liquid like solution (step 1C):
 - -Amiodarone Hydrochloride (Step 2B)
 - -Cherry Flavor

<u>Specifications</u>: Continuously mix, using high-shear mixing techniques.

End result: Homogeneous liquid-like dispersion.

Note: Add the next ingredient, once the previous one has been completely added and dispersed.



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4. **Filling to volume:**

A. Add Syrup (Simple) to the mixture (Step 3A) to fill to the required batch size (50.0 mL *plus* processing error adjustments).

Specifications: Continuously mix, using high-shear mixing techniques.

End result: Homogeneous liquid-like dispersion.

5. **pH testing:**

- A. Draw an appropriate amount of the mixture (Step 4A).
- B. Test the pH of the sample. It should lie between 4.3 and 4.5.
- C. If the pH < 4.3, carefully add in a dropwise manner the Monohydrate Potassium Citrate 20% Solution to the mixture:
 - 1. Draw and transfer 1 or 2 drops of the Monohydrate Potassium Citrate 20% Solution to the mixture.
 - 2. Stir for at least 5 minutes to evenly disperse the Monohydrate Potassium Citrate 20% Solution.
 - 3. Re-test the pH.
 - 4. Continue to add the Monohydrate Potassium Citrate 20% Solution until the pH of 4.3 to 4.5 is obtained.

IMPORTANT: Do not allow the pH to rise above 4.5.

- D. If the pH > 4.5, carefully add in a dropwise manner the Citric Acid 20% Solution to the mixture:
 - 1. Draw and transfer 1 or 2 drops of the Citric Acid 20% Solution to the mixture.
 - 2. Stir for at least 5 minutes to evenly disperse the Citric Acid 20% Solution.
 - 3. Re-test the pH.
 - 4. Continue to add the Citric Acid 20% Solution until the pH of 4.3 to 4.5 is obtained.

IMPORTANT: Do not allow the pH to fall below 4.3.

6. **Product transfer:**

Transfer the final product into the specified dispensing container (see "Packaging requirements").

<u>Note</u>: Continuously mix the final product during the transfer process into the recommended dispensing containers in order to maintain homogeneity.



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SUGGESTED PRESENTATION

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Estima Beyond-Use D		14 days, refrigerated, as per USP.	Packa Requirem		-Tightly closed, light-resistant dispensing bottleTo be administered with a metered dose-measuring device.
	1	Use as directed. Do not exceed dose.	prescribed	7	Protect from light.
	2	Keep out of reach of children.		8	Cap tightly after use.
	3	Keep refrigerated. Do not freeze.		9	Shake well before use.
Auxiliary Labels	4	May impair mental and/or physic Use care when operating a machinery.	•	10	Do not take with alcohol, sleep aids, tranquilizers or other CNS depressants.
	5	Consult your health care practition prescription or over-timedications are currently being uprescribed for future use.	he-counter	11	Do not take with grapefruit juice.
	6	Patient must avoid exposure to sartificial UV rays.	sunlight or		
Pharmacist Instructions	Ad	d any auxiliary labels specific to the	e active ingr	edien	t to the dispensing container as deemed necessary.
Patient Instructions	Co	ntact your pharmacist in the event o	of adverse re	action	ns.

REFERENCES

1.	Suspensions. In: Allen, LV, Jr. <i>The Art, Science and Technology of Pharmaceutical Compounding Fourth Edition.</i> American Pharmaceutical Association; 2012: 239.
2.	Amiodarone Hydrochloride for Injection Sandoz Standard. In: Canadian Pharmacists Association. <i>Compendium of Pharmacists and Specialties</i> , 2008. 139.
3.	Glycerin. In: Rowe RC. <i>Handbook of Pharmaceutical Excipients</i> , 7 th <i>Edition</i> . American Pharmaceutical Association; 2012: 338.
4.	Amiodarone Hydrochloride. In: Sweetman SC, ed. <i>Martindale: The Complete Drug Reference, 34th Edition.</i> London, England: The Pharmaceutical Press; 2005: 859.
5.	Amiodarone (Monograph). In: O'Neil MJ. <i>The Merck Index 15th Edition</i> . Whitehouse Station, NJ: Merck & Co, Inc.; 2013: Monograph #478.
6.	Amiodarone Hydrochloride. In: Trissel LA. <i>Trissel's Stability of Compounded Formulations</i> , 5 rd Edition. American Pharmaceutical Association; 2012: 32.



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7.	Amiodarone Systemic. Thomson Micromedex. <i>USP DI – Drug Information for the Health Care Professional</i> , 26 th <i>Edition</i> . Taunton, MA: US Pharmacopeial Convention, Inc; 2006: 106.
8.	Methylcellulose. In: Rowe RC. <i>Handbook of Pharmaceutical Excipients</i> , 7 th <i>Edition</i> . American Pharmaceutical Association; 2012: 496.
9.	Sodium Benzoate. In: Rowe RC. <i>Handbook of Pharmaceutical Excipients</i> , 7 th Edition. American Pharmaceutical Association; 2012: 718.
10.	Methylcellulose. In: Sweetman SC, ed. <i>Martindale: The Complete Drug Reference</i> , 36 th Edition. London, England: The Pharmaceutical Press; 2009: 2145.
11.	Sodium Benzoate. In: Sweetman SC, ed. <i>Martindale: The Complete Drug Reference</i> , 36 th Edition. London, England: The Pharmaceutical Press; 2009: 1630.
12.	Methylcellulose (Monograph). In: O'Neil MJ. <i>The Merck Index 15th Edition</i> . Whitehouse Station, NJ: Merck & Co, Inc.; 2013: Monograph #6111.
13.	Sodium Benzoate (Monograph). In: O'Neil MJ. <i>The Merck Index 15th Edition</i> . Whitehouse Station, NJ: Merck & Co, Inc.; 2013: Monograph #8718.
14.	Methylcellulose (Monograph). <i>United States Pharmacopeia XXXVII / National Formulary 32</i> . Rockville, MD. US Pharmacopeial Convention, Inc. 2014: 3776.
15.	USP <795>. <i>United States Pharmacopeia XXXVIII / National Formulary 33</i> . Rockville, MD. US Pharmacopeial Convention, Inc. 2015: 559.

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