

TOLL-FREE: 866-333-7811 TELEPHONE: 514-905-5096 FAX: 514-905-5097 technicalservices@medisca.net

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Suggested Formula

Mexiletine Hydrochloride 250 mg Oral Capsules (Powder Blend, 100 x Size #1 Capsules)

FIN

F 004 349v2

SUGGESTED FORMULATION

Ingredient Listing	Qty.	Unit	NDC#	Supplier	Lot Number	Expiry Date
Mexiletine Hydrochloride, USP	25.000	g				
Medisca CapsuBlend™-S	TBD					
Sodium Chloride, USP	As required					

SPECIAL PREPARATORY CONSIDERATIONS

Ingredient-Specific Information		
Light sensitive (protect from lig	ght whenever possible):	Mexiletine Hydrochloride
		5.0
Hygroscopic (protect from moi	sture whenever possible):	CapsuBlend [™] -S
Suggested Preparatory Guidelines		OP
Non-Sterile Preparat	ion Sterile Preparation	
Processing Error / Testing Considerations:		considerations during preparation, it is suggested to the required quantities of ingredients.
Special Instruction:	Protective apparel, such as a lab c should always be worn.	oat, 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 100 Size #1 Capsules)

Weigh and / or measure the following ingredients when appropriate:

Ingredient Listing	Qty.	Unit	Multiplication factor (*):	Processing Error	Qty. to measure
Mexiletine Hydrochloride, USP §	25.000	g			
Medisca CapsuBlend™-S §	TBD				
Sodium Chloride, USP	As required		⊗		

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

Preparatory Instruction

1. CapsuBlendTM-S requirements for 100 x size #1 capsules

A. Calculate the amount of CapsuBlend™-S required for the batch. Refer to attached appendix for details.

2. **Powder preparation:**

- A. By geometric addition, combine and mix the following ingredients together:
 - -Mexiletine Hydrochloride
 - -CapsuBlendTM-S (Quantity determined in appendix (**I**))

3. **Product transfer:**

Fill each of 100 Size #1 capsules with the powder blend (Step 2A). Close each capsule tightly.

Clean each capsule by placing the capsules in a container filled with Sodium chloride, and then gently rolling the container. Pour the container contents into a 10-mesh sieve, and allow the Sodium chloride to pass through. Finally, roll the capsules on a cloth-covered surface.

4. Validation technique (average capsule weight):

The final weight of each capsule (not including capsule shell) should fall between 90 and 110% of the theoretically calculated weight, in accordance to USP 795 guidelines. The theoretically calculated weight can be determined by adding the amount in appendix (\mathbf{G}) + 0.250 g together.

5. **Product transfer:**

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



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SUGGESTED PRESENTATION

Estimated Beyond-Use Date		6 months, as per USP	Packa Requirem		Tightly closed, light-resistant capsule shells and vials.
	1	Use as directed. Do not exceed prescribed dose.		6	Keep in a dry place.
	2	2 Keep out of reach of children.		7	Keep at room temperature (20°C – 23°C).
A '11'	3	Cap tightly after use.		8	Protect from light.
Auxiliary Labels	4	Consult your health care practitioner if any other prescription or over-the-counter medications are currently being used or are prescribed for future use.		9	Do not take with alcohol, sleep aids, tranquilizers or other CNS depressants.
	5	May impair mental and/or phys Use care when operating machinery.		10	Take with food or antacid.
Pharmacist Instructions	Add any auxiliary labels specific to the API to the dispensing container as deemed necessary.				
Patient Instructions	Contact your pharmacist in the event of adverse reactions.				

REFERENCES

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2.	Novo-Mexiletine. In: Canadian Pharmacists Association. Compendium of Pharmacists and Specialties, 2010. 1613.
3.	Sodium Chloride. In: Rowe RC. <i>Handbook of Pharmaceutical Excipients</i> , 6 th <i>Edition</i> . American Pharmaceutical Association; 2009: 637.
4.	Mexiletine Hydrochloride. In: Sweetman SC, ed. <i>Martindale: The Complete Drug Reference</i> , 36 th Edition. London, England: The Pharmaceutical Press; 2009: 1339.
5.	Mexiletine (Monograph). In: O'Neil MJ. <i>The Merck Index 14th Edition</i> . Whitehouse Station, NJ: Merck & Co, Inc.; 2006: Monograph #6169.
6.	Mexiletine Hydrochloride (Monograph). <i>United States Pharmacopeia XXXII / National Formulary 27</i> . Rockville, MD. US Pharmacopeial Convention, Inc. 2009: 2976.
7.	Mexiletine Hydrochloride. In: Trissel LA. <i>Trissel's Stability of Compounded Formulations, 4th Edition</i> . American Pharmaceutical Association; 2009: 380.
8.	USP <795>. <i>United States Pharmacopeia XXXII / National Formulary 27</i> . Rockville, MD. US Pharmacopeial Convention, Inc. 2009: 314.

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		Procedure		
1.	Capsule filling: a. For each ingredient powder below, de CAPSULES. Do not forget to divide	C I	<i>.</i> .	2 2
	Plug each amount into Step 2, colum	n B.		
2.	Volume Percent Occupied:		©	
	<u>Ingredients</u>	Column A Quantity Required per capsule	Column B Average capsule fill weight	Column C A/B x 100 equals percent filled
	a. Mexiletine Hydrochloride	0.250 g	g	%
	b. CapsuBlend™-S		g	
	c. Total (add column C together)			% (D)
3.	Calculate the quantity of CapsuBlend ^T	M-S required for the batch	<u>ı:</u>	
	a. Percent of CapsuBlend™-S required =	= 100% – (D)		% (E)
	b. Average capsule fill weight of Capsul	Blend™-S (from column B,	Step 2b):	g (F)
	c. Quantity of CapsuBlend™-S required	per capsule = $[(E) \div 100 \times$	(F)]	g (G)
	d. Total quantity of CapsuBlend™-S req	uired for the batch = 100 ca	apsules × (G)	g (H)
	e. Total quantity of CapsuBlend™-S <i>plu</i>	as processing error = $(H) \times 1$	1.05-1.09	g (I)

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