



MEDISCA® NETWORK INC.  
TECHNICAL SUPPORT SERVICES  
FORMULATION CHEMISTRY DEPARTMENT  
TOLL-FREE: 866-333-7811  
TELEPHONE: 514-905-5096  
FAX: 514-905-5097  
[technicalservices@medisca.net](mailto:technicalservices@medisca.net)

11/9/2022; Page 1

Suggested Formula	Amoxicillin 250 mg/5 mL Oral Liquid (Suspension, 150 mL)	FIN	F 009 743
-------------------	--	-----	-----------

### SUGGESTED FORMULATION

Ingredient Listing	Qty.	Unit	NDC #	Supplier	Lot Number	Expiry Date
Amoxicillin (250 mg) Capsule	30	Units				
Strawberry Flavor	0.3	mL				
Banana Flavor	0.3	mL				
Stevia Powder	0.75	g				
Glycerin, USP	10.0	mL				
Medisca Oral Mix (Flavored Suspending Vehicle)	50.0	mL				
Medisca Oral Mix (Flavored Suspending Vehicle)	q.s. to 150.0	mL				
Sodium Hydroxide 10% Solution	As required					
Hydrochloric Acid 10% Solution	As required					



Suggested Formula	Amoxicillin 250 mg/5 mL Oral Liquid (Suspension, 150 mL)	FIN	F 009 743
-------------------	--	-----	-----------

## SPECIAL PREPARATORY CONSIDERATIONS

### Ingredient-Specific Information

**Hygroscopic** (protect from moisture whenever possible): *Stevia Powder, Glycerin*

**Light sensitive** (protect from light whenever possible): *Amoxicillin*

### Suggested Preparatory Guidelines

Non-Sterile Preparation     Sterile Preparation

Processing Error / Testing Considerations: To account for processing error considerations during preparation, it is suggested to measure an additional **5 to 9 %** of the required quantities of ingredients.

Special Instruction: This formula may contain one or more Active Pharmaceutical Ingredients (APIs) that may be classified as hazardous, please refer & verify the current NIOSH list of Antineoplastic and Other Hazardous Drugs in Healthcare Settings. At this time, **General Chapter <800> Hazardous Drugs – Handling in Healthcare Settings** is informational and not compendially applicable unless otherwise specified by regulators and enforcement bodies. For information on the scope, intended applicability, and implementation context for USP General Chapter <800>, see: <https://www.usp.org/compounding/general-chapter-hazardous-drugs-handling-healthcare>.

This formula must be prepared within the appropriate facilities under adequate environmental conditions, following the necessary guidelines and procedures as stated within *USP 795* and *USP 800*, when handling hazardous drugs. Only trained and qualified personnel must prepare this formula.

All required personal protective equipment (hazardous if applicable), such as but not limited to, lab coat, protective sleeves, gloves both inner and outer if applicable, dedicated shoe covers, hairnet, beard cover, eyewear, appropriate face mask, respirator and face shield, etc., where applicable must be worn at all times.

If applicable, follow all required procedures for hazardous drug handling including but not limited to procurement, transport, storage, preparation, dispensing, administration, clean up (spills) & disposal.

If you are a registered 503B facility, please refer to all relevant guidance documents including but not limited to the Code of Federal Regulations (CFR), Guidance for Industry (GFI) and Compliance Policy Guides (CPGs).

This procedure requires the use of very small quantities of ingredients. All calculations and preparation techniques must be verified before dispensing the final product.



Suggested Formula	Amoxicillin 250 mg/5 mL Oral Liquid (Suspension, 150 mL)	FIN	F 009 743
-------------------	--	-----	-----------

**SUGGESTED PREPARATION (for 150 mL)**

Weigh and / or measure the following ingredients when appropriate:

Ingredient Listing	Qty.	Unit	Multiplication factor (*): ____	Processing Error	Qty. to measure
Amoxicillin (250 mg) Capsule §	30	Units			
Strawberry Flavor	0.3	mL			
Banana Flavor	0.3	mL			
Stevia Powder §	0.75	g			
Glycerin, USP §	10.0	mL			
Medisca Oral Mix (Flavored Suspending Vehicle)	50.0	mL			
Medisca Oral Mix (Flavored Suspending Vehicle)	q.s. to 150.0	mL			
Sodium Hydroxide 10% Solution	As required				
Hydrochloric Acid 10% Solution	As required				

\* Takes into account increased batch size conversions and density conversions, if required.

§ Weigh / measure just prior to use.



Suggested Formula	Amoxicillin 250 mg/5 mL Oral Liquid (Suspension, 150 mL)	FIN	F 009 743
-------------------	--	-----	-----------

Preparatory Instruction

1. **Ingredient quantification (determine the actual quantity of Amoxicillin 250 mg Capsule powder mix to weigh):**

A. Empty and weigh the contents of 33 Amoxicillin 250 mg Capsules.  
Record the total weight here: \_\_\_\_\_ g

B. Calculate the average weight of powder in each capsule:

Weight of powder from 33 capsules (from Step 1A):	_____ g
DIVIDED BY	
Number of capsules:	33
EQUALS	
Average weight of a single Amoxicillin 250 mg Capsule:	_____ g

C. Calculate the weight of powder equivalent to 30 capsules:

Average weight of a single Amoxicillin 250 mg Capsule (from Step 1B):	_____ g
MULTIPLIED BY	
Number of capsules required:	30
EQUALS	
Weight of powder equivalent to 30 capsules:	_____ g

D. Calculate the weight of powder required *plus* processing error adjustments:

Weight of powder equivalent to 30 capsules (from Step 1C):	_____ g
MULTIPLIED BY	
Processing error adjustments (5 to 9%):	1.05 to 1.09
EQUALS	
<b>Weight of powder required <i>plus</i> processing error adjustments:</b>	_____ g



Suggested Formula	Amoxicillin 250 mg/5 mL Oral Liquid (Suspension, 150 mL)	FIN	F 009 743
-------------------	--	-----	-----------

2.	<p><b><u>Powder-liquid preparation:</u></b></p> <p>A. Triturate the contents of the 33 Amoxicillin 250 mg Capsules to form a fine, homogeneous powder.</p> <p>B. Weigh the quantity Amoxicillin 250 mg Capsule powder mix required for the batch (refer to Step 1D) and discard the remaining powder.</p>
3.	<p><b><u>Powder-liquid preparation:</u></b></p> <p>A. Combine and triturate the following ingredients together to form a fine, homogeneous powder blend:</p> <ul style="list-style-type: none"><li>-Amoxicillin 250 mg Capsule powder mix (amount weighed from Step 2B)</li><li>-Stevia Powder</li></ul> <p>B. Combine and mix the following ingredients together to form a homogeneous liquid-like solution:</p> <ul style="list-style-type: none"><li>-Glycerin</li><li>-Strawberry Flavor</li><li>-Banana Flavor</li></ul> <p>C. Levigate the fine, homogeneous powder blend (Step 3A) with the homogeneous liquid-like solution (Step 3B).</p> <p><u>End result:</u> Homogeneous paste-like dispersion.</p>
4.	<p><b><u>Medium integration:</u></b></p> <p>A. Incrementally add the homogeneous paste-like dispersion (Step 3C) to the Oral Mix (Flavored Suspending Vehicle) (50.0 mL <i>plus</i> processing error adjustments).</p> <p><u>Specifications:</u> Continuously mix, using high-shear mixing techniques.</p> <p><u>End result:</u> Homogeneous liquid-like dispersion.</p>
5.	<p><b><u>Filling to volume:</u></b></p> <p>A. Add additional Oral Mix (Flavored Suspending Vehicle) to the mixture (Step 4A) to fill to the required batch size (150.0 mL <i>plus</i> processing error adjustments).</p> <p><u>Specifications:</u> Continuously mix until homogeneous.</p> <p><u>End result:</u> Homogeneous liquid-like dispersion.</p>



Suggested Formula	Amoxicillin 250 mg/5 mL Oral Liquid (Suspension, 150 mL)	FIN	F 009 743
-------------------	--	-----	-----------

6.	<p><b><u>pH testing:</u></b></p> <p>A. Draw an appropriate amount of the mixture (Step 5A).</p> <p>B. Test the pH of the sample. It should lie between 6.0 and 6.5.</p> <p>C. <u>If the pH &lt; 6.0 carefully add in a dropwise manner the Sodium Hydroxide 10% Solution to the mixture:</u></p> <ol style="list-style-type: none"><li>1. Draw and transfer 1 or 2 drops of the Sodium Hydroxide 10% Solution to the mixture.</li><li>2. Stir for at least 5 minutes to evenly disperse the Sodium Hydroxide 10% Solution.</li><li>2. Re-test the pH.</li><li>4. Continue to add the Sodium Hydroxide 10% Solution until the pH of 6.0 to 6.5 is obtained.</li></ol> <p>IMPORTANT: Do not allow the pH to rise above 6.5.</p> <p>D. <u>If the pH &gt; 6.5, carefully add in a dropwise manner the Hydrochloric Acid 10% Solution to the mixture:</u></p> <ol style="list-style-type: none"><li>1. Draw and transfer 1 or 2 drops of the Hydrochloric Acid 10% Solution to the mixture.</li><li>2. Stir for at least 5 minutes to evenly disperse the Hydrochloric Acid 10% Solution.</li><li>3. Re-test the pH.</li><li>4. Continue to add the Hydrochloric Acid 10% Solution until the pH of 6.0 to 6.5 is obtained.</li></ol> <p>IMPORTANT: Do not allow the pH to fall below 6.0.</p>
7.	<p><b><u>Product transfer:</u></b></p> <p>A. Transfer the final product into the specified dispensing container (see “Packaging requirements”).</p> <p><u>Note:</u> Continuously mix the final product during the transfer process in order to maintain homogeneity.</p>



Suggested Formula	Amoxicillin 250 mg/5 mL Oral Liquid (Suspension, 150 mL)	FIN	F 009 743
-------------------	--	-----	-----------

**SUGGESTED PRESENTATION**

Estimated Beyond-Use Date	14 days, refrigerated, as per USP 795*.	Packaging Requirements	- Tightly closed, light-resistant dispensing bottle. - To be administered with a metered measuring device.	
Auxiliary Labels	1	Use as directed. Do not exceed prescribed dose.	6	Keep out of reach of children.
	2	Keep refrigerated (2°C – 8°C). Do not freeze.	7	Consult your health care practitioner if any prescription or over-the-counter medications are currently being used or are prescribed for future use.
	3	<b>Shake well before use.</b>	8	Cap tightly after use.
	4	May impair mental and/or physical ability. Use care when operating a car or machinery.	9	Do not take with alcohol, sleep aids, tranquilizers or other CNS depressants.
	5	Protect from light.		
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.			

\* If the API or any other components in the CNSP have an expiration date that is earlier than the assigned BUD, the expiration date supersedes the assigned BUD and must be the assigned shortest date.



Suggested Formula	Amoxicillin 250 mg/5 mL Oral Liquid (Suspension, 150 mL)	FIN	F 009 743
-------------------	--	-----	-----------

## REFERENCES

1.	Suspensions. In: Allen, LV, Jr. <i>The Art, Science, and Technology of Pharmaceutical Compounding Fifth Edition</i> . American Pharmacists Association; 2016: 279.
2.	Amoxicillin. In: Sweetman SC, ed. <i>Martindale: The Complete Drug Reference, 38<sup>th</sup> Edition</i> . London, England: The Pharmaceutical Press; 2014: 216.
3.	Amoxicillin (Monograph). In: O'Neil MJ. <i>The Merck Index 15<sup>th</sup> Edition</i> . Whitehouse Station, NJ: Merck & Co, Inc.; 2012: Monograph #574.
4.	Amoxicillin. In: Trissel LA. <i>Trissel's Stability of Compounded Formulations, 5<sup>rd</sup> Edition</i> . American Pharmaceutical Association; 2012: 35.
5.	Amoxicillin (Monograph). <i>United States Pharmacopeia / National Formulary</i> . Rockville, MD. US Pharmacopeial Convention, Inc. 2022.
6.	USP <795>. <i>United States Pharmacopeia / National Formulary</i> . Rockville, MD. US Pharmacopeial Convention, Inc. 2022.

DISCLAIMER: THIS DOCUMENT IS COPYRIGHT© 2022 MEDISCA PHARMACEUTIQUE INC. MEDISCA NETWORK INC., HEREBY REFERRED TO AS 'THE NETWORK', HAS PROVIDED THE FORMULA AND INSTRUCTIONS ABOVE AS A MODEL FOR EDUCATIONAL PURPOSES ONLY ON THE BASIS OF THE RECOGNIZED COMPENDIA AND TEXTS REFERENCED AT THE END OF THIS DOCUMENT. THE NETWORK TAKES NO RESPONSIBILITY FOR THE VALIDITY, SCHEDULING OR ACCURACY OF THIS INFORMATION OR FOR ITS SAFETY OR EFFECTIVENESS, NOR FOR ANY USE THEREOF, WHICH IS AT THE SOLE RISK OF THE LICENSED PHARMACIST OR OTHER APPROPRIATELY STATE LICENSED PROFESSIONAL. ADJUSTMENTS MAY BE NEEDED TO MEET SPECIFIC PATIENT NEEDS AND IN ACCORDANCE WITH A LICENSED PRESCRIBER'S PRESCRIPTION. THE PHARMACIST OR OTHER APPROPRIATELY STATE LICENSED PROFESSIONAL MUST EMPLOY APPROPRIATE TESTS TO DETERMINE THE STABILITY OF THIS SUGGESTED FORMULA. THE NETWORK CANNOT BE HELD LIABLE TO ANY PERSON OR ENTITY CONCERNING CLAIMS, LOSS, OR DAMAGE CAUSED BY, OR ALLEGED TO BE CAUSED BY, DIRECTLY OR INDIRECTLY, THE USE OR MISUSE OF THE INFORMATION CONTAINED IN THIS SUGGESTED FORMULA. IN ALL CASES IT IS THE RESPONSIBILITY OF THE LICENSED PHARMACIST OR OTHER APPROPRIATELY STATE LICENSED PROFESSIONAL TO KNOW THE LAW, TO COMPOUND ANY FINISHED PRODUCT AND TO DISPENSE THESE PRODUCTS IN ACCORDANCE WITH FEDERAL AND STATE LAW. MEDISCA NETWORK INC. MAKES NO WARRANTIES WITH RESPECT TO INFRINGEMENT OR NON-INFRINGEMENT BY THE FORMULA OF ANY PATENT OR OTHER INTELLECTUAL PROPERTY OF ANY OTHER PARTY, AND IT IS THE RESPONSIBILITY OF THE PHARMACIST OR OTHER APPROPRIATELY STATE LICENSED PROFESSIONAL TO INVESTIGATE AND DETERMINE ANY SUCH ISSUE.