



Suggested Formula	Nystatin 100,000 IU/mL Oral Mucoadhesive Liquid (Suspension, 100 mL)	FIN	F 008 262
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SUGGESTED FORMULATION

Ingredient Listing	Qty.	Unit	NDC #	Supplier	Lot Number	Expiry Date
Nystatin, USP	10	MU				
Glycerin, USP	3.0	mL				
Cherry Flavor	1.0	mL				
Medisca NovaFilm™	30.0	mL				
Medisca Oral Mix SF (Sugar-Free Flavored Suspending Vehicle)	20.0	mL				
Medisca Oral Mix SF (Sugar-Free Flavored Suspending Vehicle)	q.s. to 100.0	mL				
Sodium Hydroxide 10% Solution	As required					

SPECIAL PREPARATORY CONSIDERATIONS

Ingredient-Specific Information

<i>Light Sensitive (protect from light whenever possible):</i>	<i>Nystatin</i>
<i>Hygroscopic (protect from moisture whenever possible):</i>	<i>Glycerin, Nystatin</i>
<i>Oxygen Sensitive (protect from oxygen whenever possible):</i>	<i>Nystatin</i>
<i>Moisture Sensitive (protect from humidity whenever possible):</i>	<i>Nystatin</i>
<i>Air Sensitive (protect from air whenever possible):</i>	<i>Nystatin</i>
<i>Heat Sensitive (protect from heat whenever possible):</i>	<i>Nystatin</i>



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SPECIAL PREPARATORY CONSIDERATIONS (CONTINUED)

Suggested Preparatory Guidelines

Non-Sterile Preparation Sterile Preparation

Processing Error / Testing Considerations: To account for processing error and pH testing 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, 2016. **General Chapter <800> Hazardous Drugs – Handling in Healthcare Settings** was formally published February 1, 2016 in the First Supplement to USP 39-NF 34 and has a delayed **official implementation date of December 31st, 2019**.

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 (GFIs) 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.



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SUGGESTED PREPARATION (for 100 mL)

Weigh and / or measure the following ingredients when appropriate:

Ingredient Listing	Qty.	Unit	Multiplication factor ^(*) : _____	Processing Error	Qty. to measure
Nystatin, USP §	10	MU			
Glycerin, USP §	3.0	mL			
Cherry Flavor	1.0	mL			
Medisca NovaFilm™	30.0	mL			
Medisca Oral Mix SF (Sugar-Free Flavored Suspending Vehicle)	20.0	mL			
Medisca Oral Mix SF (Sugar-Free Flavored Suspending Vehicle)	q.s. to 100.0	mL			
Sodium Hydroxide 10% Solution	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. Ingredient quantification:

A. Determine the quantity (in g) of Nystatin required to make a Nystatin 100,000 IU/mL Liquid, batch size (100 mL):

Quantity of Nystatin (in Units) required for 100 mL	10,000,000 IU
DIVIDED BY	
Nystatin biopotency assay result (from Certificate of Analysis)	_____ IU/mg
EQUALS	
i. Quantity of Nystatin (in milligrams) required	_____ mg
MULTIPLIED BY	
Multiplication factor – milligrams to grams	0.001
EQUALS	
ii. Quantity of Nystatin (in grams) required	_____ g
MULTIPLIED BY	
Processing error adjustments (5 to 9%)	1.05 to 1.09
EQUALS	
iii. Quantity of Nystatin needed <i>plus</i> processing error adjustments	_____ g

2. Powder-liquid preparation:

A. Triturate the Nystatin (amount determined in Step 1Aiii) to form a fine, homogeneous powder.

B. Levigate the fine, homogeneous powder (Step 2A) with the Glycerin.

End result: Homogeneous paste-like dispersion.



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3.	<p><u>Medium integration:</u></p> <p>A. In the given order, sequentially add the following ingredients to the Oral Mix SF (Sugar-Free Flavored Syrup Vehicle) (20.0 mL <i>plus</i> processing error adjustments):</p> <ul style="list-style-type: none">-Homogeneous paste-like dispersion (Step 2A)-Cherry Flavor-NovaFilm™ <p><u>Specifications:</u> Continuously mix, using high-shear mixing techniques.</p> <p><u>End result:</u> Homogeneous liquid-like dispersion.</p> <p><u>Note:</u> Add the next ingredient, once the previous one has been completely added and dispersed.</p>		
4.	<p><u>Filling to volume:</u></p> <p>A. Add additional Oral Mix SF (Sugar-Free Flavored Syrup Vehicle) to the mixture (Step 3A) to fill to the required batch size (100.0 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><u>pH testing:</u></p> <p>A. Draw an appropriate amount of the mixture (Step 4A).</p> <p>B. Test the pH of the sample. It should lie between 6.0 and 8.0.</p> <p>C. <u>If the pH < 6.0, carefully add, in a dropwise fashion, the Sodium Hydroxide 10% Solution to the mixture:</u></p> <ol style="list-style-type: none">1. Draw and transfer 1 or 2 drops of the Sodium Hydroxide 10% Solution to the mixture.2. Stir for at least 5 minutes to evenly disperse the Sodium Hydroxide 10% Solution.3. Re-test the pH.4. Continue to add the Sodium Hydroxide 10% Solution until the pH of 6.0 to 8.0 is obtained. <p>IMPORTANT: Do not allow the pH to rise above 8.0.</p>		
6.	<p><u>Product transfer:</u></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 into the recommended dispensing containers in order to maintain homogeneity.</p>		



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SUGGESTED PRESENTATION

Estimated Beyond-Use Date		Packaging Requirements	
	7 days, refrigerated, as per USP.		- Tightly closed, light-resistant dispensing bottle. - To be administered with a metered-dose measuring device.
Auxiliary Labels	1 Use as directed. Do not exceed prescribed dose.	5	Protect from light.
	2 Keep out of reach of children.	6	Shake well before use.
	3 Cap tightly after use.	7	Consult your health care practitioner if any prescription or over-the-counter medications are currently being used or are prescribed for future use.
	4 Keep refrigerated (2°C – 8°C). Do not freeze.	8	
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.		



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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.	Glycerin. In: Rowe RC. <i>Handbook of Pharmaceutical Excipients, 7th Edition</i> . American Pharmaceutical Association; 2012: 324.
3.	Nystatin. In: Sweetman SC, ed. <i>Martindale: The Complete Drug Reference, 36th Edition</i> . London, England: The Pharmaceutical Press; 2009: 543.
4.	Nystatin (Monograph). In: O'Neil MJ. <i>The Merck Index 15th Edition</i> . Whitehouse Station, NJ: Merck & Co, Inc.; 2013: Monograph #6825.
5.	Nystatin. In: Trissel LA. <i>Trissel's Stability of Compounded Formulations, 5rd Edition</i> . American Pharmaceutical Association; 2012: 357.
6.	Nystatin (Monograph). <i>United States Pharmacopeia XLII / National Formulary 37</i> . Rockville, MD. US Pharmacopeial Convention, Inc. 2019: 3186.
7.	USP <795>. <i>United States Pharmacopeia XLII / National Formulary 37</i> . Rockville, MD. US Pharmacopeial Convention, Inc. 2019: 6951.

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