



Suggested Formula	Oxytocin 40 IU Oral Transmucosal Films (Solid Suspension, 30 Films)	FIN	F 009 711
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SUGGESTED FORMULATION

Ingredient Listing	Qty.	Unit	NDC #	Supplier	Lot Number	Expiry Date
Oxytocin 2000 Units/mL Stock Solution †	0.60	mL				
Bitterness Reducing Agent (NF-01) (Natural) (Powder)	0.25	g				
Cellulose (Microcrystalline), NF	0.60	g				
Menthol (Crystals) (Levorotatory) (Natural), USP	0.06	g				
Sucralose, NF	0.05	g				
Glycerin, USP	0.075	g				
Purified Water, USP	8.0	mL				
NovaFilm™ Gel Base	18.00	g				
Purified Water, USP	q.s. to 30.0	mL				
† Oxytocin 2000 Units/mL Stock Solution						
Oxytocin, USP	TBD					
Purified Water, USP	19.0	mL				
Purified Water, USP	q.s. to 20.0	mL				



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SPECIAL PREPARATORY CONSIDERATIONS

Ingredient-Specific Information

Hygroscopic (protect from moisture whenever possible):	Oxytocin, Cellulose, Glycerin
Light Sensitive (protect from light whenever possible):	Oxytocin, NovaFilm™ Gel Base
Heat Sensitive (protect from heat whenever possible):	Oxytocin

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 **12 to 15%** 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.



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SUGGESTED PREPARATION (for 30 Films)

Weigh and / or measure the following ingredients when appropriate:

Ingredient Listing	Qty.	Unit	Multiplication factor (*): ____	Processing Error	Qty. to measure
Oxytocin 2000 Units/mL Stock Solution † §	0.60	mL			
Bitterness Reducing Agent (NF-01) (Natural) (Powder)	0.25	g			
Cellulose (Microcrystalline), NF §	0.60	g			
Menthol (Crystals) (Levorotatory) (Natural), USP	0.06	g			
Sucralose, NF	0.05	g			
Glycerin, USP §	0.075	g			
Purified Water, USP	8.0	mL			
NovaFilm™ Gel Base §	18.00	g			
Purified Water, USP	q.s. to 30.0	mL			
† Oxytocin 2000 Units/mL Stock Solution					
Oxytocin, USP §	TBD		---	---	
Purified Water, USP	19.0	mL	---	---	
Purified Water, USP	q.s. to 20.0	mL	---	---	



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Preparatory Instruction

1. Oxytocin Ingredient quantification (units per weight measure adjustment):

A. Determine the quantity (in g) of Oxytocin required to make an Oxytocin 2000 Units/mL Stock Solution, batch size (20.0 mL):

Quantity of Oxytocin (in Units) required	40 000 IU
DIVIDED BY	
Oxytocin biopotency assay result (from Certificate of Analysis)	_____ IU/mg
EQUALS	
Quantity of Oxytocin (in milligrams) required	_____ mg
MULTIPLIED BY	
Multiplication factor – milligrams to grams	0.001
EQUALS	
Quantity of Oxytocin (in grams) required for the Stock Solution	_____ g

2. † Oxytocin 2000 Units/mL Stock Solution Preparation:

A. Incrementally add the Oxytocin (amount determined in Step 1A) to the Purified Water (19.0 mL).

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

End result: Homogeneous liquid-like solution.

B. Add additional Purified Water to the mixture (Step 2A) to fill to the required batch size (20.0 mL).

Specifications: Continuously mix.

End result: Homogeneous liquid-like solution



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3.	<p><u>Powder-liquid preparation:</u></p> <p>A. By geometric addition, combine and triturate the following ingredients together to form a fine, homogeneous powder blend:</p> <ul style="list-style-type: none">-Bitterness Reducing Agent (NF-01) (Natural) (Powder)-Cellulose (Microcrystalline)-Menthol (Crystals) (Levorotatory) (Natural)-Sucralose <p>B. Levigate the fine, homogeneous powder blend (Step 3A) with the Oxytocin 2000 Units/mL Stock Solution (0.60 mL <i>plus</i> processing error adjustments), Glycerin and Purified Water (8.0 mL <i>plus</i> processing error adjustments).</p> <p><u>End result:</u> Homogeneous liquid-like dispersion.</p>
4.	<p><u>Medium incorporation:</u></p> <p>A. Incrementally add the homogeneous liquid-like dispersion (Step 3B) to the NovaFilm™ Gel Base.</p> <p><u>Specifications:</u> Continuously mix, using high-shear mixing techniques.</p> <p><u>NOTE:</u> Ensure mixing process does not incorporate any air.</p> <p><u>End result:</u> Homogeneous gel-like dispersion.</p>
5.	<p><u>Filling to volume:</u></p> <p>A. Add additional Purified Water to the mixture (Step 4A) to fill to the required batch size (30.0 mL <i>plus</i> processing error adjustments).</p> <p><u>Specifications:</u> Continuously mix, using high-shear mixing techniques.</p> <p><u>NOTE:</u> Ensure mixing process does not incorporate any air.</p> <p><u>End result:</u> Homogeneous gel-like dispersion.</p>
6.	<p><u>Mold filling and drying:</u></p> <p>A. Fill the 30 blister mold cavities with 1.00 mL of the homogeneous gel-like dispersion (Step 5A) per cavity. Spread the homogeneous gel-like dispersion in the cavity to a uniform thickness.</p> <p><u>NOTE:</u> Ensure no air bubbles are added to the mold cavities.</p> <p>B. Under a laminar air flow hood, allow the filled blister molds to dry at room temperature overnight.</p> <p><u>Specifications:</u> Homogeneous solid dispersion.</p>



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7.	<p><u>Validation technique:</u></p> <p>A. Weigh 6 films separately.</p> <p>B. The final weight of each film from Step 7A (not including the weight of the blister mold) shall not be less than 90% and not more than 110% of the theoretically calculated weight 0.141 g in accordance to USP guidelines.</p>
8.	<p><u>Product transfer:</u></p> <p>Transfer the final product into the specified dispensing container (see “Packaging requirements”).</p>

SUGGESTED PRESENTATION

Estimated Beyond-Use Date	180 days, controlled room temperature or refrigerator, as per USP 795*.	Packaging Requirements	Manually lock blister molds and put into light-resistant resealable foil pouch.	
Auxiliary Labels	1	Use as directed. Do not exceed prescribed dose.	5	Do not take with alcohol, sleep aids, tranquilizers or other CNS depressants.
	2	Keep out of reach of children.	6	Keep at controlled room temperature (20°C – 25°C) OR keep refrigerated (2°C – 8°C). Do not freeze.
	3	Keep in a dry place.	7	Protect from light.
	4	Discard container after use.	8	Consult your health care practitioner if any other prescription or over-the-counter medications are currently being used or are prescribed for future use.
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.



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REFERENCES

1.	Lozenge, Troches, and Films. In: Allen, LV, Jr. <i>The Art, Science, and Technology of Pharmaceutical Compounding Fifth Edition</i> . American Pharmacists Association; 2016: 215.
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7.	Oxytocin (Monograph). In: O'Neil MJ. <i>The Merck Index 15th Edition</i> . Whitehouse Station, NJ: Merck & Co, Inc.; 2013: Monograph #7078.
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