MEDISCA® NETWORK INC.
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FORMULATION CHEMISTRY DEPARTMENT
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Suggested Formula	Furosemide 10 mg/mL Intramuscular/Intravenous Injection (Solution, 50 mL)	FIN	F 005 095v2

SUGGESTED FORMULATION

Ingredient Listing	Qty.	Unit	NDC #	Supplier	Lot Number	Expiry Date
Furosemide, USP	0.500	g				
Sodium Chloride, USP	0.36	g				
Sterile Water for Injection, USP	40.0	mL				
Sterile Water for Injection, USP	q.s. to 50.0	mL				
Sodium Hydroxide 10% Solution	As required					

SPECIAL PREPARATORY CONSIDERATIONS

Ingredient-Specific Information

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

Moisture sensitive *(protect from humidity whenever possible):* *Furosemide*

Suggested Preparatory Guidelines

☐ Non-Sterile Preparation ■ Sterile Preparation

Processing Error / Testing Considerations: To account for processing error, sterility, endotoxin 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 must be prepared within the appropriate facilities under adequate environmental conditions, following the necessary guidelines and procedures as stated within *USP 797*. Only trained and qualified personnel must prepare this formula.

All heat stable, reusable materials and equipment must be sterilized and depyrogenated by dry heat sterilization at 250°C for 2 hours prior to use.

Every batch of final product compounded using this procedure must be sterility and endotoxin tested before being dispensed.

Protective apparel, such as a sterile gown, sterile gloves, shoe covers, head cap, eyewear and face-masks should always be worn. In addition, proper personnel cleansing must be done before entering the buffer or clean area.

Filter integrity must be validated by performing a filter stress test. If the test demonstrates that the filter might be defective, the solution must be discarded and remade.

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 50 mL)

Weigh and / or measure the following ingredients when appropriate:

Ingredient Listing	Qty.	Unit	Multiplication factor (*): ____	Processing Error	Qty. to measure
Furosemide, USP §	0.500	g			
Sodium Chloride, USP §	0.36	g			
Sterile Water for Injection, USP §	40.0	mL			
Sterile Water for Injection, USP §	q.s. to 50.0	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.

Preparatory Instruction **IMPORTANT: All preparatory procedures must be performed using proper Aseptic Technique**	
1.	**Equipment sterilization:** Following the manufacturer's specifications, sterilize and depyrogenate all heat stable, reusable materials and equipment, then return to ambient temperature.
2.	**Powder-liquid preparation:** A. In the given order, sequentially add the following ingredients to the Sterile Water for Injection (40.0 mL *plus* processing error adjustments): -Furosemide -Sodium Chloride Specifications: Continuously mix until all solid particles have completely dispersed. End result: Homogeneous liquid-like dispersion.

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3. **pH testing:**

A. Draw an appropriate amount of the mixture (Step 2A).

B. Test the pH of the sample. It should lie between 8.0 and 8.5.

C. If the pH < 8.0, carefully add, in a dropwise fashion, the Sodium Hydroxide 10% Solution to the mixture:

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 8.0 to 8.5 is obtained.

IMPORTANT: Do not allow the pH to rise above 8.5.

Note: Once the pH has been adjusted to between 8.0 and 8.5, a clear homogeneous solution will result.

4. **Filling to volume:**

A. Add additional Sterile Water for Injection to the above mixture to fill to the required batch size (50.0 mL *plus* processing error adjustments).

Specifications: Continuously mix.

End result: Homogeneous liquid-like solution.

5. **Filtering and transferring:**

Aseptically filter the solution through a 0.22-µm sterile filter into the recommended dispensing containers (see Packaging requirements). Transfer the remainder into separate dispensing containers. These are to be used as the Test samples for sterility and endotoxin testing.

6. **Filter integrity test:**

Validate filter integrity by performing a filter stress test. If the test demonstrates that the filter might be defective, the solution must be discarded and remade.

7. **Sterility testing:**

Validate the Test samples for sterility and endotoxins, in accordance to current USP 797 regulatory guidelines.

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SUGGESTED PRESENTATION

Estimated Beyond-Use Date		14 days, refrigerated, as per USP. BUD based on a successful sterility and endotoxin test result.	Packaging Requirements		Sterile, tightly closed, light-resistant, unit dose injection vials.
Auxiliary Labels	1	Use as directed. Do not exceed prescribed dose.	7		Preservative free, unit does injection. Single use only, discard container after use.
	2	Keep out of reach of children.	8		Discard in the presence of particulate matter.
	3	Keep refrigerated. Do not freeze.	9		Protect from light.
	4	Do not use if discolored.	10		Consult your health care practitioner if any prescription or over-the-counter medications are currently being used or are prescribed for future use.
	5	Equilibrate to room temperature before use.	11		Keep in a dry place.
	6	Do not take with alcohol, sleep aids, tranquilizers or other CNS depressants.	12		May impair mental and/or physical ability. Use care when operating a car or machinery.
Pharmacist Instructions	Add any auxiliary labels specific to the API to the dispensing container as deemed necessary. To be administered only under the close supervision of the prescribing physician.				
Patient Instructions	Contact your pharmacist in the event of adverse reactions.				

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