



ARROWHEAD REGIONAL MEDICAL CENTER Department of Nursing (NRS)

Policy No. 850.00 Issue 1 Page 1 of 3

SECTION:	VIII. SPECIALTY SERVICES
SUBJECT:	ULTRASONIC ENDOVASCULAR SYSTEM FOR THROMBUS MANAGEMENT
APPROVED BY:	CHIEF NURSING OFFICER

POLICY

This policy outlines the proper use, management, and safety procedures for the Ultrasonic Endovascular System in our hospital setting. It is designed to ensure consistent, safe, and effective treatment for patients with pulmonary embolism and other vascular bed thrombosis.

EQUIPMENT

- I. Ultrasonic Control Unit: The central unit that powers and controls the ultrasound delivery.
- II. Ultrasonic Endovascular Device:
 - A. Consists of two components:
 - 1. <u>Ultrasonic Assisted</u> Infusion Catheter (IC): Delivers the thrombolytic agent and coolant.
 - 2. Ultrasonic Core (USC): Generates ultrasound waves to enhance thrombolysis.
- III. Connector Interface Cable (CIC): Connects the Endovascular Device to the Control Unit.
- IV. Two Hospital-grade infusion pumps for precise delivery of thrombolytic agent and coolant.
- V. Ultrasonic cart: For secure mounting and transport of the Control Unit.

PROCEDURES

- I. Indications for use include but not limited to:
 - A. Ultrasound facilitated, controlled and selective intravascular infusion of practitionerspecified fluids, including thrombolytics, for the treatment of pulmonary embolism and other vascular bed deemed appropriate by supervising practitioner.
 - B. Controlled and selective infusion of practitioner-specified fluids, including thrombolytics, into the pulmonary arteries and/or peripheral vasculature.
- II. Contraindications for use includes but not limited to:
 - A. Patients in whom thrombolytic and/or anticoagulation therapy is contraindicated.
 - B. Any situation in which the medical judgment of the practitioner determines such a procedure may compromise the patient's condition.
- III. Equipment Preparation:

SUBJECT: ULTRASONIC ENDOVASCULAR SYSTEM FOR THROMBUS NRS Policy No. 850.00 Issue 1

MANAGEMENT Page 2 of 3

A. Follow the Manufacturer's Instructions for Use (IFU)

- IV. Order Set:
 - A. Obtain thrombolytic orders
- V. Device set-up, Insertion, Treatment Initiation and Trouble Shooting:
 - A. Follow the Manufacturer's Instructions for Use (IFU)
- XI. Device Removal
 - 1. Sheath removal by practitioner only
 - 2. Follow the IFU
- XII. Education
 - A. Registered Nurses with validated competency may perform monitoring and managing device.

REFERENCES: EkoSonic® Control Unit Instructions for Use OneSource Docs | Document

Database of IFUs and PM Service Manuals

DEFINITIONS: Control Unit (CU): The Control Unit (CU) is the main component that manages and regulates the operation of the entire system, providing power and user interface for

operator control.

Connector Interface Cable (CIC): The Connector Interface Cable (CIC) is the cable assembly that connects the Ultrasonic Assisted Infusion Catheter (IC) and

Ultrasonic Core (USC) to the Control Unit (CU).

<u>Ultrasonic Assisted</u> Infusion Catheter (IC): The <u>Ultrasonic Assisted</u> Infusion Catheter (IC) is a component used to deliver fluids, including thrombolytics, into the vasculature.

Ultrasonic Core (USC): The Ultrasonic Core is a part of the system that uses ultrasound to enhance the delivery and effectiveness of the infused fluids.

ULTRASONIC ENDOVASCULAR SYSTEM FOR THROMBUS SUBJECT: NRS Policy No. 850.00 Issue 1

MANAGEMENT

ATTACHMENTS: N/A

APPROVAL DATE: 9/18/2024 Gloria Steen, Interventional Radiology Charge Nurse

Department/Service Director, Manager or Supervisor

9/20/2024 **Nursing Standards Committee**

Applicable Administrator, Hospital or Medical Committee

Page 3 of 3

Patient Safety and Quality Committee
Applicable Administrator, Hospital or Medical Committee 9/25/2024

Quality Management Committee
Applicable Administrator, Hospital or Medical Committee 1/9/2025

1/23/2025 **Medical Executive Committee**

Applicable Administrator, Hospital or Medical Committee

9/9/2025 **Board of Supervisors**

Approved by the Governing Body

REPLACES: N/A

EFFECTIVE: 9/18/2024

REVISED: N/A

REVIEWED: N/A