

Model (Fig 1)	Deployed Funnel Diameter	Sheath Inner Diameter	Guidewire Compatibility	Usable Length
ClotTriever®	14 mm	13 Fr	0.035"	15 cm
Sheath (50-101)		(4.33 mm)	(0.9 mm)	

Model (Fig 2)	Guidewire Compat- ibility			Deployed Length (A)	•			Open Collection Bag Length (D)
ClotTriever® Catheter (40-102)	0.035"	11 Fr (3.67 mm)	80 cm	31 cm	24 cm	16 mm	45 mm	19 cm

Indications: The ClotTriever® system is indicated for (1) the non-surgical removal of soft emboli and thrombi from blood vessels, and (2) Injection, infusion and/or aspiration of contrast media and other fluids into or from a blood vessel. The ClotTriever® System is intended for use in the peripheral vasculature.

Extracting Large Clots from Large Vessels without the need for Thrombolytics or ICU stay.

Caution: Federal (USA) law restricts this device to sale by or on the order of a physician.

See Instructions for Use for complete Indications for Use, contraindications, warnings, and precautions.



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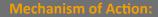
ClotTriever®

What is the ClotTriever®?

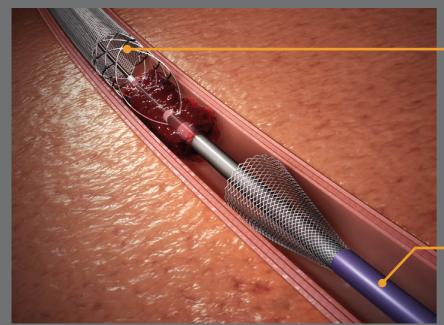
The ClotTriever® is a mechanical thrombectomy system designed to remove large clots from large vessels in a single session, without the need for thrombolytic drugs or consequent ICU stay.

Highlights & Clinical Benefits

- Immediate symptom improvement
- L Single session treatment
- Eliminates the need for thrombolytics and ICU stay
- Minimizes blood loss
- Avoids need for capital equipment



The ClotTriever® Catheter is a purely mechanical treatment option that liberates thrombus from the vessel walls and captures in a collection bag to be extracted via the ClotTriever® Sheath.



ClotTriever® Catheter

The ClotTriever® Catheter features a nitinol coring element and a braided collection bag, designed to core and collect clot for extraction from ClotTriever® Sheath.

ClotTriever® Sheath

The ClotTriever® Sheath features a self expanding nitinol mesh funnel to maximize clot removal. In addition, the sheath features a sideport for rapid aspiration and a hemostasis valve for catheter insertion.



CLOUT REGISTRY

ClotTriever® System

> 75%
Limbs with complete or near-complete

O
Device or Procedure
Related
Major-Adverse
Events

PROCEDURE

O DAYS

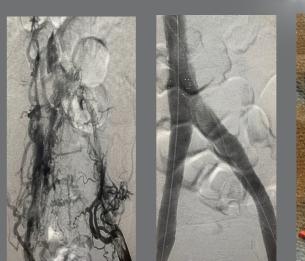
Median ICU
length of stay

30-DAY OUTCOMES 89% Patency

Initial CLOUT data (n = 50) demonstrates the ability of ClotTriever to successfully treat a range of clot chronicity in real world DVT patients in a single session, without the need for thrombolytic drugs.

Measure	Baseline	30-days	Δ	p-value
Villalta score	10	3	-7	< 0.01
PTS rate	93.9%	24.2%	-69.7%	< 0.01
Moderate/Severe PTS rate	51.5%	9.1%	-42.2%	< 0.01
rVCSS	6	3	-3	< 0.01
EQ-5D QoL	0.79	0.88	0.09	< 0.01
NPRS pain	3	0	-3	< 0.01

CLOUT is a 500-patient prospective, multicenter, single-arm registry evaluating real world patient outcomes after treatment of lower extremity DVT with ClotTriever. Three-quarters of the first 50 patients had clot older than two weeks and over half had failed other therapies before treatment. This is truly a prospective look at the "real world" DVT patient population. Nonetheless, ClotTriever still successfully removed nearly all the clot in over three-quarters of these patients resulting in improvements to disease severity and quality of life.





What types of DVT does the ClotTriever® treat?

Iliofemoral or Femoral-Popliteal DVT (or any vessel 6-16mm in diameter) for ClotTriever®.

The mechanism of clot removal for ClotTriever allows for extraction of a wide range of clot chronicity.

