



FUNNEL 7 mm diameter, 6.5 cm long nitinol funnel to capture baskets and entrain clot

Capture tibial clot -on the spot.

NEW! The **Pounce[™] LP Thrombectomy**

System is designed to remove arterial clot in **BTK** vessels 2mm–4mm in diameter.

Schedule a Product Demonstration

pouncesystem.com/jointhehunt



BASKETS

Dual **4 mm** diameter nitinol self-expanding baskets mounted in series on a **.020**" diameter core wire for capturing thrombus



DELIVERY CATHETER

.018" Guidewire compatible 150 cm Working length



42 cm of Serene[™] Hydrophilic coating on distal end of catheter for improved trackability

Successful Treatment of Infrapopliteal Arterial Thrombus

Lucas Ferrer Cardona, MD Vascular Surgeon Dell Seton Medical Center The University of Texas Hospital Austin, Texas

- Initial angiography showed complete thrombosis of left popliteal, TPT, and tibial arteries
- Pounce[™] MD Thrombectomy System cleared organized clot from popliteal and TPT arteries
- Pounce[™] LP Thrombectomy System cleared subacute clot from posterior tibial and peroneal arteries, restoring flow to plantar arch





√PSURMODICS

How It Works

The **Pounce[™] LP Thrombectomy System** is comprised of three components:

a delivery catheter, a basket wire, and a funnel catheter.

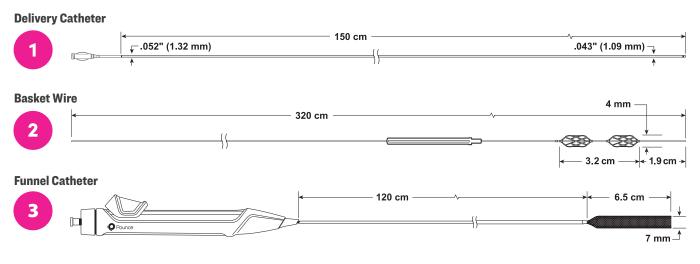


The basket wire is delivered distal to the location of the thrombus, deploying two nitinol self-expanding baskets.

The baskets capture the clot and are retracted into a nitinol collection funnel.

With the clot entrained, the system is retracted into a minimum 7 Fr guide sheath through which the clot is withdrawn and removed from the body.

Product Specifications and Ordering Information



Model Number	Sheath Compatibility	Guidewire Compatibility	Working Length	Basket Diameter	Funnel Diameter
PTS-0407-7F150	≥ 7 Fr	0.018" (0.457 mm)	150 cm	4 mm	7 mm

INDICATION FOR USE / INTENDED USE

- The Pounce Thrombectomy System is intended for the non-surgical removal of thrombi and emboli from the peripheral arterial vasculature.
- The Pounce LP Thrombectomy System is indicated for use in vessels ranging from 2 mm to 4 mm in diameter.

CONTRAINDICATIONS

- The device is not intended for venous applications.
- The device is not intended for peripheral vasculature dilatation.
- The device is not for coronary or neurovascular use.
- The device is contraindicated for use in patients who cannot receive recommended intravenous anticoagulant therapy.
- The safety and effectiveness of the device has not been established in pediatric patients (<18 years of age).
- The device is not intended to be deployed in vessels with previously implanted devices.

Caution: Federal (US) law restricts these device(s) to sale by or on the order of a physician. Please refer to Instructions for Use for indications, contraindications, warnings, and precautions.

Also available to treat peripheral arteries ranging from **3.5mm**-**6mm** in diameter.



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