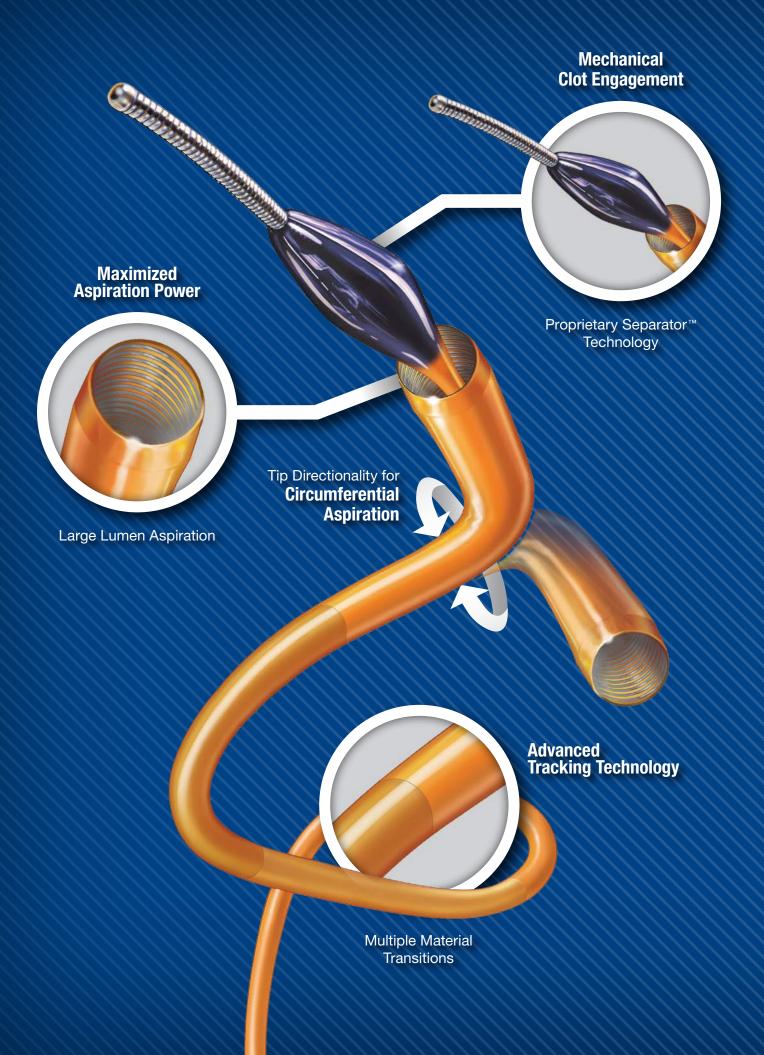
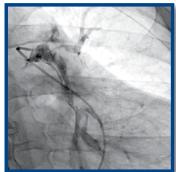


Penumbra (P)





Arterial and venous clot in bigger vessels where circumferential aspiration is desired.

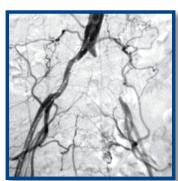


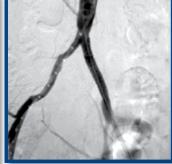


Extraction of Thrombus from Pulmonary ArteryDrs. Alex Powell & James Benenati, Miami Cardiac and Vascular Institute, FL



Arterial clot in larger arteries including: SFA, popliteal and viscerals.

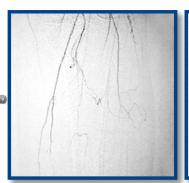


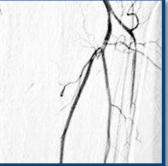


Extraction of Thrombus from Common Iliac ArteryDr. Paul Perkowski, Baton Rouge General, LA

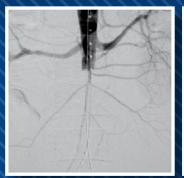


Smaller, distal reach for upper and lower arterial extremities cases.



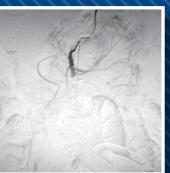


Recanalization of TrifurcationDr. Matthew Loe, St. Paul Radiology, MN





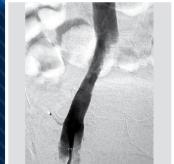
Recanalization of Occluded Aorta Dr. Frank Arko, Sanger Heart and Vascular Institute, NC



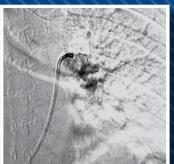


Removal of SMA Embolus Drs. Rahul Patel & Robert Lookstein, Mt. Sinai Hospital, NY





Extraction of Thrombus from Iliac Vein Dr. Lawrence Whitney, Lakeland Regional Medical Center, FL



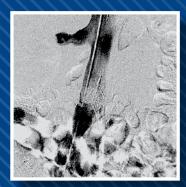


Extraction of Thrombus from Pulmonary Artery Dr. Ezell Askew Jr., St. Joseph's Medical Center of Stockton, CA



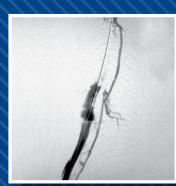


Recanalization of Thrombosed SFA Stent Dr. George Adams, Rex UNC Health Care, NC



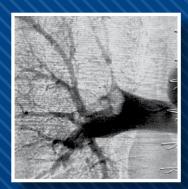


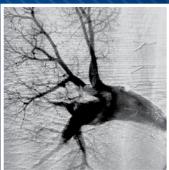
Extraction of Thrombus from IVC Filter Dr. Jonathan Ha, Hunter Holmes McGuire Virginia Medical Center, VA





Extraction of Thrombus from Iliac Vein Dr. Mahmood Razavi, St. Joseph Hospital, Orange, CA

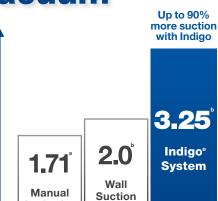




Extraction of Thrombus from Pulmonary Artery Dr. Paul Perkowski, Baton Rouge General, LA

Full Vacuum[™]

Flow Rate (cc/sec)

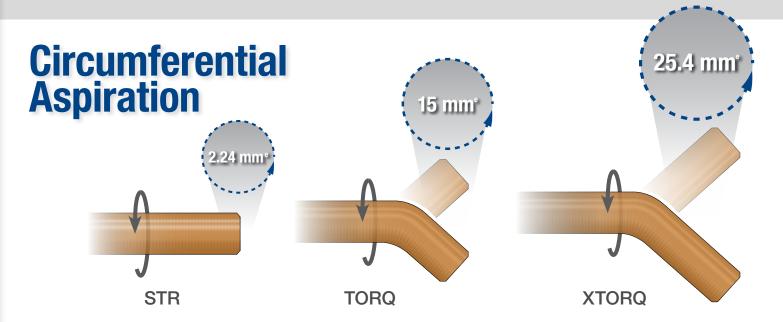




More Aspiration

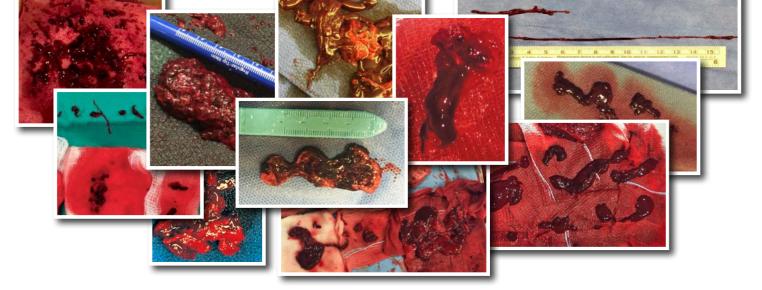
Aspiration Catheters

Aspiration Volume (mL in 20 sec)° 180 -160 140 -120 100 -56 14 AngioJet CAT3 CAT5 CAT6 CAT8 AngioJet™



- a. Vascular Solutions, Inc. Comparison of dimensions and aspiration rate of the Pronto® V3, Pronto® LP, Export® XT, Export® AP, Fetch®, Xtract™, Diver C.E.™ and QuickCat™ catheters. http://www.aquilantinterventional.com/assets/aquilantinterventional/Products/brochures/871027/ML1623_rev_F_Pronto_V3_Extraction_Lumen_Comparison.pdf. Published December 2009, Accessed March 13, 2014.
- b. Data on file at Penumbra, Inc. based on testing with CAT5. c. Data on file at Penumbra, Inc. d. Data on file at Penumbra, Inc. e. Data on file at Penumbra, Inc.

ZelanteDVT™



Ordering Information

Catalog Number	Description	Proximal OD	Distal OD	Length	Compatible Penumbra Devices
Indigo® Catheters				Working	
CAT8XTORQ115	Indigo 8 XTorq Tip	8.0 F	8.0 F	115 cm	Separator™ 8
CAT8TORQ85	Indigo 8 Torq Tip	8.0 F	8.0 F	85 cm	Separator 8
CAT8STR85	Indigo 8 Straight Tip	8.0 F	8.0 F	85 cm	Separator 8
CAT6	Indigo 6	6.0 F	6.0 F	135 cm	Separator 6
CAT5	Indigo 5	6.0 F	5.0 F	132 cm	Separator 5
CAT3	Indigo 3	4.1 F	3.4 F	150 cm	Separator 3
Indigo Separators				Total	
SEP8	Separator 8	n/a	.072"	150 cm	CAT8
SEP6	Separator 6	n/a	.055"	175 cm	CAT6
SEP5	Separator 5	n/a	.045"	175 cm	CAT5
SEP3	Separator 3	n/a	.028"	190 cm	CAT3
Accessories					
PMX110	Pump MAX ™				MAX Canister
IAPS2	MAX Canister				Pump MAX
IST3	Large Lumen Aspiration Tubing	g			All Indigo Catheters

Indigo® Aspiration Catheters and Separators – Indication For Use
As part of the Indigo® Aspiration System, the Indigo Aspiration Catheters and Separators are indicated for the removal of fresh, soft emboli and thrombi from vessels of the peripheral arterial and venous systems.

Indigo Aspiration Tubing - Indication For Use

As part of the Indigo Aspiration System, the Indigo Sterile Aspiration Tubing is indicated to connect the Indigo Aspiration Catheters to the Penumbra Pump MAX™.

Contraindications

Not for use in the coronaries or the neurovasculature.

- The Indigo Aspiration System should only be used by physicians who have received appropriate training in interventional techniques.

 Do not advance, retract or use any component of the Indigo System against resistance without careful assessment of the cause using fluoroscopy. If the cause cannot be determined, withdraw the device or system as a unit. Unrestrained torquing or forced insertion of the catheter or separator against resistance may result in damage to the device or vessel.
- Do not use the Indigo Aspiration System with a pump other than the Penumbra Aspiration Pump

- The device is intended for single use only. Do not resterilize or reuse. Resterilization and/or Reuse may result in ineffective catheter coating

- The device is intended for single use only. Do not resterlize or reuse. Resterlization and/or Reuse may result in ineffective catheter coating lubrication, which may result in high friction and the inability to access the target vasculature location.
 Do not use kinked or damaged devices. Do not use opened or damaged packages. Return all damaged devices and packaging to the manufacturer/distributor.
 Use prior to the "Use By" date.
 Use the Indigo Aspiration System in conjunction with fluoroscopic visualization.
 When performing aspiration ensure the Indigo Aspiration Tubing valve is open for only the minimum time needed to remove thrombus. Excessive aspiration or failure to close the Indigo Aspiration Tubing valve when aspiration is complete is not recommended.
 The Indigo Separator" is not intended for use as a neurovascular guidewire. If repositioning of the Indigo Aspiration Catheter is necessary during the revascularization procedure, such repositioning should be performed over an appropriate guidewire using standard microcatheter and quidewire techniques. catheter and quidewire techniques.
- Do not use automated high-pressure contrast injection equipment with the Indigo Aspiration Catheter because it may damage the device.

Indigo Potential Adverse Events

Possible complications include, but are not limited to, the following: allergic reaction and anaphylaxis from contrast media, acute occlusion, air embolism, arteriovenous fistula, death, device malfunction, distal embolization, emboli, false aneurysm formation, hematoma or hemorrhage at access site, inability to completely remove thrombus, infection, hemorrhage, schemia, kidney damage from contrast media, neurological deficits including stroke, vessel spasm, thrombosis, dissection, or perforation, intimal disruption, myocardial infarction, emergent surgery, fibrillation, hypotension, respiratory failure, peripheral thromboembolic events.

Pump MAX™

Indication For Use

The Penumbra Pump MAX™ is indicated as a vacuum source for Penumbra Aspiration Systems.

Contraindications

There are no known contraindications.

Warnings/Precautions

- The canister/tubing is intended for single use only. Do not reuse. Reuse may result in the inability to aspirate.
 Do not block bottom or back air vents. Unit may overheat and shut off or fail to restart if run for extended periods without airflow.
- without arritow.

 To avoid risk of electric shock, this equipment must only be connected to a supply mains with protective earth.

 Do not position the pump so that it is difficult to operate the power cord disconnection device.

 Remove and service the pump if liquids or solids have been drawn into the vacuum pump.

 Do not use in the presence of flammable anaesthetic mixture with air or nitrous oxide.

 Do not use in oxygen rich environment.

 To prevent fire or shock hazard, use replacement fuses of equal size and rating.

- To prevent fire or shock hazard, use a replacement power cord of equal rating.
 To prevent fire or shock hazard, use a replacement power cord of equal rating.
 To not re-influse blood or fluid from the canister back into the patient.
 To not use petroleum base compounds, acids, caustics, or chlorinated solvents to clean or lubricate any parts. It will reduce service life of the pump. Use only water-base solvents for cleaning.
- Federal law (USA) restricts this device to sale by or on the order of a physician.
 No modification of this equipment is allowed.



www.penumbrainc.com

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Penumbra Neuro Australia Pty Ltd Suite 3, Level 5, 1 Oxford Street Darlinghurst NSW 2010 Australia T +61-1300 817 025 F +61-1300 817 026 order.anz@penumbrainc.com

Product availability varies by country. Images used with permission. Consents on file at Penumbra, Inc. Caution: Federal (USA) law restricts this device to sale by or on the order of a physician. Prior to use, please refer to the Instructions for Use for the Indigo Aspiration System and Pump MAX for complete product indications, contraindications, warnings, precautions, potential adverse events and detailed instructions for use. Please contact your local