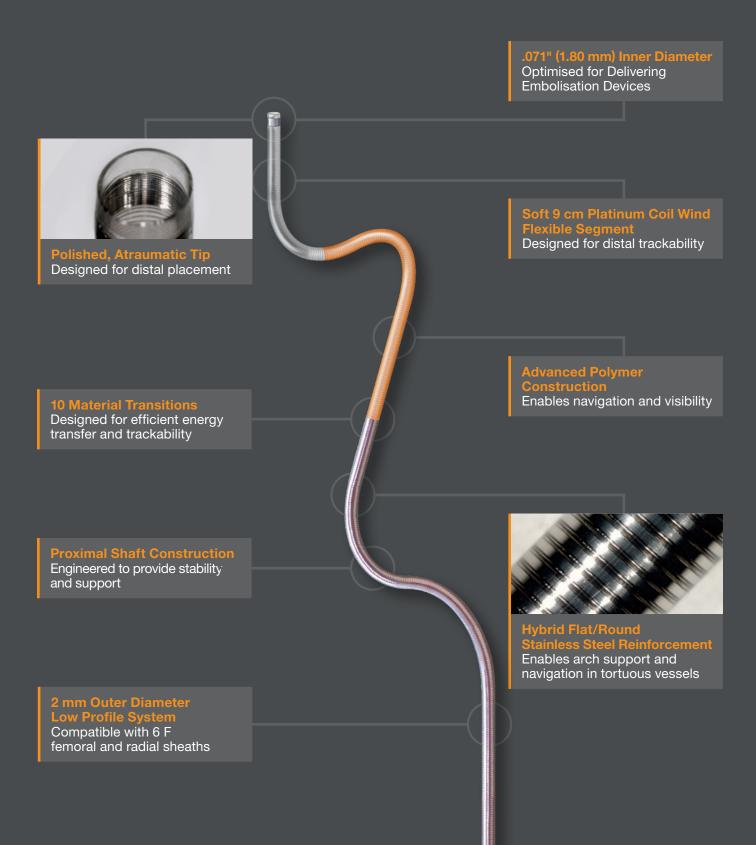






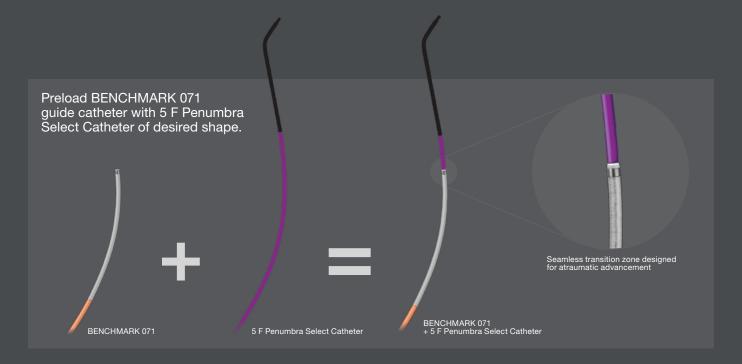
BENCHMARK 071Key Design Features



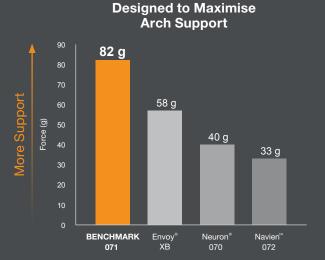
BENCHMARK 071 packaged with 5 F Penumbra Select Catheter

Advantages of Rapid Primary Access

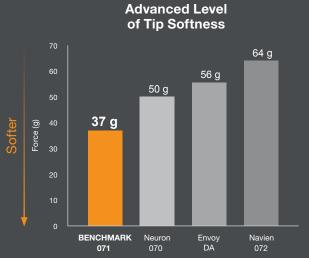
- Faster procedures no need for an exchange
- Allows easy selection off of arch into desired vessel
- Facilitates atraumatic placement into distal vasculature
- Can be used for diagnostic angiogram .040" (1.02 mm) lumen
- Compatible with .035"-.038" (.89 mm-.97 mm) guidewires



Performance Testing



Buckling Test: Higher values correspond with more proximal support^a



Deflection Test: Lower values correspond to softer distal tip^b

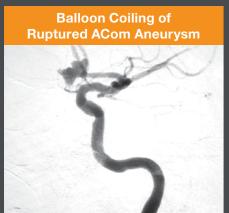
Femoral Access, Defined

Designed for stability and compatibility with atraumatic navigation

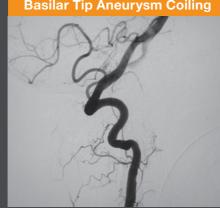
Standalone or compatible with 6 F long sheath



BENCHMARK 071 Intracranial Access via Femoral Approach





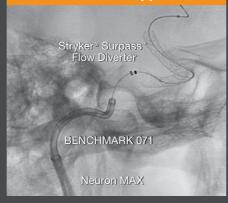




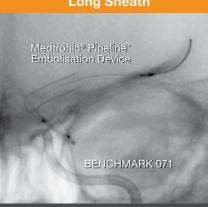
Full distal shaft

Images courtesy Drs. Yasha Kayan and Josser Delgado Abbott Northwestern Hospital, Minnesota, USA

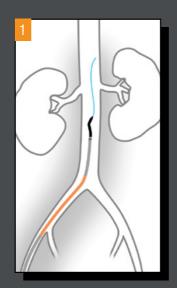
Compatible with Neuron MAX® for Added Support



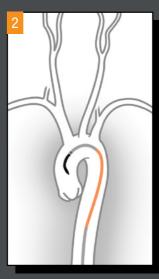
Stability Without Long Sheath



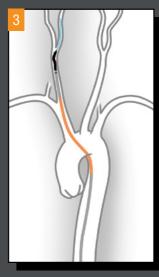
Rapid Primary Access — Typical Approach via Femoral Artery



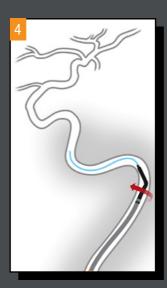
Introduce the preloaded system over a guidewire and advance to a straight section of the abdominal aorta



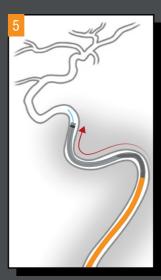
Advance the tip of 5 F Select over the guidewire into the ascending aorta while maintaining the position of BENCHMARK 071



Advance 5 F Select and BENCHMARK 071 over the guidewire into the internal carotid artery



With sufficient wire purchase, torque 5 F Select towards the first major turn



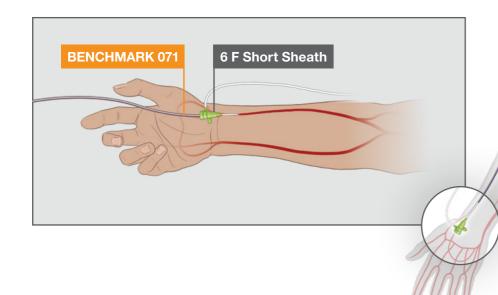
While maintaining position of 5 F Select and guidewire, advance BENCHMARK 071 into desired position



Remove 5 F Select while holding BENCHMARK 071 in position

Radial Advantage

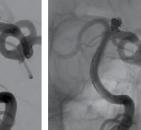
Low profile 2 mm OD compatible with the typical radial artery



Hybrid stainless steel reinforcement engineered to provide kink resistance in aortic arch



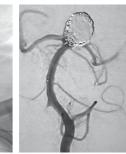




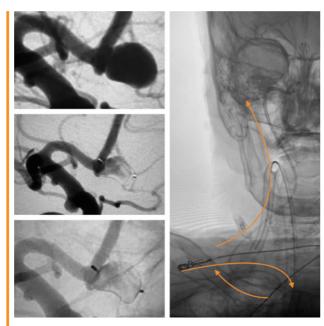
Stent-Assisted Coiling Dr. Levansri Makalanda The Royal London Hospital London, UK







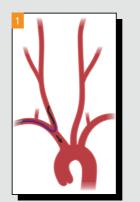
Primary Coiling Dr. Jean Delbrune Northside Hospital Florida, USA



MicroVention® WEB® Embolisation Dr. Justin Singer Spectrum Health

Michigan, USA Images used with permission. Consents on file at Penumbra, Inc. Case examples presented are for informational purposes only. Results may not be predictive for all patients and may vary based on patient-specific attributes and other factors.

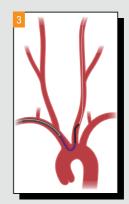
Rapid Primary Access — Typical Approach via Radial Artery



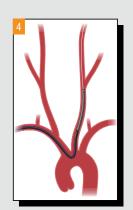
With BENCHMARK 071 and 5 F SIM Select in right subclavian artery, 5 F SIM Select is advanced forward into aortic arch



BENCHMARK 071 is moved forward over 5 F SIM Select in right subclavian artery, 5 F SIM Select is



.035" wire is moved through 5 F SIM Select into left ICA



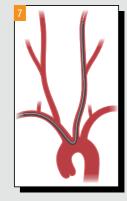
place, BENCHMARK 071 is tracked into CCA



5 F SIM Select is removed, while BENCHMARK 071 and .035" wire stay in CCA/ICA



Switch to 5 F BER Select, which is advanced into ICA while BENCHMARK 071 and .035" wire stay in CCA/ICA



over BER Select catheter into ICA, then 5 F Select is removed

Procedural and operative techniques and considerations are illustrative examples from physician experience. Physicians' treatment and technique decisions will vary based on their medical judgment.

Data Presented at SNIS 2018

Radial Access for Cerebrovascular Intervention Using Penumbra BENCHMARK 071 Guiding Catheter[®]

36	Patients
10	Primary Coiling
8	Stent-Assisted Coiling
5	Flow-Diverter Embolisation
2	Balloon-Assisted Coiling
2	Wingspan™ Stent-Assisted Coiling
2	AVM or Dural AV Fistula
2	Vessel Sacrifice
1	Subclavian Stent
1	Vasospasm Treatment
3	Other
o Cotti C Doot	oni C. Edan T. et al. E. 021 Padial access for carebravascular intervention

Key Results

No catheter-related complications

No major radial access site complications

BENCHMARK 071 Kits

		BENCHMARK 071		5 F Select™ Catheter	
Catalog Number	Description	Length (cm)	Shape	Length (cm)	Shape
BMK6F95BER120	BENCHMARK 071 KIT	95	Straight	120	BER
BMK6F95MBER120	BENCHMARK 071 KIT	95	MP	120	BER
BMK6F105BER130	BENCHMARK 071 KIT	105	Straight	130	BER
BMK6F105MBER130	BENCHMARK 071 KIT	105	MP	130	BER

BENCHMARK 071

		BENCHM	IARK 071
Catalog Number	Description	Length (cm)	Shape
BMK6F95	BENCHMARK 071	95	Straight
BMK6F95M	BENCHMARK 071	95	MP
BMK6F105	BENCHMARK 071	105	Straight
BMK6F105M	BENCHMARK 071	105	MP
BMK6F115	BENCHMARK 071	115	Straight
BMK6F115M	BENCHMARK 071	115	MP

Tip Shapes

BENCHMARK 071	Select Catheter		
Straight MP	H1 BER SIM		

5 F Select Catheters

Catalog Number	Description	Working Length (cm)	Inner Diameter (in / mm)	Wire Compatibility (in / mm)	Shape
PNS5F120BER	5F Select Catheter	120	.040 (1.02)	.035038 (.8997)	BER
PNS5F130BER	5F Select Catheter	130	.040 (1.02)	.035038 (.8997)	BER
PNS5F130SIM	5F Select Catheter	130	.040 (1.02)	.035038 (.8997)	SIM
PNS5F120H1	5F Select Catheter	120	.040 (1.02)	.035038 (.8997)	H1
PNS5F130H1	5F Select Catheter	130	.040 (1.02)	.035038 (.8997)	H1

Prior to use, please refer to the Instructions for Use for complete product indications, contraindications, warnings, precautions, potential adverse events, and detailed instructions for use.

BENCHMARK Intracranial Access System -

The BENCHMARK Intracranial Access System is intended for the introduction of interventional devices into the peripheral, coronary, and neuro vasculature.

Potential Adverse Events
Possible complications include, but are not limited to, the following: acute occlusion; air embolism;

death; distal embolization; emboli; false aneurysm formation; hematoma or hemorrhage at puncture site; infection; intracranial hemorrhage; ischemia; neurological deficits including stroke; vessel spasm, thrombosis, dissection, or perforation.

NEURON MAX System – Intended Use The NEURON MAX System is intended for the introduction of interventional devices into the

peripheral, coronary, and neuro vasculature.

Potential Adverse Events
Possible complications include, but are not limited to, the following: acute occlusion; air embolism;

death; distal embolization; emboli; false aneurysm formation; hematoma or hemorrhage at puncture site; infection; intracranial hemorrhage; ischemia; neurological deficits including stroke; vessel spasm, thrombosis, dissection, or perforation.

NEURON Intracranial Access System – Intended Use The NEURON Intracranial Access System is intended for the introduction of interventional devices into the peripheral, coronary, and neuro vasculature. Potential Adverse Events

Possible complications include, but are not limited to, the following: acute occlusion; air embolism;

death; distal embolization; emboli; false aneurysm formation; hematoma or hemorrhage at puncture site; infection; intracranial hemorrhage; ischemia; neurological deficits including stroke; vessel spasm, thrombosis, dissection, or perforation.



One Penumbra Place Alameda, CA 94502 IISA 1.888.272.4606 T 1.510.748.3200 F 1.510.748.3232 order@penumbrainc.com info@penumbrainc.com

Penumbra Europe GmbH Am Borsigturm 4413507 Berlin Germany T +49 30 2005 676-0 F +49 30 2005 676-10 de-order@penumbrainc.com info@penumbrainc.de

Penumbra Neuro Penumbra weuro
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

Penumbra Latin America Distribuidora de Equipamentos e Produtos Médicos Ltda Avenida Brigadeiro Luís Antônio 3421 cj 201 CEP 01401-001 São Paulo, Brazil T 5511.2883.5825 order.la@penumbrainc.com

Prior to use, please refer to the Instructions for Use for complete product indications, contraindications, warnings, precautions, potential adverse events, and detailed instructions for use. Renderings for illustrative purposes only. Images used with permission. Consents on file at Penumbra, Inc. Please contact your local Penumbra representative for more information.