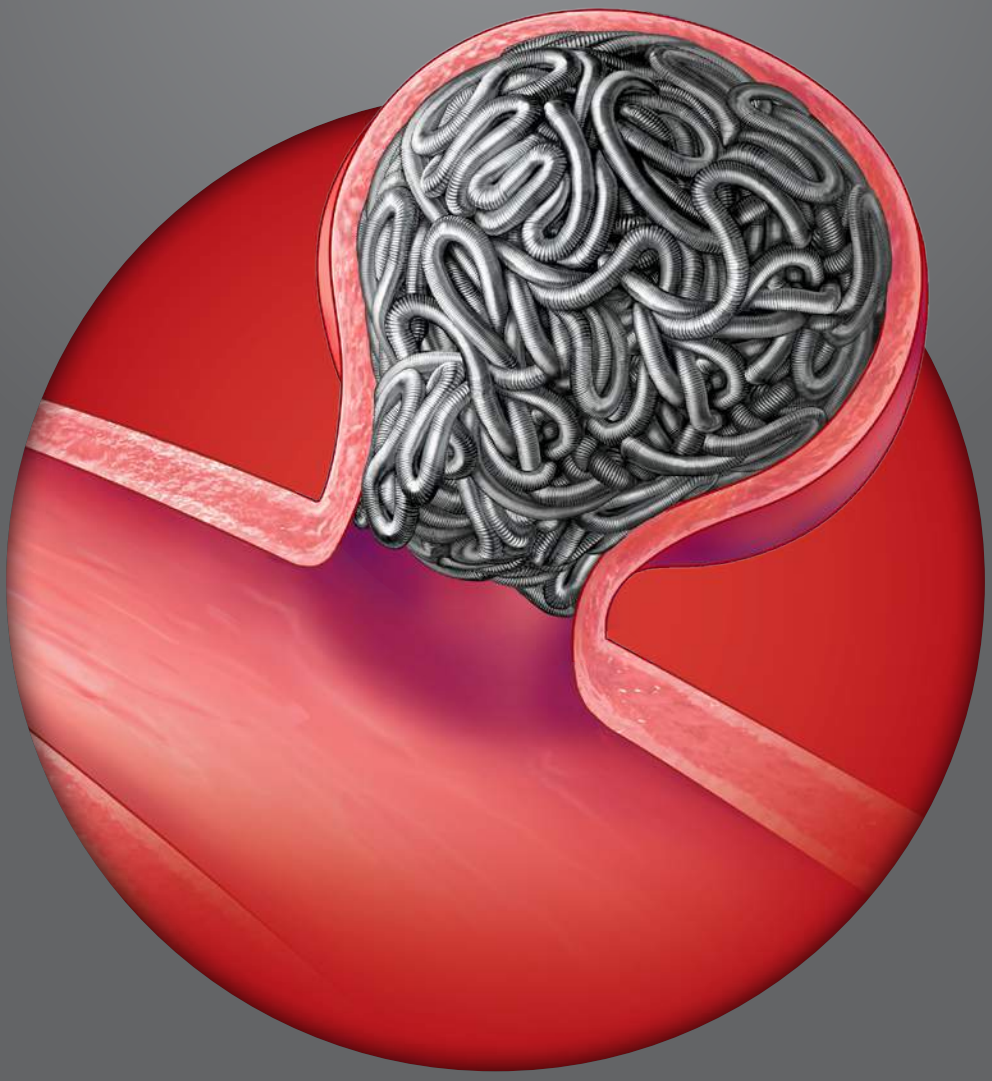
Ruby® Coil

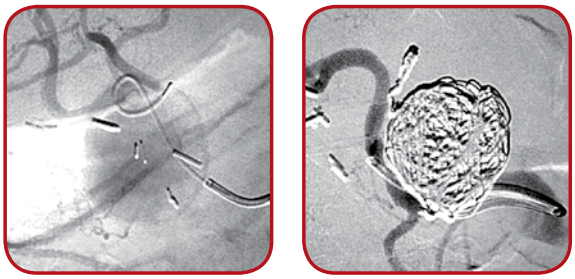
Large Volume Detachable Coils



Penumbra®

Ruby® Case Examples

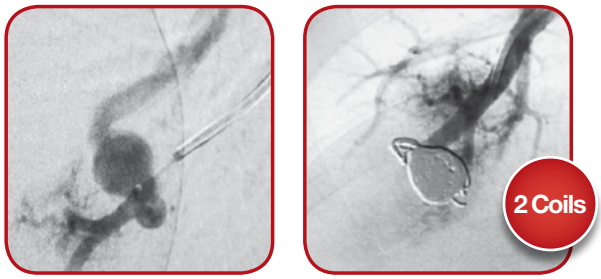
38 mm Hepatic Artery Aneurysm



Pre-Treatment | Post-Treatment

Dr. James Benenati
Miami Cardiac and Vascular Institute, FL

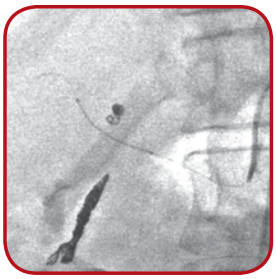
Pulmonary AVM



Pre-Treatment | Post-Treatment

Dr. J Moskovitz
Florida Hospital, Orlando, FL

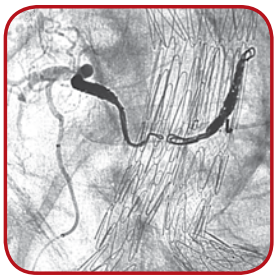
Y90 Embolization



Post-Treatment

Dr. Dmitri Samoilov
Medical Center Radiologists, VA

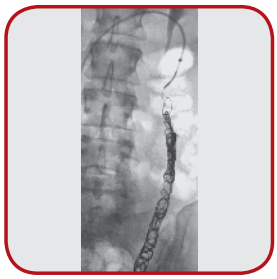
Type 2 Endoleak



Post-Treatment

Dr. Corey Teigen
Sanford Health, ND

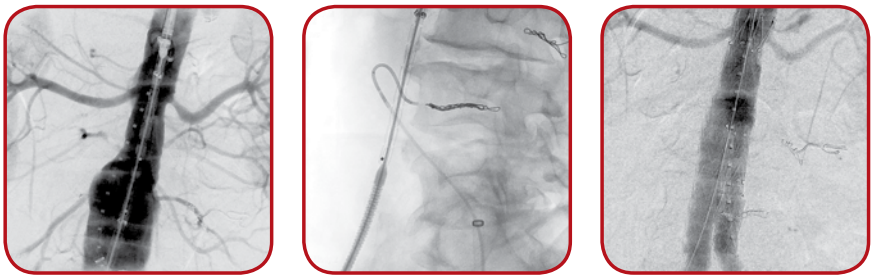
Ovarian Vein Embolization



Post-Treatment

Dr. Olga Brook
Beth Israel Deaconess Medical Center, MA

Type 2 Endoleak



Pre-Treatment | Intra-Operative | Post-Treatment

Dr. Frank Arko
Sanger Heart and Vascular Institute, NC

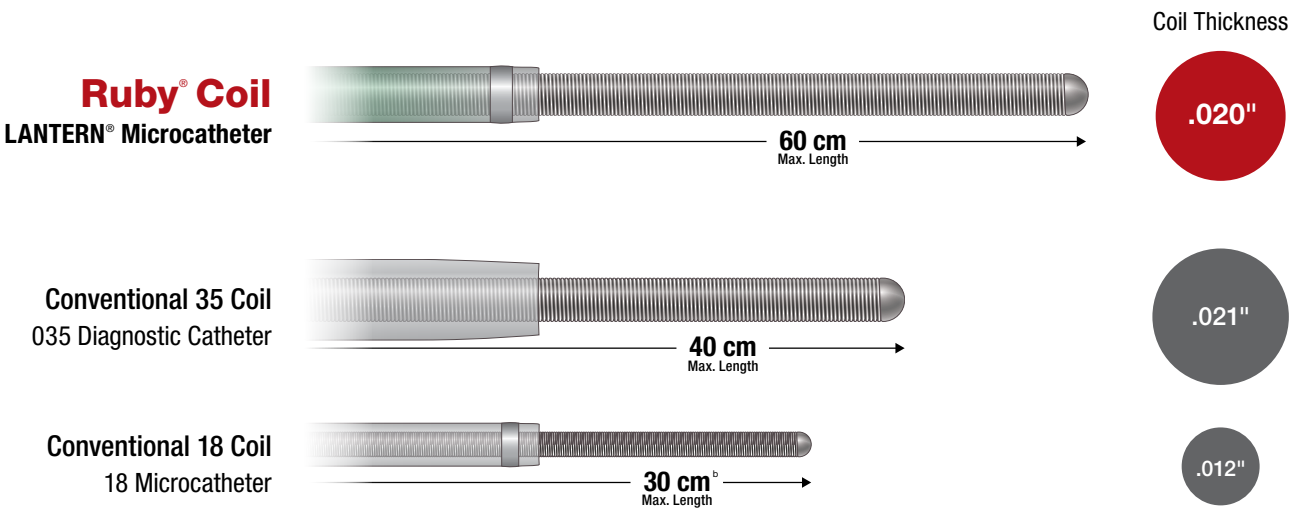
Volume Advantage

30 cm Ruby® Coil

30 cm Conventional Detachable Coil

Reduces the number of coils and procedure time by maximizing length and coil thickness.[a]

Only Coil to Provide 60 cm Length and .020" Thickness



a. Data on file at Penumbra, Inc.
b. Based on 8 mm coil.

Large Volume Detachable Coils

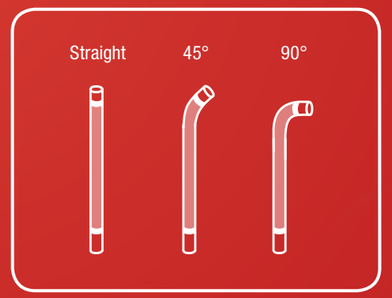
Standard
32 mm × 60 cm

Soft
2 mm × 1 cm

EXTREME DELIVERABILITY

Only coil designed to be delivered through a .025–.027" high-flow microcatheter.

Tip Shapes



LANTERN®
Microcatheter

Low Profile
2.6 F Distal Shaft Enables Access to Small Tortuous Vasculature

High-Flow
Large Lumen for Optimal Contrast Injections

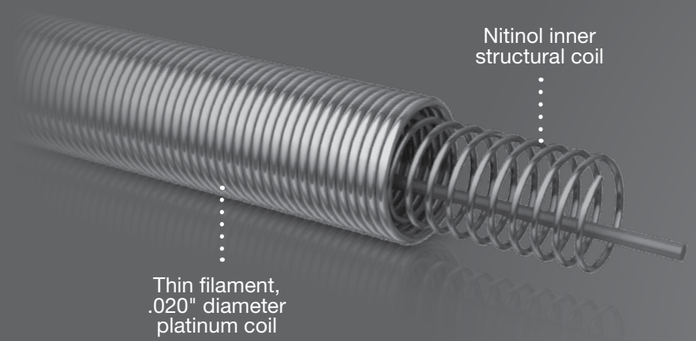
Increased Visibility
Provides Confident Placement

Dual Markers
Facilitates Precise Coil Deployment

8 Transition Zones
for Advanced Trackability

SOFTNESS

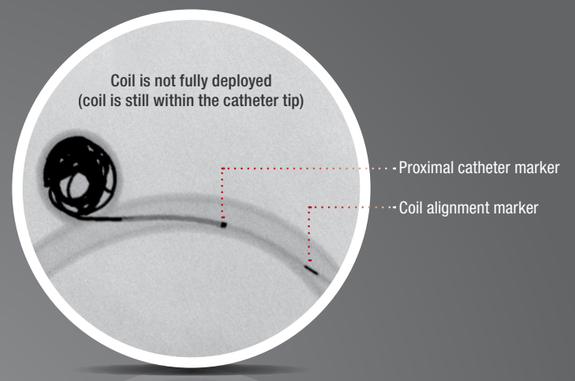
Choice of Standard and Soft coils provide superior packing density while avoiding catheter kick-out.



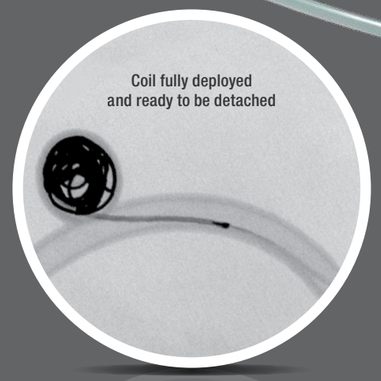
Unique Tri-Layer Coil Design

MECHANICAL DETACHMENT

Ruby® Coil deployed from LANTERN® High-Flow Microcatheter



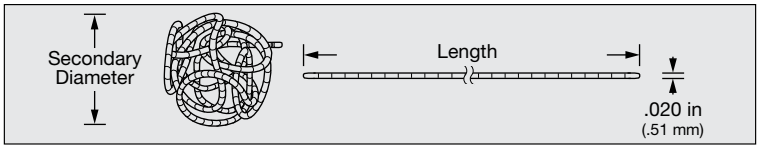
Ruby® Coil is fully retractable and resheathable, allowing user to achieve ideal placement in the target vessel.



Coil alignment marker passes proximal catheter marker to form “T,” indicating that Ruby Coil can now be detached.

Ordering Information

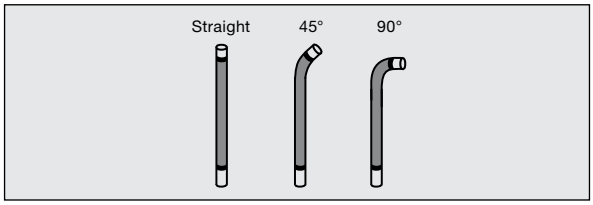
Ruby® Coil



Catalog Number	Secondary Diameter (mm)	Length (cm)
Frame – COMPLEX **STANDARD**		
RBY2C0305	3	5
RBY2C0312	3	12
RBY2C0320	3	20
RBY2C0410	4	10
RBY2C0415	4	15
RBY2C0420	4	20
RBY2C0430	4	30
RBY2C0435	4	35
RBY2C0512	5	12
RBY2C0520	5	20
RBY2C0530	5	30
RBY2C0620	6	20
RBY2C0630	6	30
RBY2C0725	7	25
RBY2C0825	8	25
RBY2C0840	8	40
RBY2C0930	9	30
RBY2C1035	10	35
RBY2C1240	12	40
RBY2C1260	12	60
RBY2C1460	14	60

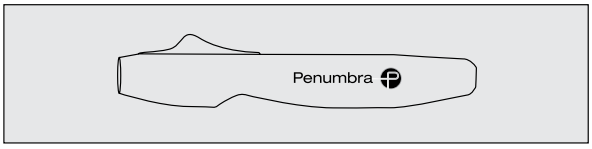
Catalog Number	Secondary Diameter (mm)	Length (cm)
Frame – COMPLEX **STANDARD** (cont.)		
RBY2C1660	16	60
RBY2C1857	18	57
RBY2C2060	20	60
RBY2C2457	24	57
RBY2C2860	28	60
RBY2C3260	32	60
Fill – COMPLEX **SOFT**		
RBY4C0201	2	1
RBY4C0202	2	2
RBY4C0204	2	4
RBY4C0305	3	5
RBY4C0315	3	15
RBY4C0406	4	6
RBY4C0415	4	15
RBY4C0620	6	20
RBY4C0630	6	30
RBY4C0835	8	35
RBY4C0860	8	60
RBY4C1650	16	50
RBY4C2060	20	60

LANTERN® Delivery Microcatheter



Catalog Number	Tip Shape	Length (cm)	ID	OD Distal / Proximal
PXSLIMLAN115STR	Straight Tip	115	.025"	2.6F / 2.95F
PXSLIMLAN115T45	45° Tip			
PXSLIMLAN115T90	90° Tip			
PXSLIMLAN135STR	Straight Tip	135	.025"	2.6F / 2.95F
PXSLIMLAN135T45	45° Tip			
PXSLIMLAN135T90	90° Tip			
PXSLIMLAN150STR	Straight Tip	150	.025"	2.6F / 2.95F
PXSLIMLAN150T45	45° Tip			
PXSLIMLAN150T90	90° Tip			

Ruby® Coil Detachment Handle



Catalog Number	Description
RH1	Detachment Handle

Indication For Use
The Ruby Coil System is indicated for arterial and venous embolizations in the peripheral vasculature.

Contraindications
There are no known contraindications.

Warnings
The Ruby Coil System should only be used by physicians who have received appropriate training in interventional techniques.

Precautions
- The device is intended for single use only. Do not resterilize or reuse. Resterilization and/or reuse may compromise the structural integrity of the device or increase the risk of contamination or infection leading to device failure and/or cross-infection and potential patient injury, illness or death.
- 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 device in conjunction with fluoroscopic guidance
- Do not advance or withdraw the device against resitance without careful assessment of the cause using fluoroscopy
- Moving or torquing the device against resistance may result in damage to the vessel or device
- Maintain a constant infusion of an appropriate flush solution

Potential Adverse Events
Potential complications include but are not limited to: Acute occlusion, air embolism, death, distal embolization, emboli, false aneurysm formation, hematoma or hemorrhage at puncture/access site/site of entry, infection, intracranial hemorrhage, ischemia, neurological deficits including stroke, vessel spasm, thrombosis, dissection or perforation, thrombo-embolic episodes, neurological deficits including stroke and possibly death, vascular thrombosis, post-embolization syndrome. revascularization, recanalization, inadequate occlusion, aneurysm rupture, parent artery occlusion, incomplete aneurysm filling

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Indication For Use
The Penumbra Delivery Microcatheters are indicated to assist in the delivery of diagnostic agents, such as contrast media, and therapeutic devices, such as occlusion coils, to the peripheral and neuro vasculature.

Contraindications
There are no known contraindications.

Warnings
The Penumbra Delivery Microcatheters should only be used by physicians who have received appropriate training in interventional techniques.

Precautions
- The devices are intended for single use only. Do not resterilize or reuse. Resterilization and/or reuse may result in ineffective catheter coating lubrication, which may result in high friction and the inability to access the target 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 Penumbra Delivery Microcatheters in conjunction with fluoroscopic visualization
- Do not advance or withdraw the Penumbra Delivery Microcatheters against resistance without careful assessment of the cause using fluoroscopy. If the cause cannot be determined, withdraw the device. Moving or torquing the device against resistance may result in damage to the vessel or device.
- Maintain a constant infusion of appropriate flush solution
- If flow through the device becomes restricted, do not attempt to clear the lumen by infusion. Remove and replace the device.

Potential Adverse Events
Possible complications include but are not limited to, the following: Acute occlusion, hematoma or hemorrhage at access site, death, intracranial hemorrhage, hemorrhage, infection (at access site), distal embolization, ischemia (cardiac and/or cerebral), embolus (air, foreign body, thrombus, plaque), aneurysm perforation, false aneurysm formation, neurological deficits including stroke, vessel spasm, thrombosis, dissection, perforation or rupture, air embolism, emboli

Penumbra

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6728, Rev. D 05/17 USA