### Venovo<sup>™</sup> Venous Stent System

Dia. (mm)	Length (mm)	Sheath Size (F)	80 cm Catheter Length	120 cm Catheter Length
10	40	8	VENUM10040	U VENUL10040
	60	8	□ VENUM10060	□ VENUL10060
	80	8	UENUM10080	U VENUL10080
	100	8	□ VENUM10100	□ VENUL10100
	120	8	□ VENUM10120	□ VENUL10120
	140	8	□ VENUM10140	□ VENUL10140
	160	8	□ VENUM10160	U VENUL10160
	40	8	□ VENUM12040	U VENUL12040
	60	8	□ VENUM12060	U VENUL12060
	80	8	□ VENUM12080	□ VENUL12080
12	100	8	□ VENUM12100	U VENUL12100
	120	8	□ VENUM12120	U VENUL12120
	140	8	□ VENUM12140	UVENUL12140
	160	8	□ VENUM12160	□ VENUL12160
14	40	9	U VENUM14040	VENUL14040
	60	9	□ VENUM14060	□ VENUL14060
	80	9	UENUM14080	U VENUL14080
	100	9	□ VENUM14100	□ VENUL14100
	120	9	□ VENUM14120	U VENUL14120
	140	9	□ VENUM14140	U VENUL14140
	160	9	U VENUM14160	U VENUL14160

Venovo<sup>™</sup> Venous Stent System

Dia. mm)	Length (mm)	Sheath Size (F)	80 cm Catheter Length	120 cm Catheter Length
16	40	10	□ VENUM16040	□ VENUL16040
	60	10	VENUM16060	U VENUL16060
	80	10	□ VENUM16080	□ VENUL16080
	100	10	VENUM16100	U VENUL16100
	120	10	□ VENUM16120	□ VENUL16120
	140	10	VENUM16140	U VENUL16140
	160	10	□ VENUM16160	□ VENUL16160
	40	10	UENUM18040	U VENUL18040
	60	10	□ VENUM18060	□ VENUL18060
18	80	10	UENUM18080	U VENUL18080
	100	10	□ VENUM18100	□ VENUL18100
	120	10	□ VENUM18120	U VENUL18120
	140	10	□ VENUM18140	□ VENUL18140
	160	10	UENUM18160	U VENUL18160
20	40	10	VENUM20040	U VENUL20040
	60	10	VENUM20060	U VENUL20060
	80	10	□ VENUM20080	U VENUL20080
	100	10	VENUM20100	VENUL20100
	120	10	□ VENUM20120	U VENUL20120
	140	10	UENUM20140	VENUL20140
	160	10	□ VENUM20160	□ VENUL20160

The Venclose<sup>™</sup> EVSRF Catheter is intended to be used with the Venclose<sup>™</sup> digiRF<sup>™</sup> Generator as a system. The Venclose<sup>™</sup> EVSRF catheter is intended for endovascular coagulation of blood vessels in patients with superficial vein reflux. The Venclose<sup>™</sup> EVSRF Catheter is contraindicated in patients with thrombus in the vein segment to be treated. Potential adverse events include but are not limited to the following: vessel perforation; skin discoloration; nerve injury; temporary paresthesia; thrombosis; deep vein thrombosis; phlebitis; hematoma; infection; skin burn; pulmonary embolism; and pain.

The Venclose Maven<sup>™</sup> Catheter is intended for endovascular coagulation of blood vessels in patients with perforator and tributary vein reflux. The Venclose Mayen<sup>™</sup> Catheter is contraindicated in patients with thrombus in the vein segment to be treated. Potential adverse events include, but are not limited to the following: vessel perforation; skin discoloration; nerve injury; temporary paresthesia; thrombosis; deep vein thrombosis; phlebitis; hematoma; infection; skin burn; pulmonary embolism; and pain.

The Venovo<sup>™</sup> Venous Stent System is indicated for the treatment of symptomatic iliofemoral venous outflow obstruction. The Venovo<sup>™</sup> Venous Stent is not designed for repositioning or recapturing. The Venovo<sup>™</sup> Venous Stent System is contraindicated for use in patients with a known hypersensitivity to nitinol (nickel-titanium) and tantalum and patients who cannot receive intraprocedural anti-coagulation therapy. Potential Adverse Events include, but are not limited to: Allergic/anaphylactic reaction · Death related/unrelated to procedure · Dissection · Embolization · Extravasation · Hematoma, puncture site ·Hypotension/hypertension  $\cdot$  Incorrect positioning of the stent requiring further stenting or surgery  $\cdot$ Intimal injury/dissection · Ischemia/infarction of tissue/organ · Malposition (failure to deliver the stent to the intended site) · Open surgical repair · Pulmonary embolism · Stent Fracture · Stent Migration · Venous occlusion/thrombosis/restenosis Please consult product labels and instructions for the use of indications, contraindications, hazards, warnings, and precautions.

The Aspirex<sup>®</sup> Thrombectomy System is indicated for the removal of acute emboli and thrombi from vessels of the peripheral venous system. The 6F and 8F Aspirex<sup>™</sup> Thrombectomy Catheters are indicated for the removal of acute emboli and thrombi from hemodialysis access grafts and native arteriovenous fistulas. The Aspirex<sup>™</sup> Mechanical Aspiration Thrombectomy System is not for use in the vessels of the cardiac, pulmonary, coronary and neurovasculature. Potential adverse events include, but are not limited to: air embolism; vessel spasm; thrombosis; dissection or perforation; or valve damage.

The Denali" Filter is indicated for use in the prevention of recurrent pulmonary embolism via placement in the vena cava in the following situations: Pulmonary thromboembolism when anticoagulants are contraindicated · Failure of anticoagulant therapy for thromboembolic disease · Emergency treatment following massive pulmonary embolism where anticipated benefits of conventional therapy are reduced · Chronic, recurrent pulmonary embolism where anticoagulant therapy has failed or is contraindicated. The Denali<sup>®</sup> Filter may be removed according to the instructions supplied in the Instructions for Use under the section labeled: "Optional Procedure for Filter Removal". The Denali" Vena Cava Filter should not be implanted in: Patients with an IVC diameter larger than 28 mm. Pregnant patients when fluoroscopy may endanger the fetus. Risks and benefits should be assessed carefully. Patients with risk of septic embolism. · Patients with uncontrolled sepsis. · Patients with known hypersensitivity to nickel-titanium alloys. The Denali<sup>™</sup> Vena Cava Filter should not be retrieved if significant thrombus is in or near the filter.

The Atlas<sup>®</sup> Gold PTA Dilatation Catheter is indicated for use in Percutaneous Transluminal Angioplasty of the peripheral vasculature, including the iliac arteries and iliac and femoral veins, and for the treatment of obstructive lesions of native or synthetic arteriovenous dialysis fistulae. This device is also indicated for post-dilatation of stents and stent grafts in the peripheral vasculature. This catheter is not for use in coronary arteries. Contraindications: None known. Potential Adverse Reactions: The complications which may result from a peripheral balloon dilatation procedure include: • Acute thrombotic occlusion • Additional intervention · Allergic reaction to drugs or contrast medium · Aneurysm or pseudoaneurysm · Arrhythmias · Balloon rupture · Balloon getting stuck on stent · Distal embolization (PE) · Hematoma · Hemorrhage, including bleeding at the puncture site · Hypotension/hypertension · Inflammation · Leg edema · Occlusion · Pain or tenderness · Pneumothorax or hemothorax · Sepsis/infection · Shock · Short term hemodynamic deterioration · Stent disruption or dislodgement with balloon insertion · Stroke · Thrombosis · Vessel dissection, perforation, rupture, or spasm.

Please consult product labels and instructions for use for indications, contraindications, hazards, warnings, and precautions.

Venous disease impacts millions of people all around the world. The BD venous portfolio consists of category-leading products, innovative solutions and dedicated patient resources – all designed to help improve clinical efficiency and enhance patient care. Learn more by scanning the QR code.



# Your Partner in Venous Care



# Simplified **RF** Ablation

# Venclose<sup>™</sup>

**RF** Ablation Catheter



long and short refluxing vein segments with one catheter.\* **Flexible** Torquable Curved

Together with its flexible and curved catheter design, the torquable Venclose<sup>™</sup> RF Ablation Catheter helps to navigate tortuous anatomies.

**DUAL HEATING LENGTHS** 10 CM & 2.5 CM

**6F** LOW-PROFILE

## **Patient Outcomes**





Images courtesy of Matthew Wise, MD, Advanced Vein Center, Orange, CA. Individual results may vary. After treatment image is representative of a patient typically 2 weeks post-op. \* As of January 2025

# **Modernized** 360° Solution

# Venclose Maven<sup>™</sup>

**Perforator Catheter** 



Curved



# **Engineered for Strength & Flexibility**

## Venovo<sup>™</sup>

Venous Stent System

### **Designed for Veins**

The Venovo<sup>™</sup> Venous Stent is designed to offer a balance between radial strength, compression resistance, and flexibility.

### **Triaxial Delivery**

Triaxial delivery system designed for placement accuracy

### Flared Ends

Unique 3 mm flared ends designed to help reduce stent migration and maximize wall apposition

### **Radial Force**

Highest mean radial resistive force among tested iliofemoral venous stents<sup>1</sup>

### **Patient Outcomes**

Post thrombotic patient treated at the common iliac vein.





Case by David Dexter, MD. Individual treatment results may vary.

Radial resistive force was tested at 13 mm crimp diameter (1 mm oversizing) using a radial expansion force gauge. Results shown are averages measured in N/mm as follows (n=6): Venovo® Venous Stent System (0.126), Medtronic Abre® Venous Stent (0.104), Cook Zilve® Vena® (0.063) and Boston Scientific VICI VENOUS STENT® (0.054). Data on file. BD, Tempe, AZ. Bench tests may not be indicative of clinical performance. Different test methods may yield different results

# **3-in-1** Thrombectomy Aspirate · Macerate · Transport

# **Aspirex**<sup>™</sup>

Mechanical Aspiration Thrombectomy System

## **Complete Mechanism of Action**



The Aspirex<sup>™</sup> Mechanical Aspiration Thrombectomy System offers a 3-in-1 mechanism of action designed to aspirate, macerate, and transport clot out of the vessel.





Aspiration

Maceration

## **Real Results**

Recanalization of an Acute Iliofemoral Deep Vein Thrombosis Using the Aspirex<sup>™</sup> 10F Catheter Dr. Michael Lichtenburg, Chief Medical Officer, Karolinen Hospital, Arnsberg, Germany

Venogram shows complete thrombotic occlusion of the left iliac vein (Figure 1). Mechanical thrombectomy was performed with the 10F Aspirex<sup>™</sup> Catheter (*Figure* 2). At the 3-month clinical follow up, the patient presented symptomfree. Venous outflow was shown to be patent on the treated side.





**Transportation** 

Individual treatment results may vary



# Advanced Filter Design

## Denali<sup>™</sup> Vena Cava Filter

### Help Protect Against Recurrent Pulmonary Embolism

The advanced design of the Denali<sup>™</sup> Filter is engineered to help filter stability, strength, and retrieval.



## **BD Reach**<sup>™</sup> Program

SV NOV

The BD Reach<sup>™</sup> Program is an industry-leading initiative designed to help physicians contact their patients with a Denali<sup>™</sup> Vena Cava Filter.

reach.bd.com

# **Built for Large Diameter Vessels**

# Atlas<sup>™</sup> Gold

**PTA Dilatation Catheter** 

### Indicated for Iliofemoral Veins

The Atlas<sup>™</sup> Gold PTA Dilatation Catheters are the **only** fiber, ultra non-compliant balloons designed for large diameter vessels.<sup>1</sup>

### Predictable

Virtually no balloon growth beyond stated diameter, which helps reduce the risk of overdilation<sup>2</sup>

**High Pressure** Dilation up to 18 ATM on select sizes

<sup>1</sup> On the U.S. market, as of January 2025. <sup>2</sup> Bench testing (n=150; 30 samples each of 12 mm x 2 mm x 80 cm, 12 mm x 6 mm x 80 cm, 18 mm x 6 mm x 80 cm, 26 mm x 2 mm x 80 cm, 26 mm x 4 mm x 80 cm) may not be indicative of clinical performance. Atlas" Gold Catheter exhibited 2% or less mean growth in outer diameter between nominal pressure and rated burst pressure. Data on file, BD, Tempe, AZ. Different test methods may yield different results.

Aspirex<sup>™</sup> Catheter Set

Size Length (cm)

135 85

110

110

6F

8F

10F

Aspirex<sup>™</sup> Drive System

Drive System 📃 80300

Aspirex<sup>™</sup> Catheter Set includes

catheter, guidewire, sterile drape, and collecting bag

Description

Product Codes

Spare Aspirex<sup>™</sup> Guidewires (5-Pack)

Diameter	Length (cm)	Тір	Flexible Tip	Product Codes
	220	Angled	40 mm	80320
0.018"	270	Angled	40 mm	80321
	320	Angled	40 mm	80322
0.025"	270	Angled	60 mm	80323

### Atlas<sup>™</sup> Gold PTA Dilatation Catheter

Produc Codes

80325

80326

80327

80328

Diameter (mm)	Length (cm)	Sheath (F)	Nominal Pressure (atm)	RBP (atm)	80 cm Shaft Codes	120 cm Shaft Code
12	2	7	6	18	ATG80122	ATG120122
	4	7	6	18	🗆 ATG80124	🗆 ATG120124
	6	7	6	18	ATG80126	ATG120126
	2	7	6	18	□ ATG80142	🗌 ATG120142
14	4	7	6	18	🗌 ATG80144	🔲 ATG120144
	6	8	6	18	🗌 ATG80146	🗆 ATG120146
	2	8	6	18	ATG80162	ATG120162
16	4	8	6	18	🗆 ATG80164	🗌 ATG120164
	6	8	6	16	ATG80166	ATG120166
	2	8	6	16	🗆 ATG80182	□ ATG120182
18	4	8	6	16	🗌 ATG80184	ATG120184
	6	9	6	16	🗆 ATG80186	□ ATG120186
20	2	9	6	16	ATG80202	ATG120202
	4	9	6	16	🗆 ATG80204	🗌 ATG120204
22	2	10	4	14	ATG80222	ATG120222
22	4	10	4	14	□ ATG80224	□ ATG120224
24	2	10	4	14	ATG80242	ATG120242
	4	10	4	14	🗆 ATG80244	🗌 ATG120244
26	2	12	4	12	ATG80262	ATG120262
	4	12	4	12	□ ATG80264	□ ATG120264

### Denali<sup>™</sup> Vena Cava Filter

Description	Qty	Product Code
Femoral Delivery Kit	1 ea	DL900F
Jugular Delivery Kit	1 ea	🗆 DL900J

### Venclose<sup>™</sup> RF Ablation Catheter

Description	Product Codes
Venclose <sup>™</sup> RF Ablation Catheter (60 cm)	UC10A256F6
Venclose <sup>™</sup> RF Ablation Catheter (100 cm)	UVC10A256F1

Venclose Maven<sup>™</sup> Perforator Catheter

Description	Product Co
Venclose Maven <sup>™</sup> Perforator Catheter (40 cm)	VC056F

### **Generator & Accesories**

Description	Product Codes
Venclose <sup>™</sup> RF Generator	VCRFG1
Venclose <sup>™</sup> Procedure Pack (No Access)	U VCPK
7 cm Micro Introducer Kit	🗆 NISO2A
12 cm Micro Introducer Kit	🗆 NISO2
12G Angiocath <sup>™</sup> IV Catheter	382277
Venclose <sup>™</sup> System Foot Pedαl	UVCFP1
Venclose <sup>™</sup> System US Power Cord	VCPCB