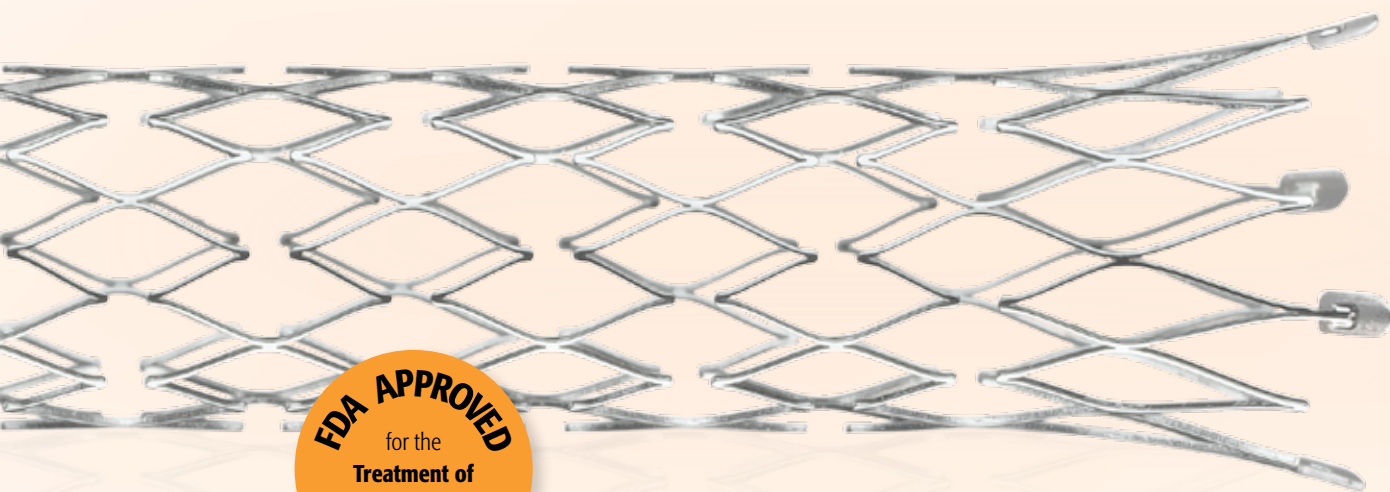


LIFESTAR[®]

Vascular Stent System



FDA APPROVED
for the
**Treatment of
Iliac Occlusive
Disease***

Simply Brilliant

Highly Visible Stent System

Optimized Pin-and-Pull Delivery System
for **Simple Deployments**

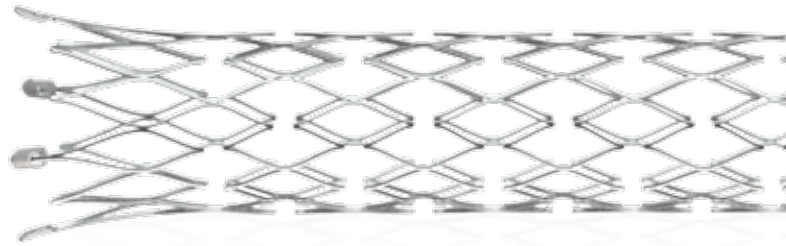
Designed for Easy, **Precise Placement**

*See back for complete indication.

BAIRD | PERIPHERAL
VASCULAR

LIFESTAR® Vascular Stent System Ordering Information

Stent Diameter (mm)	Catheter Length (cm)	Stent Length (mm)	Product Code
7	80	20	VIES07020 <input type="checkbox"/>
		30	VIES07030 <input type="checkbox"/>
		40	VIES07040 <input type="checkbox"/>
		60	VIES07060 <input type="checkbox"/>
		80	VIES07080 <input type="checkbox"/>
8	80	100	VIES07100 <input type="checkbox"/>
		20	VIES08020 <input type="checkbox"/>
		30	VIES08030 <input type="checkbox"/>
		40	VIES08040 <input type="checkbox"/>
		60	VIES08060 <input type="checkbox"/>
9	80	80	VIES08080 <input type="checkbox"/>
		100	VIES08100 <input type="checkbox"/>
		20	VIES09020 <input type="checkbox"/>
		30	VIES09030 <input type="checkbox"/>
		40	VIES09040 <input type="checkbox"/>
10	80	60	VIES09060 <input type="checkbox"/>
		80	VIES09080 <input type="checkbox"/>
		100	VIES09100 <input type="checkbox"/>
		20	VIES10020 <input type="checkbox"/>
		30	VIES10030 <input type="checkbox"/>
		40	VIES10040 <input type="checkbox"/>
		60	VIES10060 <input type="checkbox"/>
		80	VIES10080 <input type="checkbox"/>
		100	VIES10100 <input type="checkbox"/>



LIFESTAR® Vascular Stent System

Indications: The BARD® LIFESTAR® Vascular Stent System is indicated for the treatment of: • Iliac occlusive disease in patients with symptomatic vascular disease of the common and/or external iliac arteries • Residual stenoses with impaired perfusion (pressure gradient) following balloon dilatation, especially in stages III and IV according to Fontaine • Dissection • Detached arteriosclerotic plaque material and luminal obstruction following balloon dilatation • Occlusion after thrombolysis or after aspiration and before dilatation • Restenosis or reocclusion

Contraindications: Contraindications for the BARD® LIFESTAR® Vascular Stent System include, but may not be limited to: • Uncorrected coagulopathies • Functionally relevant obstruction of the inflow path, poor outflow or no distal runoff • Fresh, soft thrombotic or embolic material • Placement in the distal superficial femoral artery • Placement in the popliteal artery

Warnings: Should unusual resistance be felt at any time during the procedure, the entire system (introducer sheath or guiding catheter and stent delivery system) should be removed as a single unit. • Patients with known hypersensitivity to nickel-titanium may suffer an allergic reaction to this implant. • Stenting across a major bifurcation may hinder or prevent future diagnostic or therapeutic procedures. • Overstretching the artery may result in spasm, dissection, and/or perforation that may result in serious complications.

Precautions: This device is intended for use only by physicians who are familiar with the principles, clinical applications, complications, side effects, and risks commonly associated with vascular stenting. It is strongly recommended that physician operators adhere to all applicable institutional, local, state, and federal guidelines and protocols regarding adequate procedural training.

Potential Complications: Potential adverse events associated with the use of the BARD® LIFESTAR® Vascular Stent System include, but may not be limited to

the usual complications reported for vascular procedures such as: • Adverse and / or allergic reactions to antiplatelet agents / contrast medium / drugs / implant material • Aneurysm • Arrhythmia • Arterial occlusion / thrombosis at puncture site or remote site • Arteriovenous fistula • Bacteremia or septicemia • Bleeding from anticoagulant or antiplatelet medications • Detachment of a component of the system • Embolization, distal (air, tissue or thrombotic emboli) • Emergent surgery to remove stent • Fever • Groin hematoma, with or without surgical repair • Hemorrhage, with or without transfusion • Hyperperfusion syndrome • Hypotension / hypertension • Infection and pain at insertion site • Ischemia / infarction of tissue / organ • Pseudoaneurysm • Restenosis, recurrent narrowing or occlusion of stented segment • Stent embolization • Stent fracture • Stent malposition (failure to deliver the stent to the intended site) • Stent migration • Stent thrombosis / occlusion • Vasospasm • Vessel spasm or recoil • Vessel tear, dissection, perforation, or rupture • Vessel total occlusion

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