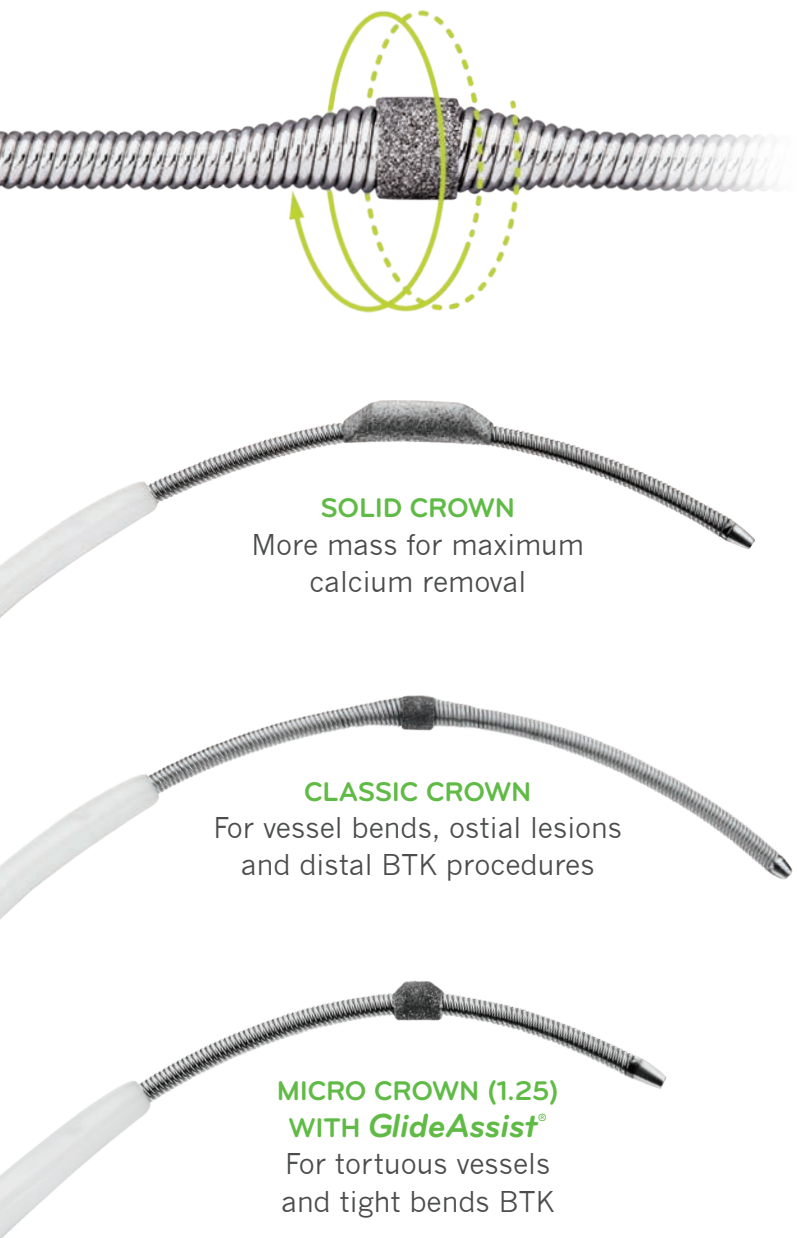


A SPECIALIZED APPROACH FOR PAD.

Our proprietary orbital technology is designed to facilitate crown contact with 360 degrees of the vessel wall. The diamond-coated crown differentially sands away arterial calcium from atherosclerotic tissue, leveraging centrifugal force to facilitate compliance change,^{1,2} while the orbital action allows continuous blood flow during treatment.¹

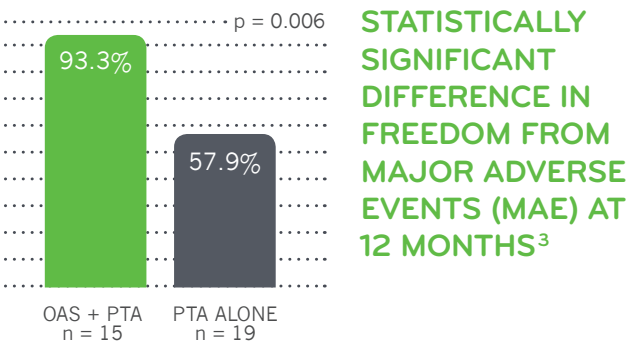
See it at work at csi360.com.



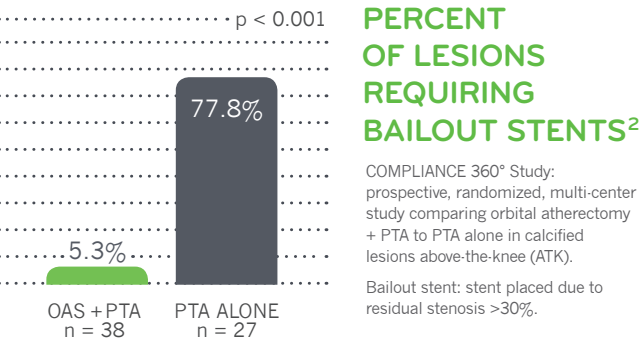
FEWER COMPLICATIONS

Using orbital atherectomy + low pressure PTA resulted in low complications³ and less bailout stents², which is especially critical for popliteal lesions.

STUDIES DEMONSTRATE OAS IS THE SMARTER SOLUTION.



Calcium 360° Study: randomized, multi-center study comparing orbital atherectomy + PTA to PTA alone in calcified lesions below-the-knee (BTK).
MAE: Device- or procedure-related major amputation (above-the-ankle), all-cause mortality and TLR/TVR.



COMPLIANCE 360° Study: prospective, randomized, multi-center study comparing orbital atherectomy + PTA to PTA alone in calcified lesions above-the-knee (ATK).
Bailout stent: stent placed due to residual stenosis >30%.

CHOOSE THE RIGHT DEVICE FOR THE JOB.

With a portfolio of three unique crown designs in multiple sizes and varying lengths, the Diamondback 360° Peripheral Orbital Atherectomy System is redefining minimally invasive.


DESCRIPTION	CROWN SIZE AND SHAFT LENGTH	SHEATH SIZE	MODEL NUMBER
Standard Length	1.25 MICRO, 145 CM with <i>GlideAssist</i> ®	4 Fr	DBP-125MICRO145
	1.25 SOLID, 145 CM	4 Fr	DBP-125SOLID145
	1.50 SOLID, 145 CM	5 Fr	DBP-150SOLID145
	2.00 SOLID, 145 CM	6 Fr	DBP-200SOLID145
	1.50 CLASSIC, 145 CM	5 Fr	DBP-150CLASSIC145
Radial-Extended Length	2.00 CLASSIC, 145 CM	6 Fr	DBP-200CLASSIC145
	1.25 SOLID, 200 CM	5 Fr	DBP-125SOL200
	1.50 SOLID, 200 CM	5 Fr	DBP-150SOL200
	1.75 SOLID, 180 CM	5 Fr	DBP-175SOL180

The CSI® Peripheral Orbital Atherectomy System is a percutaneous orbital atherectomy system indicated for use as therapy in patients with occlusive atherosclerotic disease in peripheral arteries and stenotic material from artificial arteriovenous dialysis fistulae. Contraindications for the system include use in coronary arteries, bypass grafts, stents, or where thrombus or dissections are present. Although the incidence of adverse events is rare, potential events that can occur with atherectomy include: pain, hypotension, CVA/TIA, death, dissection, perforation, distal embolization, thrombus formation, hematuria, abrupt or acute vessel closure, or arterial spasm.

Caution: Federal law (USA) restricts this device to sale by, or on the order of, a physician.

THAT'S THE POWER OF CONSTANT PROGRESS.

For more information, please contact your local CSI representative or call **1-877-274-0901**.

**CARDIOVASCULAR SYSTEMS, INC.**

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877-274-0901 **W:** csi360.com

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DIAMONDBACK 360®
PERIPHERAL ORBITAL ATHERECTOMY SYSTEM



PROVEN PERFORMANCE
IN CALCIFIED LESIONS.

THE SMART SOLUTION TO TREAT YOUR PATIENTS.

Expand your PAD treatment options with multiple access solutions and low profile advantages of the revolutionary Diamondback 360° Peripheral Orbital Atherectomy device. The low profile design facilitates minimally invasive treatment options for those with peripheral artery disease.



CROWN SIZING GUIDE

- 2.00 mm Solid Crown
- 1.75 mm Solid Crown

- 2.00 mm Solid Crown
- 1.75 mm Solid Crown
- 1.50 mm Solid Crown
- 2.00 mm Classic Crown

- 1.25 mm Micro Crown with GlideAssist®
- 1.50 mm Solid Crown
- 1.25 mm Solid Crown
- 2.00 mm Classic Crown

- 1.25 mm Micro Crown with GlideAssist®
- 1.25 mm Solid Crown
- 1.50 mm Classic Crown

- 1.25 mm Micro Crown with GlideAssist®

ILIAC AND COMMON FEMORAL ARTERIES
VESSEL DIAMETER RANGE: 5.0-10.0 mm

SUPERFICIAL FEMORAL ARTERY
VESSEL DIAMETER RANGE: 4.0-7.0 mm

POPLITEAL ARTERY
VESSEL DIAMETER RANGE: 4.0-6.0 mm

TIBIAL AND PERONEAL ARTERIES
VESSEL DIAMETER RANGE: 2.0-4.0 mm

DISTAL TIBIAL AND PEDAL ARTERIES
VESSEL DIAMETER RANGE***: 2.0-2.5 mm

* CSI data on file.
** Set-up times may vary.

*** Do not use the device in a vessel that is too small for the crown. The reference vessel diameter at the treatment area must be at least 2.00 mm in diameter for the 1.25 mm Micro Crown.