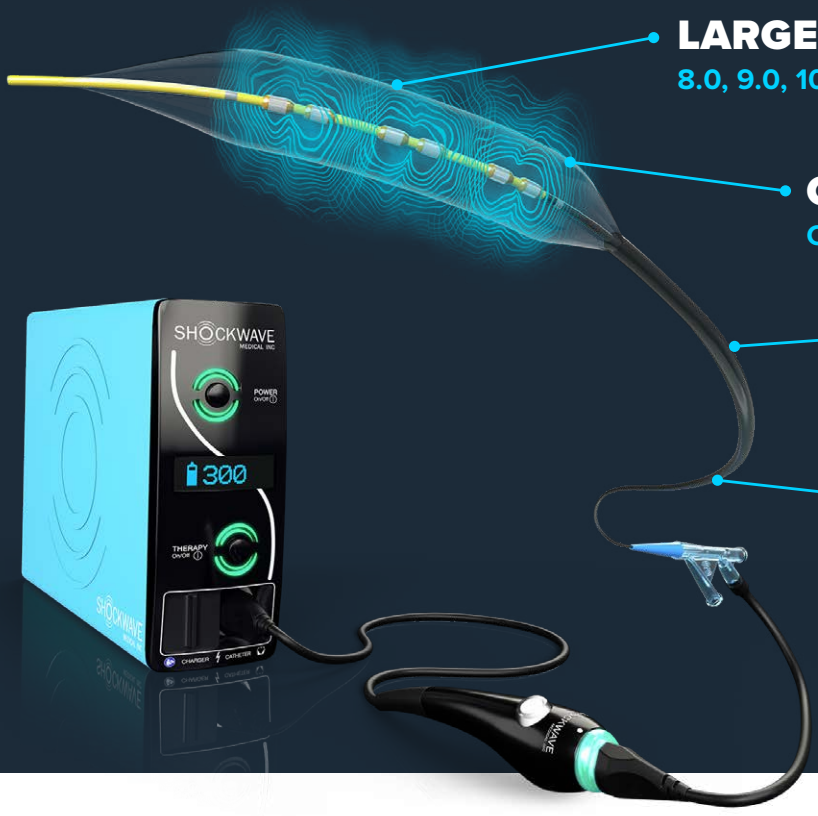
SHOCKWAVE | L6

CRACK CALCIUM IN LARGE VESSELS

Count On It



LARGE VESSEL TREATMENT
8.0, 9.0, 10.0 & 12.0 mm diameter sizes

CONSISTENT HIGH ENERGY
Compact emitter design

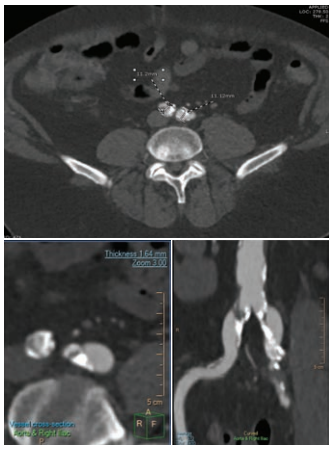
ULTRA-LOW PRESSURE
IVL therapy at 2-4 atm

SUPPORT YOU NEED
.018" guidewire

APPROPRIATE SIZING OPTIMIZES ENERGY TRANSFER

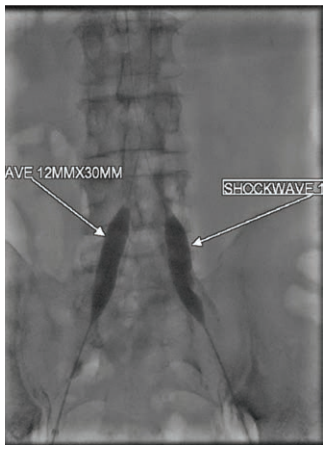
Case: Calcified Bilateral Iliacs

Pre-Procedural CTA



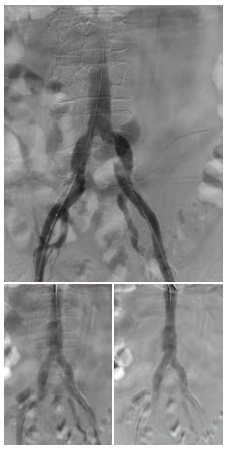
CTA Measurements
RCIA: 11.2 mm/LCIA: 11.1 mm

IVL Treatment Angio



Bilateral **Shockwave L6 12.0 mm**
300 pulses delivered

Final Angio

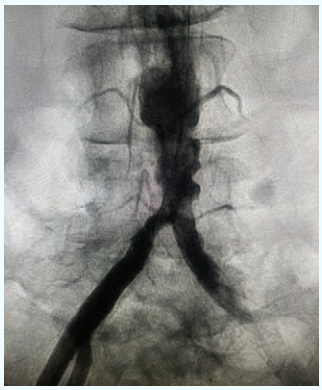


Post-op widely patent iliac stents

Case Courtesy of Dr. Paul Foley

PRE-TAVR ACCESS

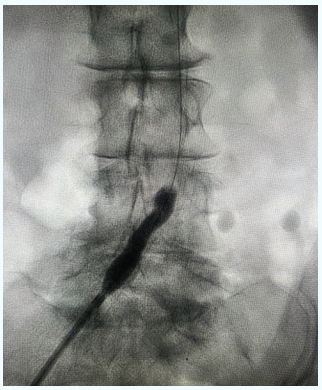
Pre-Treatment Angiogram

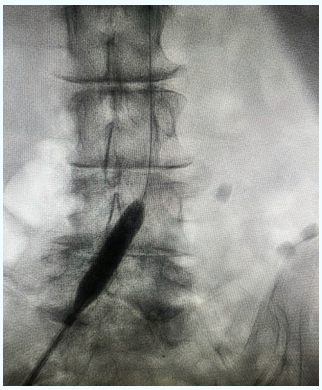


25 mm calcified focal ostial lesion

Case Courtesy of Dr. Samer Garas

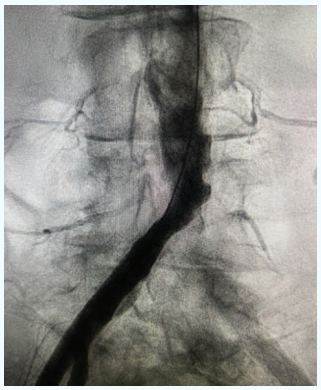
IVL Treatment





Shockwave L6 9.0 mm x 30 mm, 300 pulses

Post-IVL Treatment Angiogram



IVL successfully changed vessel compliance for valve delivery

SHOCKWAVE L6 PERIPHERAL IVL CATHETER SPECIFICATIONS

Catalog Number	Diameter (mm)	Length (mm)	Sheath Compatibility	Working Length (cm)	Pulses/ Cycle	Cycle	Pulses (Max)	Crossing Profile (in)
L6IVL080030	8.0	30	7 F	110	30	10	300	.086
L6IVL090030	9.0	30	7 F	110	30	10	300	.087
L6IVL100030	10.0	30	8 F	110	30	10	300	.091
L6IVL120030	12.0	30	8 F	110	30	10	300	.093

Visit www.shockwavemedical.com for more information.

Drs. Foley and Garas are paid consultants of Shockwave Medical.
All cases featured throughout were performed with Shockwave L6 at 1 Pulse per Second.

Peripheral IVL Important Safety Information

In the United States: Rx only.

Indications for Use—The Shockwave Medical Intravascular Lithotripsy (IVL) System is intended for lithotripsy-enhanced balloon dilatation of lesions, including calcified lesions, in the peripheral vasculature, including the iliac, femoral, ilio-femoral, popliteal, infra-popliteal, and renal arteries. Not for use in the coronary, carotid or cerebral vasculature.

Contraindications—Do not use if unable to pass 0.014" (M5, M5+, S4, E8) or 0.018" (L6) guidewire across the lesion-Not intended for treatment of in-stent restenosis or in coronary, carotid, or cerebrovascular arteries.

Warnings—Only to be used by physicians who are familiar with interventional vascular procedures—Physicians must be trained prior to use of the device—Use the generator in accordance with recommended settings as stated in the Operator's Manual.

Precautions—use only the recommended balloon inflation medium—Appropriate anticoagulant therapy should be administered by the physician—Decision regarding use of distal protection should be made based on physician assessment of treatment lesion morphology.

Adverse effects–Possible adverse effects consistent with standard angioplasty include–Access site complications –Allergy to contrast or blood thinner–Arterial bypass surgery—Bleeding complications—Death—Fracture of guidewire or device—Hypertension/Hypotension—Infection/sepsis—Placement of a stent—renal failure—Shock/pulmonary edema—target vessel stenosis or occlusion—Vascular complications. Risks unique to the device and its use—Allergy to catheter material(s)— Device malfunction or failure—Excess heat at target site.

Prior to use, please reference the Instructions for Use for more information on indications, contraindications, warnings, precautions and adverse events. www.shockwavemedical.com/IFU

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