

CASE STUDY

Embolization for High-Flow Priapism

BY MARK HORVATH, DO, AND DANIEL LOCASCIO, MS, MD

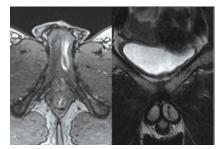


Figure 1. Dynamic contrast-enhanced MRI showing an arteriocavernosal fistula involving the left corpora cavernosa.

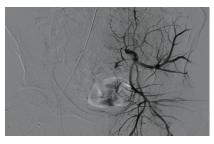


Figure 2. Angiogram demonstrating a large arteriocavernosal fistula with intense arterial blush over the left corpora.

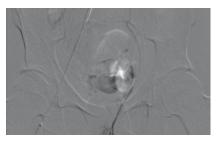


Figure 3. Angiogram obtained after using the Direxion™ Torqueable Microcatheter and Fathom™-16 Guidewire showing similar findings as in Figure 2.

CASE PRESENTATION

A 24-year-old man was referred from urology for possible endovascular treatment of high-flow priapism. He originally presented to urology clinic following symptoms from perineal impalement from a skateboard injury. The patient had complaints of persistent partial erections and inability to achieve and maintain full erections.

An initial sonogram with Doppler was obtained, which did not demonstrate any abnormality. Dynamic contrast-enhanced MRI revealed an arteriocavernosal fistula involving the left corpora cavernosa (**Figure 1**).

Potential treatment options for this patient were endovascular embolization and conservative management. According to the American Urologic Association guidelines on the management of high-flow (nonischemic) priapism, up to 62% of cases will resolve spontaneously; however, up to one-third will have an associated complaint of erectile dysfunction.¹ Given the patient's age and desire for a rapid resolution of symptoms, endovascular embolization was chosen.

Procedure Description

After using standard techniques to access the right common femoral artery, a 5-F (1.67-mm) sheath was placed, and a 4-F (1.33-mm) hockey stick catheter was used to select the left internal iliac artery. An angiogram demonstrated a large arteriocavernosal fistula with intense arterial blush over the left corpora (**Figure 2**). Next, a 0.021-inch (0.53-mm) Direxion™ Torqueable Microcatheter and Fathom®-16 Guidewire were used to select the left internal pudendal artery, and subsequent angiogram revealed similar findings (**Figure 3**). The microcatheter was then advanced just beyond the site of the arteriocavernosal fistula and two 2-mm X 4-cm IDC™ Soft Microcoils were deployed.

Repeat angiogram revealed complete resolution of the fistula and preservation of the dorsal penile artery (**Figure 4**). Of note, the patient's erection was visible on fluoroscopic spot images and immediately detumesced following coil deployment (**Figure 5**).

The patient had an uneventful overnight hospital stay and was discharged in good condition. The morning following the procedure, he endorsed nocturnal penile tumescence but otherwise the priapism continued to resolve. The patient was seen approximately 7 weeks following embolization and remained very pleased with the result, endorsing full erections and no further priapism.

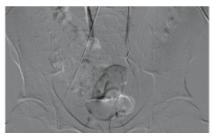


Figure 4. Repeat angiogram showing complete resolution of the fistula and preservation of the dorsal penile artery.

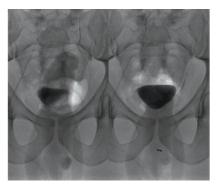


Figure 5. Fluoroscopic image after coil deployment.

Discussion

Because of the softness and packing ability of the IDC™ Soft Microcoils, they were an ideal choice for shutting down the fistula while preserving flow to the dorsal penile artery and perforating branches. The postembolization angiogram not only showed preservation of the dorsal penile artery, but also improved flow compared to the pre-embolization angiogram.

Mark Horvath, DO

Assistant Professor of Vascular and Interventional Radiology University of Florida College of Medicine Gainesville, Florida

Daniel LoCascio, MS, MD

Vascular and Interventional Radiology Fellow University of Florida College of Medicine Gainesville, Florida

Disclosures: Received no compensation for this article and is not a consultant to Boston Scientific Corporation.

1. Montague DK, Jarow J, Broderick GA, et al. American Urological Association guideline on the management of priapism. J Urol. 2003;170(4 Pt 1):1319-1324.

DIREXION™ AND DIREXION HI-FLO™

CAUTION: Federal law (USA) restricts this device to sale by or on the order of a physician. Rx only. Prior to use, please see the complete "Directions for Use" for more information on Indications, Contraindications, Warnings, Precautions, Adverse Events, and Operator's Instructions.

INTENDED USE/INDICATIONS FOR USE: The Direxion and Direxion HI-FLO Torqueable Microcatheters are intended for peripheral vascular use. The pre-loaded Fathom and Transend Guidewires can be used to selectively introduce and position the microcatheter in the peripheral vasculature. The microcatheter can be used for controlled and selective infusion of diagnostic, embolic, or therapeutic materials into the vessel. CONTRAINDICATIONS: None known WARNINGS: • Never advance or withdraw an intravascular device against resistance until the cause of resistance is determined by fluoroscopy. Movement of the microcatheter or guidewire against resistance may result in damage or separation of the microcatheter or guidewire tip, or vessel perforation. • This Direxion Microcatheter family is not intended for use in the coronary vasculature or neurovasculature. • The Direxion HI-FLO Microcatheter is not designed for the delivery of embolic coils. • Use of excessive force to manipulate the microcatheter against resistance can cause a fracture in the inition shaft. Take care not to over-torque the microcatheter, and to relieve any tension before withdrawal by rotating the microcatheter in the opposite roit. On PRECAUTIONS: • This device should be used only by physicians thoroughly trained in percutaneous, intravascular techniques and procedures. • Do not introduce the microcatheter without guidewire support as this may cause damage to the proximal shaft of the catheter. • Because the microcatheter may be advanced into narrow sub-selective vasculature, repeatedly assure that the microcatheter has not been advanced so far as to interfere within time tension. ADVERSE EVENTS: The Adverse Events include, but are not limited to: • Allergic reaction • Death • Embolism • Hemorrhage/Hematoma • Infection • Pseudoaneurysm • Stroke • Vascular thrombosis • Vessel occlusion • Vessel spasm • Vessel trauma (dissection, perforation, rupture)

FATHOM-16 STEERABLE GUIDEWIR

CAUTION: Federal law (USA) restricts this device to sale by or on the order of a physician. Rx only. Prior to use, please see the complete "Directions for Use" for more information on Indications, Contraindications, Warnings, Precautions, Adverse Events, and Operator's Instructions.

INTENDED USE/INDICATIONS FOR USE: The FATHOM -16 Steerable Guidewire is intended for general intravascular use in the peripheral vasculature. It can be used to selectively introduce and position catheters and other interventional devices swithin the peripheral vasculature. This device should be used only by physicians trained in percutaneous, intravascular techniques and procedures. CONTRAINDICATIONS: None known. WARNINGS: The FATHOM Steerable Guidewire is not intended for use in the coronary vasculature or the neuro vasculature. ADVERSE EVENTS: Complications attributed to endovascular procedures are the following: • Vessel trauma • Vessel damage • Embolism (catheter/device, air bubble, plaque, thrombus, air embolism, thromboembolism) • Pseudaneurysm • Seizure/stroke • Vessel dissection • Hemorrhage • Vascular thrombosis • Vessel occlusion • Death • Bleeding • Failed treatment • Inability to position guidewire • Damage to the catheter

INTERLOCKING DETACHABLE COIL

CAUTION: Federal law (USA) restricts this device to sale by or on the order of a physician. Rx only. Prior to use, please see the complete "Directions for Use" for more information on Indications, Contraindications, Warnings, Precautions, Adverse Events, and Operator's Instructions.

INTENDED USE/INDICATIONS FOR USE: The IDC Coil is a modified interlocking detachable coil indicated to obstruct or reduce rate of blood flow in the peripheral vasculature. This device is not intended for neurovascular use. CONTRAINDICATIONS: None known. GENERAL PRECAUTIONS: Do not advance the IDC Interlocking Detachable Coil if it is becomes lodged within the microcatheter and coil if necessary. ADVERSE EVENTS: The complications that may result from a peripheral embolization procedure include, but are not limited to: • Complications related to catheterization (e.g., hematoma at the site of entry, vessel injury, etc.) • Death • Emboli • Foreign body reactions necessitating medical intervention • Hemorrhage • Infection necessitating medical intervention • Temporary neurological deficit • Tissue necrosis • Undersiable lot formation of the vasculature • Vascupature

Direxion, Direxion HI-FLO, Fathom, IDC are unregistered or registered trademarks of Boston Scientific Corporation or its affiliates. All other trademarks are property of their respective owners.



Peripheral Interventions

300 Boston Scientific Way Marlborough, MA 01752-1234 www.bostonscientific.com

To order product or for more information contact customer service at 1.888.272.1001.

© 2016 Boston Scientific Corporation or its affiliates. All rights reserved.

PI-379504-AA MAY2016