

Charlie Mudge, RDMS, RDCS, RVT Sonographer, Digirad Corporation, Atlanta, GA Khusrow Niazi, MD Associate Professor, Director of Peripheral Vascular Intervention Emory Heart and Vascular Center, Atlanta, GA

# Significant improvement of reflux and symptoms in a young woman following Varithena treatment

### **Patient Characteristics**

- 36-year-old mother presented to the clinic with ankle swelling, leg cramps, heaviness and tender varicose veins. She sought treatment years ago but was advised to postpone treatment until she had completed her family. The patient was classified as CEAP 4 upon examination.
- Duplex ultrasound assessment revealed reflux involving the right Great Saphenous Vein (GSV) from the Saphenofemoral Junction (SFJ) to the ankle. Peak vein diameter was 8.3mm. Reflux (4.4 seconds) was present in the right common femoral vein (CFV). The deep venous system was patent.

## **Treatment and Results**

- Varithena treatment was planned for the right GSV tributary that extended medially from the mid-thigh to the ankle.
- The patient was prepped, and the skin at the access site was anesthetized with 0.2mL 1% lidocaine. Two access points were used: the tributary just above the ankle and at the knee. For both sites, a 21G micropuncture needle was inserted into the tributary followed by the passage of an 0.018 guidewire and a 4F sheath. The patient was placed in Tredelenberg and 5mL of Varithena was administered at each site for a total treatment volume of 10mL.
- Following administration of Varithena, venospasm of the treated segments was documented. Ultrasound images were obtained to confirm that the deep system was patent and that the right common femoral vein and SFJ were compressible.



Right tributary varicosity prior to Varithena treatment



Above the knee varicosity prior to treatment.



Above the knee varicosity showing an abnormal, refluxing Doppler signal.



Above the knee varicosity. Pre-treatment reflux.



Varithena filling vein during treatment.



Image of vein two-weeks post-treatment.

## **Outcome**

- The patient returned post-treatment and indicated that her edema, heaviness, swelling, tiredness, and cramps had subsided in the treated leg. The brownish discoloration continued to fade and the skin returned to a natural pink color within six weeks.
- Due to similar problems in her left leg, and the effectiveness of the procedure, the patient has since undergone treatment in that leg with similar results.



*"We have found that infusing microfoam, used in Varithena, easy to control into targeted vessel segments with near 100% closure rates and minimal complications".* 

Charlie Mudge, RVT Sonographer



#### Varithena (polidocanol injectable foam)

variation approach to the second seco

IMPORTANT SAFETY INFORMATION The use of Varithena is contraindicated in patients with known allergy to polidocanol and those with acute thromboembolic disease. Severe allergic reactions have been reported following administration of liquid polidocanol, including anaphylactic reactions, some of them fatal. Observe patients for at least 10 minutes following injection and be prepared to treat anaphylaxis appropriately. Intra-arterial injection or extravasation of polidocanol can cause severe necrois; schemia or gangene. Patients with underlying arterial disease may be at increased risk for tissue ischemia. If intra- arterial injection or extravasation of polidocanol exact severe necrois; ischemia or gangene. Patients with underlying arterial disease may be at increased risk for tissue ischemia. If intra- arterial injection or polidocanol occurs, consult a vascular surgeon immediately. Varithena can cause venous thrombosis. Follow administration instructions closely and monitor for signs of venous thrombosis after treatment. Patients with reduced mobility, history of deep vein thrombosis or pulmonary embolism, or recent (within 3 months) major surgery, prolonged hospitalization, or pregnancy are at increased risk for developing thrombosis. The most common adverse events beserve enter entity. Internot common adverse events beserve enter entity. Internot be experienced with venous procedures, posses a detailed working knowledge of the use of the duplex ultrasound in venous disease and be trained in the administration of Varithena.

For Full Prescribing Information visit Varithena.com

Varithena™ is a registered trademark of Boston Scientific. All other trademarks are property of their respective owners.

PI-1263705-AA



Advancing science for life<sup>™</sup>

#### **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.

© 2023 Boston Scientific Corporation or its affiliates. All rights reserved.

PI-796208-AA