Angio-Seal®
Vascular Closure Device

LEAD THE LAB

to OPTIMAL OUTCOMES with ANGIO-SEAL Vascular Closure Device

ACE

ANGIO-SEAL
CLINICAL
EXPERT
PROGRAM

INCREASE & EXPAND

your knowledge of vascular closure devices to improve patient care

EDUCATE & SHARE

your knowledge on proper device usage and important factors for optimal patient selection

TRAIN & CERTIFY

clinicians on proper closure techniques with the ANGIO-SEAL Vascular Closure Device

STRENGTHEN & EXTEND

your partnership with your colleagues and Terumo as an on-site expert for ANGIO-SEAL Vascular Closure Device

TERUMO
INTERVENTIONAL SYSTEMS

ABOUT THE ANGIO-SEAL CLINICAL EXPERT PROGRAM

Terumo has developed a program to train and certify licensed physicians and/or healthcare professionals (HCPs) to be an ANGIO-SEAL Clinical Expert (ACE) on ANGIO-SEAL Vascular Closure Devices.

ACE TRAINING OBJECTIVES

Upon completion of the ACE training program, the ACE will be able to:

- Deliver education and provide clinical support to ensure the safe and effective use of the ANGIO-SEAL Vascular Closure Device in accordance with the Instructions For Use (IFU)
- Identify and advise proper anatomy assessment, patient selection and deterrents for the use of the ANGIO-SEAL Vascular Closure Device, including completion of the certification process
- Highlight and avoid potential clinical factors that could result in unexpected or adverse events by way of troubleshooting education

ACE QUALIFICATIONS

The ACE program participant:

- Is either a licensed physician or other HCP who is duly authorized by the state and the facility to independently, or under the direction of a licensed physician, deploy ANGIO-SEAL Vascular Closure Devices
- Is a certified user of the ANGIO-SEAL Vascular Closure Device platform being utilized at the facility
- Has at least two (2) years of experience with vascular closure devices
- Is a full-time or part-time employee at the facility
- Is in good standing at the facility

ACE ROLE AND RESPONSIBILITIES

The ACE will have the following responsibilities at the facility:

- Provide educational or clinical support on ANGIO-SEAL Vascular Closure Devices to other HCPs duly authorized by the state and the facility to independently, or under the direction of a physician, deploy ANGIO-SEAL Vascular Closure Devices
- Be knowledgeable and have a high level of understanding of the various resources available on ANGIO-SEAL Vascular Closure Devices, including but not limited to the IFU
- Recognize and report to Terumo any adverse events that occur with ANGIO-SEAL Vascular Closure Devices in accordance with Terumo's Product Performance Reporting (PPR) process
- Train and certify other licensed HCPs duly authorized by the state and the facility on ANGIO-SEAL Vascular Closure Devices in accordance with Terumo's training and certification process
- There is no compensation for participation in this program
- Any reasonable and necessary travel expenses related to training will be reimbursed by Terumo

---

FIND OUT MORE Phone: 800.862.4143 terumois.com Fax: 800.411.5870

Indications: The Angio-Seal Vascular Closure Device product family, including the VIP and Evolution platforms, is indicated for use in closing and reducing time to hemostasis of the femoral arterial puncture site in patients who have undergone diagnostic angiography procedures or interventional procedures using an 8 French or smaller procedural sheath for the 8 F Angio-Seal device and a 6 French or smaller procedural sheath for the 6 F Angio-Seal device. The Angio-Seal VIP and Evolution platform devices are also indicated for use to allow patients who have undergone diagnostic angiography to safely ambulate as soon as possible after sheath removal and device placement, as well as to allow patients who have undergone an interventional procedure to safely ambulate after sheath removal and device placement.

Important Safety Information: Possible adverse events for vascular closure devices include, but are not limited to: bleeding or hematoma, AV fistula or pseudoaneurysm, infection, allergic reaction, foreign body reaction, inflammation or edema. This device should only be used by a licensed physician (or other health care professional authorized by or under the direction of such physician) possessing adequate instruction in the use of the device, e.g., participation in an Angio-Seal physician instruction program or equivalent.

RX ONLY. Refer to the product labels and package insert for complete warnings, precautions, potential complications, and instructions for use.

TERUMO INTERVENTIONAL SYSTEMS