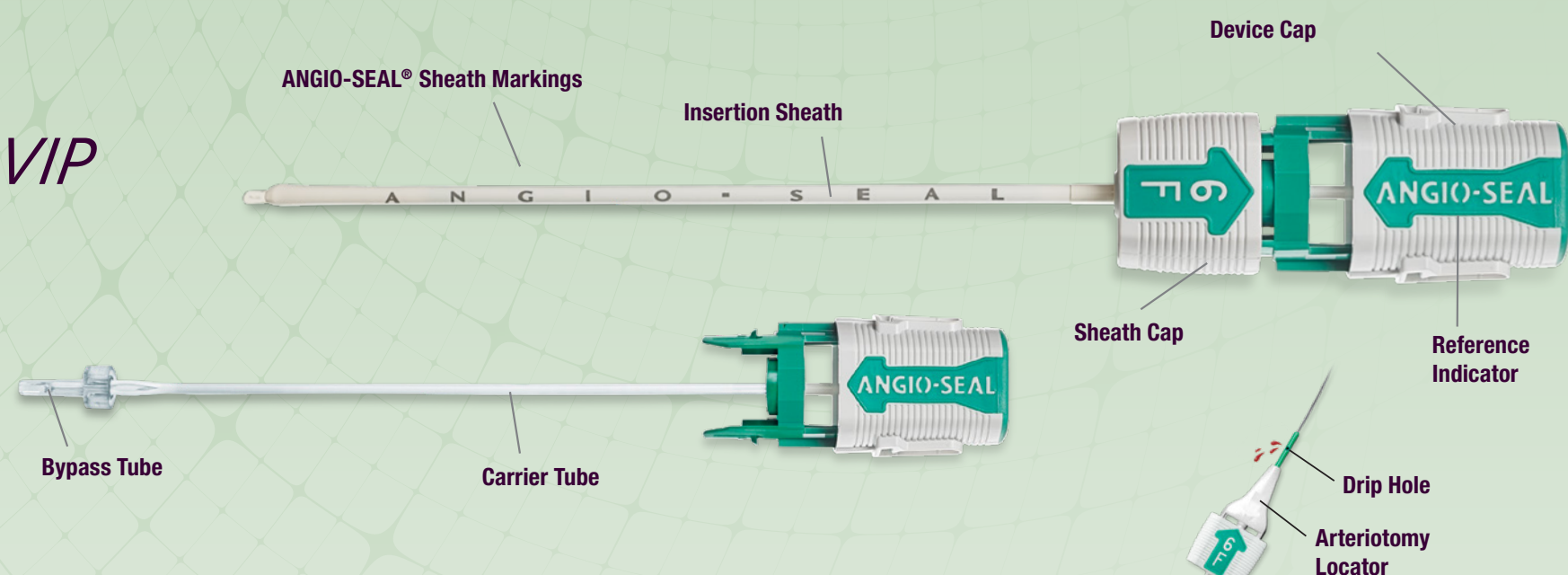


Angio-Seal® VIP

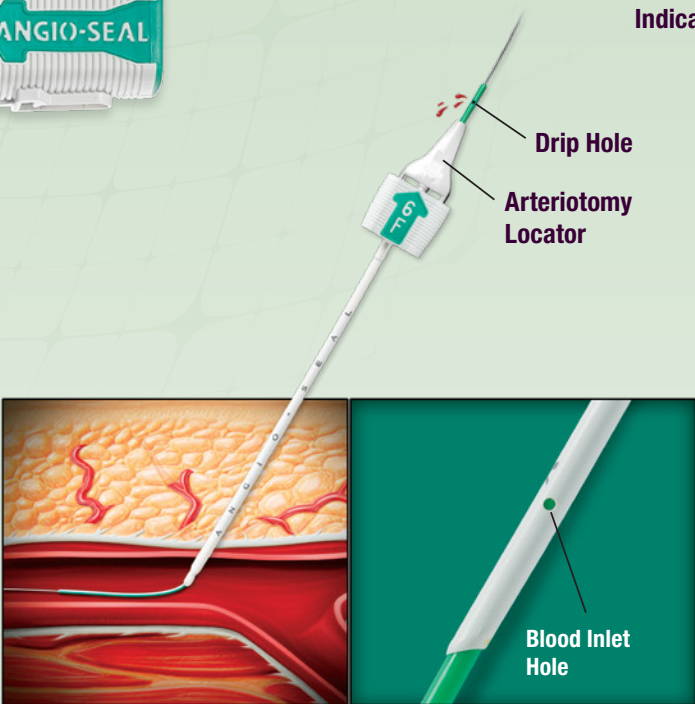
Vascular Closure Device



ANGIO-SEAL® VIP Quick Deployment Guide¹

LOCATE the Artery

1. Prior to sheath exchange, insert the arteriotomy locator into the Angio-Seal insertion sheath, making sure the two pieces snap together securely.
2. Begin by exchanging the procedural sheath with the Angio-Seal insertion sheath and locator system.
3. Confirm entry into the artery with blood flow through the locator.
4. Note sheath marking visible at skin level. The sheath markings should be used as a guide to help confirm that the device position has not changed prior to setting the anchor.
5. Hold the insertion sheath in place and remove the locator and wire.



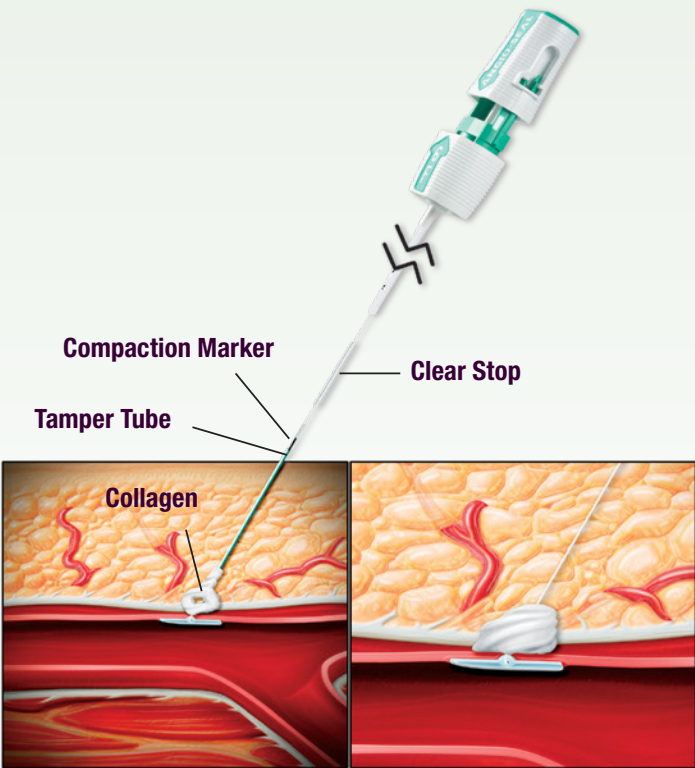
SET the Anchor

1. Carefully grasp the Angio-Seal device just behind the bypass tube and slowly insert into the insertion sheath until the sheath cap and device cap snap together. Be sure the arrow on the sheath and device sleeve align.
2. Maintaining the insertion sheath position, grasp the device cap and gently pull straight back into the full rear locked position. Resistance will be felt as the device handle and sleeve snap lock. The device sleeve bands should now be completely visible, confirming attainment of full rear locked position.



SEAL the Puncture

1. Slowly withdraw the device assembly along the angle of the puncture tract until the clear stop appears.
2. Maintain tension and advance the knot and collagen with the tamper tube. A seal occurs when resistance is felt and hemostasis is achieved. As a guide, a compaction marker will appear in most cases.
3. Cut the suture below the clear stop and remove the tamper tube.
4. Gently pull up on the suture. Push down on the skin using a sterile instrument. Cut the suture below the skin level, making sure to cut below the compaction marker.



ORDERING INFORMATION

ANGIO-SEAL® VIP		
PRODUCT CODE	FRENCH SIZE	GUIDEWIRE DIAMETER (in)
610130	6	0.035
610131	8	0.038

Contents: Vascular Closure Device, Insertion Sheath, Arteriotomy Locator and 70 cm Guidewire with “J” Straightener (10 units per box).

FIND OUT MORE



800.888.3786



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Indications:

The Angio-Seal Vascular Closure Device 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 8F Angio-Seal device and a 6 French or smaller procedural sheath for the 6F Angio-Seal device. Angio-Seal is 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 physicians with training qualifying them to perform arterial access and closure for endovascular procedures through the common femoral artery and have participated in a Terumo Medical Corporation Angio-Seal physician instruction program.

Exception (applicable to US and China only): 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. This advertisement is directed to physicians only, and not to consumers. Refer to the product labels and package insert for complete warnings, precautions, potential complications, and instructions for use.

References

1. ANGIO-SEAL® VIP Instructions for Use. ASIN0004. 2018-09-01.

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