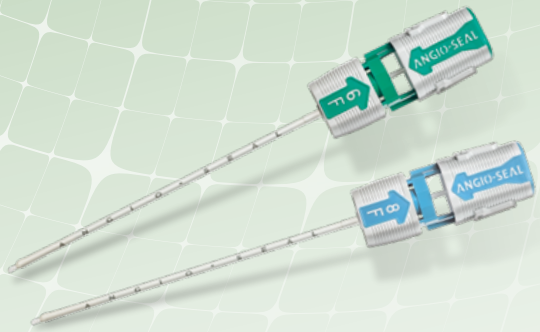
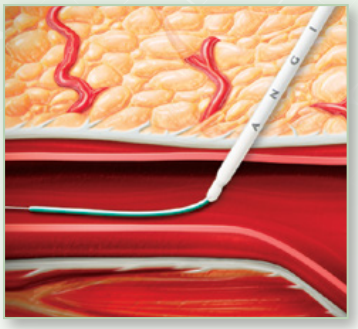
Angio-Seal®

Vascular Closure Device

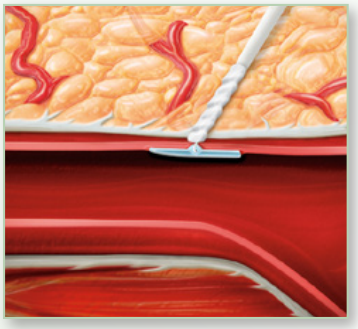


ENSURE SUCCESSFUL HEMOSTASIS



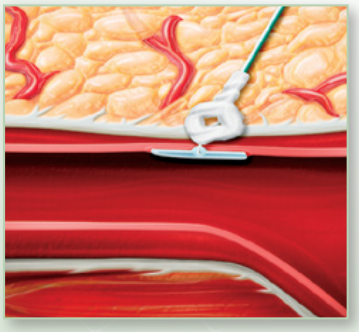
Locate the Artery

1. Exchange the procedure sheath with the Angio-Seal locator system until blood flow through the locator is observed, indicating position in the artery.
2. Withdraw the locator system until blood slows or stops, denoting the indicator is outside the artery.
3. Advance the locator system until blood flow is observed and remove the arteriotomy locator and guidewire from the insertion sheath.



Set the Anchor

1. Maintaining sheath position, insert the Angio-Seal device into the sheath, until the components snap together indicating proper fitment.
2. Maintaining sheath position, gently pull back on the device cap until you feel resistance, indicating you are in rear-lock position.
3. The anchor is now locked in place and the device is ready to be deployed.



Seal the Puncture

1. Slowly pull back on the Angio-Seal device until the suture has stopped spooling.
2. Maintain upward tension on the device and gently advance the tamper tube until resistance is felt and the compaction marker is revealed.
3. Cut the suture and remove the device.

For complete deployment steps, and nursing and ambulation guidelines

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INTERVENTIONAL SYSTEMS

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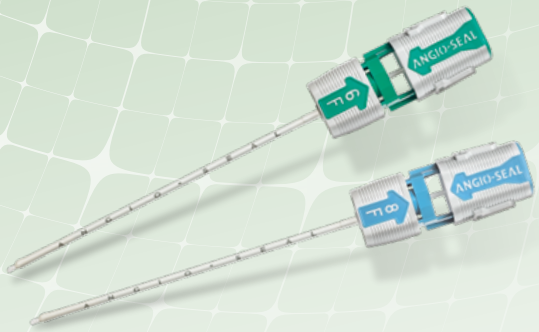

THE INSIDE ADVANTAGE™

Access Terumo's extensive education programs

For more information, please contact your local sales representative.

Indications: The Angio-Seal Vascular Closure Device is indicated for use in closing and reducing time to hemostasis of the common 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. 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 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.

References

1. ANGIO-SEAL® VIP Instructions for Use. ASIN0018. 2025-10-07.

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