FINISH STRONG
**STEP 1: ACHIEVE TEMPORARY HEMOSTASIS**

**INSERT MYNXGRIP**
- Insert MynxGrip into existing procedural sheath up to the white shaft marker

**INFLATE THE BALLOON**
- Inflate the balloon until the black marker is fully visible on the inflation indicator and close stopcock

**GENTLY PULL BACK TWO STOPS**
- Grasp black handle and withdraw catheter until the balloon abuts the distal tip of the procedural sheath (first point of resistance)
- Continue to withdraw until the balloon abuts the arteriotomy site (second point of resistance)
- While holding adequate tension on device handle, open stopcock on procedural sheath

**STEP 2: PLACE THE SEALANT**

**ADVANCE THE SEALANT**
- With stopcock open, detach shuttle and advance until resistance is felt

**UNSHEATH THE SEALANT**
- A. Lighten hold on black handle
- B. Grasp procedural sheath and withdraw it from tissue tract
- C. Continue retracting until shuttle locks onto black handle

**ADVANCE PAST SINGLE GREEN MARK**
- Ensure adequate tension is employed on the black handle to keep balloon abutted against the arteriotomy
- Immediately grasp advancer tube at skin and gently advance until single marker is fully visible
- Hold for up to 30 seconds
- Lay device down for up to 90 seconds

**STEP 3: REMOVE THE DEVICE**

**LOCK, STABILIZE, DEFlate**
- Lock syringe to maximum negative position

**STABILIZE ARTERY**
- Stabilize by applying light-fingertip compression proximal to the insertion site
- Lightly grasp advancer tube at skin with thumb and forefinger; realign with tissue tract

**DEFLATE THE BALLOON**
- Open stopcock to deflate balloon
- To ensure complete balloon deflation, wait until air bubbles and fluid have stopped moving through the inflation tubing

**REMOVE CATHETER AND ADVANCER TUBE**
- Withdraw catheter through the advancer tube lumen
- NOTE: If unusual resistance is felt during catheter withdrawal, pull the advancer tube and balloon catheter together through the tissue tract
- Remove advancer tube from the tissue tract
- Fingerfip compression can be applied for up to 60 seconds or as needed
- Assess for hemostasis and apply additional fingertip compression until sterile dressing is applied and hemostasis is achieved

**RESULT: SEALANT IS IN PLACE**

**RESULT: CONFIRM POSITION AT THE ARTERIOTOMY**

**RESULT: SECURE EXTRAVASCULAR CLOSURE**

**PREP MYNXGRIP**
- Hold the MynxGrip Vascular Closure Device by the shuttle while removing from the tray

**PREPARE BALLOON**
- Fill locking syringe with 2-3mL of sterile saline
- Attach to stopcock and draw vacuum
- Inflate balloon until black marker on inflation indicator is fully visible
- Deflate balloon and leave syringe at neutral
- Do not remove sealant sleeve

**INSERT MYNXGRIP**

**ADVANCE THE SEALANT**

**UNSHEATH THE SEALANT**

**ADVANCE PAST SINGLE GREEN MARK**

**LOCK SYRINGE**

**STABILIZE ARTERY**

**DEFLATE THE BALLOON**

**REMOVE CATHETER AND ADVANCER TUBE**

**RESULT**
- Sealant is in place
- Confirm position at the arteriotomy
- Secure extravascular closure
ORDERING INFORMATION

THE MYNXGRIP VASCULAR CLOSURE DEVICE INCLUDES:

(1) Balloon catheter with integrated sealant
(1) 10ml locking syringe

INDICATIONS FOR USE

The MynxGrip Vascular Closure Device is indicated for use to seal femoral arterial access sites while reducing times to hemostasis and ambulation in patients who have undergone diagnostic or interventional endovascular procedures utilizing a 5F, 6F or 7F procedural sheath.

WARNINGS

Do not use if the sterilization indicator dot on the pouch is yellow/gold.
Do not use if components or packaging appear to be damaged or defective or if any portion of the packaging has been previously opened.
Do not reuse or resterilize. The MynxGrip device is for single use only. The catheter is loaded with a single hydrogel sealant. Reuse of the device would result in no delivery of hydrogel sealant.

Do not use the MynxGrip device if the puncture site is located above the most inferior border of the inferior epigastric artery (IEA) and/or above the inguinal ligament based upon bony landmarks, since such a puncture site may result in a retroperitoneal hematoma/bleed. Perform a femoral angiogram to verify the location of the puncture site.
Do not use the MynxGrip device if the puncture is through the posterior wall or there are multiple punctures, as such punctures may result in a retroperitoneal hematoma/bleed.

IN THE UNITED STATES, FAX YOUR MYNXGRIP ORDER TO ACCESSCLOSURE AT (877) 933-0133.
OUTSIDE OF THE UNITED STATES, CONTACT YOUR LOCAL MYNX DISTRIBUTOR.

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PRECAUTIONS

The MynxGrip device should only be used by a trained licensed physician or healthcare professional.
The MynxGrip device should not be used in patients with a known allergy to PEG.

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AccessClosure.com

Learn More

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