Femoral vascular access-site complications in the cardiac catheterization laboratory: diagnosis and management

Vascular access-site complications remain a major cause of morbidity and mortality with cardiac catheterization and percutaneous intervention using the femoral approach. Complications may be divided into major and minor. Major complications include bleeding that requires transfusion, retroperitoneal hemorrhage and nonhemorrhagic complications such as pseudoaneurysm, arteriovenous fistula, arterial dissection, thrombosis and limb ischemia. Minor complications include minor bleeding, ecchymosis and hematoma. The incidence of access complications varies by the population studied. Numerous patient-related and procedure-related risk factors have been associated with vascular access-site complications. Alternate access sites, the use of fluoroscopic guidance, focus on anticoagulant and antiplatelet therapy, and arterial closure devices are all methods being investigated to prevent and reduce complications. History and physical examination are important in identifying vascular access-site complications and imaging is helpful to confirm the diagnosis.

Cardiac catheterization and percutaneous intervention can result in vascular access-site complications. It is important for individuals caring for the patient returning from the cardiac catheterization laboratory to be aware of and recognize the various complications that can occur and how they should be treated. This article will focus on the vascular access-site complications via the femoral approach, which can lead to significant morbidity and mortality.

Anatomy & technique

The external iliac artery crosses under the inguinal ligament to become the common femoral artery. The inferior epigastric artery branches off the external iliac artery inferiorly [1]. The common femoral artery bifurcates below the inguinal ligament into the superficial and profunda arteries. The common femoral artery is housed in the femoral triangle and has the benefit of being a large superficial vessel that can be compressed against the femoral head to achieve hemostasis [2]. Three key landmarks have traditionally been used in obtaining femoral access: the inguinal crease, the maximal pulsation of the femoral artery and bony landmarks (such as a line drawn from the anterior superior iliac crest to the pubic symphysis) [3]. The femoral artery can be cannulated at the base of the femoral triangle, just inferior to the inguinal ligament but above the bifurcation of the femoral artery [4,5]. This location corresponds to 2–3 cm below the mid-point of the pubic symphysis and the anterior superior iliac crest [6]. Because the inguinal skin crease is located below the femoral artery bifurcation in over 70% of patients, using it as a landmark for cannulation can lead to low femoral puncture [3,6]. Garrett et al. found that the femoral artery bifurcation was below the inguinal ligament and the middle of the femoral head in nearly all patients studied. The bifurcation of the common femoral artery was below the middle of the femoral head in 99% of cases [7]. Fluoroscopy is helpful in locating the position of the femoral head and therefore aids in more accurately guiding femoral puncture. Using the location of maximal femoral pulsation has been shown to ensure more consistent puncture of the common femoral artery as the maximal femoral pulse was found over the femoral artery in 93% of cases [6,8]. The goal of femoral puncture is to access the femoral artery where the artery overlies the middle third of the femoral head as depicted in Figure 1 [9].

Definitions

Vascular complications of cardiac catheterization and coronary intervention can be divided into minor and major complications. Minor complications include minor bleeding, ecchymosis and stable hematoma. Major complications include pseudoaneurysm, arteriovenous (AV) fistula, hematoma requiring transfusion, retroperitoneal

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hemorrhage, arterial dissection, embolism, thrombosis, infection, vessel rupture/perforation and limb ischemia. An illustration of a pseudoaneurysm and AV fistula can be seen in Figure 2. Bleeding is the most common complication of the transfemoral approach to catheterization and can manifest as a stable or unstable hematoma, uncontrolled bleeding, pseudoaneurysm or retroperitoneal hemorrhage. Hematomas can expand in 10–15% of patients and can manifest as hemorrhagic shock [10]. A retroperitoneal hemorrhage is a potential life threatening complication of femoral artery puncture that should be suspected in any postcatheterization patient who develops hypotension, ipsilateral flank, abdominal or back pain, or a drop in hemoglobin without a source. A pseudoaneurysm (false aneurysm) is a communication between the femoral artery and the overlying fibromuscular tissue resulting in a blood-filled cavity [11]. It is formed when blood escapes from the lumen of an artery through a defect in one or more layers of the arterial wall and forms a pocket beneath the adventitia of the artery or in the surrounding tissue near the site of arterial puncture. A pseudoaneurysm has continuity with the arterial lumen whereas a hematoma does not [10,12]. Pseudoaneurysms frequently result from failure to achieve adequate hemostasis after the catheter or sheath is removed [12]. An AV fistula is an abnormal connection between a vein and artery that is generally asymptomatic. It is more likely to result from arterial puncture below the femoral artery bifurcation and is typically created between the superficial or deep femoral artery and the adjacent lateral circumflex vein [10]. Dissection can occur in the femoral or iliac artery, but is most common in the iliac arteries due to atherosclerosis and tortuosity in these vessels [10]. Figure 3 shows an external iliac artery dissection caused by the vascular sheath. Embolic complications after catheterization are uncommon, but can lead to emboli lodging into small vessels and causing tissue ischemia [10]. Local thrombosis leading to limb ischemia is rare [9].

**Incidence**

The incidence of vascular access-site complications following catheterizations varies and in general depends on the study population. Femoral access-site complications are generally higher for interventional procedures than diagnostic procedures, which is likely related to anticoagulant therapy and sheath diameter. Femoral access-site complications have been reported to range from 0 to 17% in patients undergoing diagnostic and interventional cardiovascular procedures [2]. Chandrasekar et al. reported that femoral access-site complications were 1.8% for diagnostic and 4% for interventional procedures [13]. The incidence of major bleeding complications ranges from 2 to 6% after percutaneous coronary angioplasty [10]. Access-site injury rates requiring procedural or surgical intervention or bleeding requiring transfusion range from 2.6 to 6.6% [14]. Rates for major bleeding (bleeding requiring transfusion of more than two units of blood) in trials of glycoprotein (GP) IIb/IIIa inhibitors ranged from 1.9 to 14% [14]. Tsetis et al. reported that a significant hematoma or uncontrollable bleeding requiring transfusion or invasive procedure occurs in <1% of catheterizations [15]. Yatskar et al. found that a hematoma is the most frequent periprocedural complication and can
occur in anywhere from 2 to 12% of cases [16]. The incidence of pseudoaneurysm ranges from 0.5 to 6.3% and many sources cite an approximate incidence of 1% [17–22]. Samal et al. cites that femoral pseudoaneurysms can occur in 0.1–1.5% of diagnostic angiography and in up to 7.7% of interventional procedures [10]. The incidence of AV fistulas has been reported to be 0.2–2.1% [17–22]. Retroperitoneal hemorrhage has an incidence from 0.15 to 0.5% [11,23–25]. Trimarchi et al. studied 112,340 patients undergoing percutaneous coronary intervention (PCI) and found the incidence of retroperitoneal hemorrhage to be 0.4% [26]. Arterial dissection occurs in 0.01–0.4% of procedures [9]. Arterial thrombosis can occur in <0.5% of cases [15,21]. Risks of vascular access-site complications have changed with the advent of arterial closure devices and additional possible complications with these devices include the risk of infection, stenosis and embolism [10].

**Risk factors**

Risk factors for vascular complications of cardiac catheterization can be divided into patient-related and procedure-related factors. Patient-related risk factors include female gender, low body weight, obesity, lower body surface area, older age, peripheral vascular disease, renal failure or elevated creatinine, and low platelet count [2,9,14]. Some studies report conflicting results on other associations with vascular complications, such as myocardial infarction, cardiacogenic shock, diabetes mellitus and hypertension [14]. Procedural-related factors include previous catheterization at the same site, high doses and longer duration of anticoagulation, use of thrombolytic agents, use of GP IIb/IIIa inhibitors (particularly abciximab), larger arterial sheaths, concomitant venous sheaths, prolonged indwelling sheath duration, prolonged procedure duration, repeat PCI and location of the arterial puncture [2,9,14,27]. Sherev et al. found that 71% of all vascular access-site complications were due to low or high femoral artery puncture sites [2]. Fewer femoral complications have been noted in patients undergoing elective PCI using 6 F compared with 7- or 8-F guiding catheters [9].

Independent predictors of hematomas requiring blood transfusions include smaller patients, older age, female gender, chronic renal insufficiency, complex coronary lesions, patients with acute myocardial infarction or cardiacogenic shock presenting for an emergent procedure, use of GP IIb/IIIa inhibitors and the use of thrombolytics [16]. There is a higher risk of the formation of a pseudoaneurysm with interventional procedures than diagnostic procedures because of the length of the procedure, larger devices (and size of puncture) and more aggressive anticoagulation [28]. Low arterial access, female gender, older age, diabetes and obesity have been associated with pseudoaneurysm formation [9]. Risk of bleeding into the retroperitoneal space is increased with a high femoral puncture (particularly above the inguinal ligament and puncturing the back wall of the vessel) and postcatheterization anticoagulation [3,9,23]. Sherev et al. found that arterial puncture above the most inferior border of the inferior epigastric artery in patients undergoing PCI was associated with 100% of all retroperitoneal hemorrhages [2]. It is more difficult to compress the artery against the femoral head when the arterial

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**Figure 2. Complications of vascular access site in the femoral artery.**

Pseudoaneurysm (blood-filled cavity communicating with the vessel lumen), arterial perforation and arteriovenous fistula (abnormal communication between the femoral artery and the common femoral vein).

puncture site is below the bifurcation because the pressure is applied primarily against soft tissue rather than the bone of the femoral head in this situation [2,6]. Above the inguinal ligament, the artery resides in the retroperitoneal space making compression and hemostasis difficult. Female sex and preprocedural GP IIb/IIIa inhibitors had an odds ratio over 2 for being independent predictors of retroperitoneal hemorrhage. The use of bivalirudin was associated with a lower risk of the development of a retroperitoneal hemorrhage [26]. A high or low femoral arterial puncture, multiple puncture attempts and prolonged clotting times can increase the risk of AV fistula formation. The risk factors for lower extremity ischemia include the use of larger catheters or sheaths in relatively smaller arteries, peripheral vascular disease, older age, cardiomyopathy, hypercoagulable states and vessel dissection [9].

Prevention

There are several approaches in preventing vascular access-site complications. The radial artery has been investigated as an alternate route for cardiac catheterization. Fluoroscopic guidance has been proposed to allow for better visualization of anatomic landmarks. Advances in anticoagulant and antiplatelet agents offer reduction in bleeding complications. Arterial closure devices have been developed and are currently being studied to determine their efficacy in reducing vascular access-site complications.

■ Choice of access site

Alternative vascular access sites, such as the radial, brachial and axillary artery have been investigated. Complications associated with brachial artery access are similar to those seen with femoral access, but ischemic complications are generally more common with brachial access [10]. Radial access has been associated with lower complication rates (as low as <0.7%), improved patient comfort and reduced hospital costs [6,29]. Disadvantages of radial access include a higher incidence of procedure failure that necessitates resorting to the femoral approach, longer procedural times (at least in the ‘learning curve’ phase), postprocedural radial artery spasm and occlusion [29].

Radial access has been shown to significantly reduce bleeding complications. The PRESTO-ACS study revealed that the radial approach was associated with a significant decrease in bleeding and a nonsignificant decrease in the net clinical outcome during hospitalization compared with the femoral approach [30]. The radial approach has been found to have a decrease on death or reinfarction rates, bleeding and net clinical outcome at 1-year follow-up compared with the femoral approach. This was true despite more intense procedural use of antiplatelet therapy in patients undergoing the radial approach. The RIVAL trial investigated whether radial access was superior to femoral access in patients undergoing angiography [31]. Large hematomas and pseudoaneurysms requiring closure occurred more frequently with femoral access than radial access. Procedure time, length of hospital stay, pain at the access site and PCI contrast volume were similar in both routes, but fluoroscopy time was longer for radial access by 1.8 min. Patients were nine times more likely to prefer the radial approach for catheterization compared with the femoral approach if they were to have another procedure. There was no difference in death, myocardial infarction, stroke or noncoronary artery bypass graft-related major bleeding in the two groups.

The rate of bleeding complications requiring transfusions for radial access is 0.15% [32]. Pseudoaneurysm and AV fistula are even more uncommon with the radial approach compared with the femoral approach [33]. Radial artery occlusion is more common than femoral
artery occlusion because of the smaller artery to sheath ratio and prolonged high pressure compression. Pancholy et al. found the incidence of radial artery occlusion to be 9% [34] but in the newer series (and with the use of patent hemostasis) the incidence is closer to 3%. The decision to use the radial approach must be balanced by weighing the advantages and disadvantages. Radial access may be associated with fewer hematomas, pseudoaneurysms and AV fistulas, but it carries its own risk of arterial occlusion and arterial spasm. It is preferred by patients, costs less, may be associated with shorter hospital stays and has similar (if not better) outcomes as compared with femoral access. However, operators may not be as experienced with this route leading to longer procedures. The radial approach is a promising technique that can significantly reduce the incidence of access complications and, as its use will be more widely adopted, more long-term results will become available.

**Fluoroscopy of the femoral head**

Determining the ideal site of femoral artery cannulation with fluoroscopy can reduce a number of vascular access-site complications. Fluoroscopy can be used to identify the location of the medial border of the femoral head, which is the most reliable landmark for the common femoral artery [6]. Jacobi et al. found that the use of fluoroscopy resulted in more ideal arterial sheath placement, particularly in obese patients, but did not dramatically reduce complication rates [35]. Abu-Fadel et al. showed that fluoroscopy decreased the number of low arteriotomies, but did not decrease the number of arterial punctures or complications rates. This study did show benefit in the use of fluoroscopy in obese and in female patients [36]. Fitts et al. aimed to reduce vascular access-site complications in patients undergoing PCI and found that the use of fluoroscopy was associated with a significantly lower incidence of pseudoaneurysms and arterial injury, but found no significant difference in the incidence of bleeding complications [37]. Fluoroscopy is quick and appears to result in more successful arteriotomies while only adding a minimal amount of radiation. The routine use of fluoroscopy could be beneficial in most patients undergoing catheterization and in our experience fluoroscopy of the femoral head especially prevents the sticks above the inguinal ligament, which are most often associated with retroperitoneal bleeds.

**Ultrasound guidance**

The FAUST trial studied procedural and clinical outcomes of femoral arterial access using ultrasound guidance compared with standard fluoroscopic guidance. In general, ultrasound guidance reduced the number of attempts, time to access, risk of accidental venipuncture and vascular complications (i.e., fewer hematomas) when obtaining femoral arterial access. However, the study found that ultrasound guidance improved cannulation rates only in patients who had a high common femoral artery bifurcation compared with standard fluoroscopic guidance [38].

**Type of anticoagulant & antiplatelet agents used**

Vascular access-site complications are associated with the use of anticoagulant and antiplatelet agents. The use of GP IIb/IIIa inhibitors, particularly abciximab, has been shown to increase vascular access-site complications [39,40]. The combination of abciximab and unfractionated heparin was associated with an increased risk of bleeding in the EPIC trial [41]. The combination of abciximab and low-dose weight-adjusted heparin conferred substantial clinical benefit without increased bleeding complications in the EPILIG trial [42]. The reversal of abciximab’s antiplatelet action is slow. On the contrary, eptifibatide and tirofiban selectively bind to the GP IIb/IIIa receptor and their effect on platelets is rapidly reversed when discontinued. If bleeding occurs, the effects of eptifibatide and tirofiban can be reversed with platelet transfusion [43]. Bivalirudin binds specifically and reversibly to both fibrin bound and unbound thrombin [44]. In the REPLACE-2 trial, bivalirudin with selective GP IIb/IIIa inhibition provided similar protection from periprocedural ischemia and was associated with less major bleeding compared with low-dose heparin plus GP IIb/IIIa inhibition [24]. Bivalirudin was associated with similar rates of bleeding and ischemia as heparin in patients with moderate or high-risk acute coronary syndromes who were undergoing intervention with GP IIb/IIIa inhibitors. Bivalirudin alone was associated with significantly lower rates of bleeding and similar rates of ischemia [45]. Compared with heparin plus GP IIb/IIIa inhibition, bivalirudin alone resulted in significantly reduced 30-day rates of major bleeding and major adverse cardiovascular events in patients with ST-segment elevation myocardial infarctions undergoing primary PCI [46]. This was true despite the significantly higher peak activated coagulation time among patients treated with bivalirudin.
**Arterial closure devices**

Arterial closure devices were initially designed as active closure methods. The two main types of active arterial closure devices are collagen plug closure devices (e.g., AngioSeal™; St. Jude Medical) and suture-mediated closing devices (e.g., Perclose™; Abbott Vascular) [10,47]. StarClose™ (Abbott Vascular) is a newer active closure method involving surgical staple/clip technology. Passive closure devices focus on enhanced manual compression utilizing assisted compression with mechanical clamps (e.g., FemoStop™; St. Jude Medical), external patches or plugs (e.g., the Mynx™ Vascular Closure Device [AccessClosure] and the ExoSeal™ device [Cordis]) and wire-stimulated track thrombosis (e.g., the Boomerang™ Catalyst Wire; Cardiva Medical). Some of these devices have the disadvantage of not inducing immediate hemostasis [48]. An overview of closure devices and their main features is provided in Table 1 [6,49–54].

Some of the complications that arise from the use of arterial closure devices are similar to complications with manual compression: bleeding (from incomplete closure), hematoma, pseudoaneurysm (from a persisting arterial leak) and AV fistulas. Other complications include thrombus formation, embolization of the collagen plug or clip, late stenosis of the artery (which can lead to limb ischemia), compression of the femoral vein access site infections, septic emboli, femoral endarteritis and limb ischemia [6,10,55]. Infectious complication rates with closure devices have been observed to be anywhere from 0.0 to 5.1%, with *Staphylococcus aureus* being the most common infectious agent followed by coagulase-negative *Staphylococcus* [55]. The foreign material in the intravascular space and arterial wall may serve as a nidus for infection. The median time from arterial closure device usage to site infection is 8 days. Mycotic pseudoaneurysm is the most common complication of infection from arterial closure devices.

Arterial closure devices are generally safe and allow for earlier mobility, improved patient comfort, earlier discharge, potentially reduced healthcare costs and arguably reduced groin complications (particularly hematomas and pseudoaneurysms). They are effective in small arteries and vessels affected by connective tissue disorders [6,15].

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<tr>
<th>Closure device</th>
<th>Mechanism</th>
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<tr>
<td>AngioSeal™</td>
<td>Utilizes a collagen plug and anchor mechanism. A suture cinches the anchor and collagen component to form a secure seal</td>
<td>Collagen plug breaks down over months and anchor is absorbed by hydrolysis over months Similar or lower complications as manual compression</td>
</tr>
<tr>
<td>Perclose™</td>
<td>A suture on each side of the arterial puncture site is tied using a preloaded knot tying device. Knot is cinched and suture is cut close to the arterial wall</td>
<td>Suture material is not absorbable</td>
</tr>
<tr>
<td>StarClose™</td>
<td>Extravascular clip deployed through peel-away sheath onto the extravascular arterial surface, achieving closure through primary healing</td>
<td>Similar major vascular complications and fewer minor complications as manual compression The clip is not absorbable</td>
</tr>
<tr>
<td>FemoStop™</td>
<td>A belt and air pressure filled bulb is used to compress the artery</td>
<td>Complications result from inadequate compression time or poorly placed compression Requires significant staff time Applying device for too long can result in increased risk of arterial and venous thrombosis and injury to femoral nerve</td>
</tr>
<tr>
<td>Boomerang™</td>
<td>Percutaneous vascular closure device that promotes hemostasis via intravascular tamponade with complete removal of the system after hemostasis is achieved</td>
<td>Requires significantly shorter staff time than manual compression</td>
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<tr>
<td>Mynx™</td>
<td>Utilizes a water soluble polyethylene glycol sealant that expands inside the tissue tract by absorbing blood and subcutaneous fluids to provide a seal over the arteriotomy site and within the tissue tract</td>
<td>Sealant is fully reabsorbed within 30 days Avoids cinching or tugging of the artery</td>
</tr>
<tr>
<td>ExoSeal™</td>
<td>Visually guided deployment mechanism that delivers a bioabsorbable polyglycolic acid plug atop the femoral artery</td>
<td>Plug is hydrolyzed within 3 months Avoids cinching or tugging of the artery</td>
</tr>
<tr>
<td>Femoral Introducer Sheath and Hemostasis (FISH)™</td>
<td>FISH uses an extracelular matrix closure patch premounted on a 5, 6, or 8 Fr access sheath</td>
<td>Serves as an access and closure device Patch is resorbed within 90 days</td>
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Taken from [6,49–54].
Earlier experience with arterial closure devices revealed higher vascular complication rates than with manual compression [56]. A meta-analysis conducted by Koreny et al. in 2004 found that time to hemostasis was shorter with arterial closure devices, but they potentially conferred a higher risk of hematoma and pseudoaneurysm formation [57]. Another meta-analysis in 2004 conducted by Nikolsky et al. showed that vascular complications were similar with AngioSeal, Perclose, and VasoSeal™ (Datascope Corporation) compared with mechanical compression for diagnostic cardiac catheterization; vascular complications with AngioSeal and Perclose were similar to mechanical compression for PCI, but complication rates were higher with VasoSeal compared with mechanical compression [58]. Arora et al. found that arterial closure devices reduced the risk of vascular complications compared with manual compression in patients undergoing both diagnostic angiography and PCI [59]. Newer generations of vascular closure devices and more operator experience are likely reducing the vascular complication rates and may be reducing cost. One study found that the routine use of AngioSeal was associated with a lower cost per PCI procedure compared with manual compression [60]. A study comparing AngioSeal to Mynx found that the rates of major vascular complications were not significantly different between the two devices, but Mynx was associated with a higher rate of device failure [61]. The main advantage of vascular closure devices used after diagnostic coronary angiography is shortening the time to ambulation. This may become less relevant as more procedures are performed using the radial access route. After PCI, where the patient is at much higher risk of bleeding, there may be some advantage with the use of closure devices, although transradial PCI may change this paradigm as well.

**Diagnosis**

The diagnosis of the various vascular complications is based on history and physical examination findings and can typically be confirmed by imaging. Ultrasound is usually the initial imaging procedure in the evaluation of patients with suspected minor vascular access-site complications as it can accurately and reliably diagnose AV fistulas and pseudoaneurysms [61]. Femoral angiography performed during catheterization can assist in identifying complications and some operators advocate performing a femoral angiogram for every case. An overview of the clinical findings and secondary complications of groin access complications is presented in Table 2 [9–12,15,62].

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**Treatment of vascular access-site complications**

Hematomas can progress to shock in some cases. If a hematoma develops ensued by hypotension during catheterization, the femoral artery may have been nicked. A larger sheath size should be utilized and if this is not successful, then manual pressure should be applied. Anticoagulation should be stopped and reversed and hypotension should be managed with fluids and blood products as needed. If bleeding cannot be controlled with manual compression, the bleeding site can be localized with angiography with a sheath inserted in the contralateral femoral artery and potentially tamponaded with balloon inflation [10].

Small pseudoaneurysms, less than 3 cm, can typically be observed as they can clot spontaneously over time (within 4 weeks). Ultrasound-guided compression can compress the site of communication between the pseudoaneurysm and the native artery and allow thrombosis of the aneurysmal sac. A FemoStop device can also be used to compress a pseudoaneurysm. Ultrasound-guided percutaneous thrombin injection into the pseudoaneurysm can cause thrombosis of the pseudoaneurysm within a few seconds after the injection. However, this method has the risk of thrombosis of the native artery and in some cases the femoral vein. Thrombin injection appears to be more effective than ultrasound-guided compression for femoral pseudoaneurysms. Other methods that have been investigated include para-aneurysmal saline where saline is injected into the communication between the false aneurysm and the native artery resulting in elimination of blood flow. Endovascular techniques such as coil embolization, detachable balloons and stent-grafts have also been used to treat femoral pseudoaneurysms.

Surgery is the gold standard treatment for pseudoaneurysms requiring treatment. Indications for surgical treatment include rapid expansion of the pseudoaneurysm, concomitant distal ischemia or neurological deficit, mycotic pseudoaneurysm, failure of percutaneous intervention and compromised soft tissue viability. Surgical options include simply suturing the defect after hematoma evacuation or by patch angioplasty where surgical repair is achieved by using a patch [11,12,28,63,64].

Retroperitoneal hemorrhages can typically be treated conservatively with fluid administration, blood transfusions, and discontinuation and reversal of antiplatelet and anticoagulant agents. This is typically sufficient in many patients as accumulated blood will tamponade the site of bleeding. Ongoing active bleeding should prompt immediate vascular surgery consultation. Endovascular
methods are available for treating the bleeding. The main endovascular method is balloon tamponade [65]. Other endovascular methods include selective intra-arterial embolization or stent-grafting. Embolization can use a combination of agents including coils, gelatin, and polyvinyl alcohol, and these agents should be placed proximal and distal to the site of bleeding [11]. One proposed indication for embolization was based on hemodynamic instability and degree of blood loss (requiring four or more units of blood within a 24-h period or six or more units within a 48-h period) [66]. Surgical intervention is indicated for unstable patients not responding to fluid and blood product administration if interventional radiology is not successful or available, or for the development of abdominal compartment syndrome due to a large retroperitoneal hemorrhage, or for femoral neuropathy [11,66]. In one study, 16% of patients who developed a retroperitoneal hemorrhage required surgical intervention [67].

AV fistulas can be treated conservatively by monitoring in most patients. They are generally small, not hemodynamically significant and close spontaneously. When AV fistulas become symptomatic, they require closure to prevent increased shunting and distal swelling and tenderness. Options include prolonged bandaging, ultrasound-guided compression, percutaneous implantation of covered stents, percutaneous coil embolization and surgical repair. In many cases, the fistula may be too short or long to be compressed effectively by ultrasound. Surgical intervention is generally reserved for patients who fail a less invasive approach [9,11,15].

If patients develop lower extremity ischemia, angiography should be obtained to characterize the anatomic basis of ischemia. Blood flow can be restored by balloon angioplasty with or without thrombolytic therapy, stents or catheter thrombectomy. Surgical thrombectomy and repair may be required if percutaneous methods are not successful [9]. When arterial dissection is identified, angiography should be obtained to characterize the extent of dissection. Removal of catheters and wires could potentially allow spontaneous resolution of the dissection when identified. If catheter removal is unsuccessful, treatment options include balloon angioplasty, endovascular stent placement or surgical repair to stabilize a flow limiting dissection [9,10]. Treatment of arterial thrombosis consists of thrombectomy performed

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<td>Retroperitoneal hemorrhage</td>
<td>Hypotension, Back, flank, hip or abdominal pain, Anemia</td>
<td>Computed tomography, Arteriography, Local exploration in the operating room</td>
<td>Anemia, Neuropathy, Mortality rate of 6.6%</td>
</tr>
<tr>
<td>Thrombosis or embolism</td>
<td>Pain, Pallor, Parasthesia, Pulselessness, Poikilothermia</td>
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Table 2. Clinical findings and secondary complications of groin access complications.

Taken from [9–12,15,26,62].
either percutaneously or surgically. Thrombolytic therapy with tissue plasminogen activator can be considered if a large thrombus burden persists after thrombectomy [15].

Clinical consequences of vascular access-site complications

Bleeding complications during cardiac catheterization have been found to result in longer hospital stays, additional treatments and higher healthcare costs. Yatskar et al. found that patients undergoing PCI who developed hematomas requiring transfusion had a fivefold increase in in-hospital death and a fourfold increase in 1-year mortality [16]. Kinnaird et al. studied patients between 1991 and 2000 and found that major and minor bleeding complications were associated with an increase in in-hospital death and major adverse cardiac events [68]. Blood transfusions were an independent predictor of in-hospital and 1-year mortality. Patients with major bleeding complications had an average hospital stay of 8.9 days compared with 3.1 days for those without any bleeding complications. Patients undergoing PCI from 1992 to 2003 who developed a retroperitoneal hemorrhage had a higher rate of transfusion, increased length of hospital stay and higher mortality rate [1]; this may be due to the different anticoagulation regimen used at that time. Bleeding is a complication with economic implications as well: the increase in cost was found to range from US$6022 to US$11,767 for each bleeding event [69]. Major vascular access-site complications can lead to the discontinuation of dual antiplatelet therapy in patients who have had recent PCI. Discontinuation of antiplatelet therapy in patients with minor vascular access-site complications is not always necessary and can result in other complications such as recurrent ischemia, myocardial infarction and stent thrombosis [69].

Other vascular access-site complications

Infections and neurologic complications can also occur. The frequency of such complications is not well defined, perhaps because they are rarely observed. Infectious complications include local abscess or cellulitis and sepsis. Less than 1% of patients develop local infection at the site of femoral arterial access and the most common organisms isolated are Staphylococcus aureus and Staphylococcus epidermidis. Neurologic complications include femoral neuropathy, foot drop, paresthesia and chronic pain. Femoral neuropathy occurs in approximately 0.2% of patients. It is typically seen in association with a large hematoma or pseudoaneurysm, which causes pressure on the femoral nerve. When caused by a hematoma or pseudoaneurysm, pressure is exerted on the femoral cutaneous nerve resulting in a sensory neuropathy; this tends to resolve when the pressure is relieved. When caused by retroperitoneal bleeding, pressure is exerted on the femoral nerve resulting in weakness in the upper leg and pain in the thigh and calf; this may not be entirely reversible [4].

Conclusion

Vascular access-site complications continue to pose a significant challenge for invasive and interventional cardiologists. The preventive measures discussed earlier offer promise for reduction of such complications and improvement in patient outcomes.

Future perspective

Fortunately, the incidence of bleeding complications appears to be decreasing. The incidence of major bleeding complications after PCI at the Mayo Clinic decreased from 8.4% in 1994–1995 to 3.5% in 2000–2005 [55]. This decrease is likely due to a decrease in the mean sheath size (from 8.2-F to 6.4-F), decreased use of a venous sheath (13% to 6%), decreased intensity of anticoagulation with heparin as assessed by peak activated coagulation time (405 s to 312 s), decreased use of postprocedure heparin (80% to 27%) and a decrease in procedure time (1.7 h to 1.4 h) [69]. This decrease in major bleeding complications was found despite the increased use of GP IIb/IIIa inhibitors and procedures performed on older patients. Advancements in anticoagulant and antiplatelet agents and newer generations of arterial closure devices offer some promise in decreasing vascular access-site complications. Alternative access sites, particularly the radial approach, are gaining favor and could result in further decreases in complication rates.

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Vascular access-site complications of cardiac catheterization range from minor complications such as stable hematomas and pseudoaneurysms to major conditions including arteriovenous fistulas, hematomas requiring transfusion, retroperitoneal hemorrhage, arterial dissection, embolism, thrombosis, vessel rupture and limb ischemia.

The incidence of vascular access-site complications is generally higher for interventional procedures than for diagnostic procedures. Risk factors for vascular complications of cardiac catheterization can be divided into patient-related and procedural-related risk factors. Newer generations of arterial closure devices have been shown to reduce vascular access-site complications.


Access complications in the catheterization laboratory


Analysis of the economic advantages of angio-seal for vascular closure.


Study reporting the decreasing incidence of femoral access complications over the past decade.