External iliac artery pseudoaneurysm in patient with history of failed kidney transplant

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CASE SUMMARY

A 66-year-old African-American male presented with a four-month history of asymptomatic hematuria. Interventional Radiology was consulted for embolization of bilateral native arteries in this patient on hemodialysis. His past medical history was significant for end-stage renal disease secondary to polycystic kidney disease. Patient underwent right kidney transplantation 21 years ago, which ultimately failed 15 years ago.

IMAGING FINDINGS

CT scan of the pelvis showed multiple avidly enhancing lesions of various sizes throughout bilateral native kidneys (Figure 1). Subsequently, angiography was done in order to evaluate for embolization. A digitally subtracted oblique image of the right external iliac artery demonstrated a pseudoaneurysm (Figure 2). Finally, a digitally subtracted image after covered stent placement showed no pseudoaneurysm filling (Figure 3).

DIAGNOSIS

Multiple bilateral hypervascular masses were suspected to be the source of the hematuria. Angiogram also showed an incidental finding of a pseudoaneurysm in the right external iliac artery.

DISCUSSION

A pseudoaneurysm occurs due to a disruption in an arterial wall. The high-pressure arterial blood stretches the weakened arterial wall and creates a communicating sac within the lumen of the artery. The sac is surrounded by the adventitia and media of the vessel. There are many causes of pseudoaneurysms with the most common being due to inflammation, iatrogenic surgical or catheterization injuries, and trauma.¹ There have also been a few rare incidences where pseudoaneurysms have been discovered in patients with failed renal transplants. One particular study looked at 843 renal transplants performed over a 37-year period. Only three, or 0.35 percent, of those patients developed a pseudoaneurysm at the anastomotic site.² This is the case in the patient presented in this report. The incidental pseudoaneurysm discovered in this patient was at the site of anastomoses of the right external iliac artery to the graft renal artery.

It is also important to note that this patient's pseudoaneurysm was not associated with some of the common symptoms that are seen in patients with pseudoaneurysms. These common symptoms include but are not limited to pain, swelling, pulsatile mass, and symptoms associated with nerve and vein compression, such as venous thrombosis and neuropathy.³

In the few instances that a pseudoaneurysm has been described at the anastomotic site of a failed renal transplant, the discovery of the pseudoaneurysm has been within several weeks of transplantation or at the time of failure. In one particular case, bruits were heard over an anastomotic site only five weeks after transplantation. Further investigation using arteriography in this

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right external iliac artery demonstrating pseudoan-

eurysm in the superior lateral aspect of the image.



FIGURE 1. Multiple avidly enhancing lesions are FIGURE 2. Digitally subtracted oblique image of the noted throughout bilateral native kidneys.

particular study showed a 15 mm by 10 mm pseudoaneurysm on the end-to-end anastomosis between the internal iliac artery and renal artery.² In another case, a pseudoaneurysm and transplant rejection were discovered in the internal iliac artery anastomosis seven weeks after transplantation.4

The patient in this case had a unique presentation. Angiography used to aid in the discovery of the site of the bilateral hypervascular renal masses thought to be the source of hematuria led to the incidental finding of a 3.9 cm by 2.3 cm pseudoaneurysm at the site of anastomosis between the right external iliac artery and graft renal artery. Our patient did not have any signs or symptoms that were directly caused by the pseudoaneurysm. The patient did not have pain in the area, swelling, pulsatile mass, or bruits. Also, our patient's case was unusual compared to the two cases previously mentioned in that our patient was presenting 21 years after transplantation and 15 years after rejection. Most published research describes pseudoaneurysm and its complications being discovered within several weeks after transplantation or failure. There is no definitive way of confirming how long the pseudoaneurysm had been present in this patient. However, based on the literature where pseudoaneurysms

were discovered along with the failure of the renal graft, it is to be assumed that the pseudoaneurysm was present in our patient for 15 years. Because of this, a prompt therapeutic approach was undertaken. There is much debate on how or whether to treat asymptomatic pseudoaneurysms. Due to the uncertainty of our patient's pseudoaneurysm, it was determined that a placement of a covered stent in the right external iliac artery would be the best option.

In the procedure, a 10 mm by 40 mm covered stent was extended beginning just distal to the take-off of the right internal iliac artery. An arteriogram after the deployment of the 10mm x 40mm covered stent showed persistent blood flow within the pseudoaneurysm with take-off near the distal aspect of the stent. An additional 10mm x 80 mm covered stent was then deployed beginning distal to the take-off from the right internal iliac artery, landing proximal to the takeoff of the right inferior epigastric artery. Post stent deployment angiogram of the right external iliac artery demonstrated no flow into the pseudoaneurysm.

CONCLUSION

Our patient had an unusual presentation of a pseudoaneurysm. Pseudoaneurysms associated with failed

FIGURE 3. Digitally subtracted image showing covered stent placement with no pseudoaneurysm filling.

renal transplants are typically detected within weeks after transplantation and along with failure of the transplant. Our patient's pseudoaneurysm was detected 21 years after transplantation and 15 years after transplant failure. Rupture of a pseudoaneurysm can occur at any time, and treating it with a covered stent is a feasible option.

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