Report of a rare fistula between a Studer neobladder and external iliac artery

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Abstract

A neobladder-arterial fistula is a very rare complication following cystectomy, with only 1 previously reported case. Delay in diagnosis can be rapidly fatal and requires prompt intervention. We report the case of a 63-year-old male who developed massive hematuria, and was found to have a fistula between the right external iliac artery and Studer neobladder during emergent exploratory laparotomy. Treatment success relies on a high index of suspicion and may include open operative intervention.

Background

Urovascular fistulas are well-documented in the urologic literature, although only in small case series. These usually occur between the ureter and iliac artery in patients with predisposing risk factors, such as prior pelvic surgery, radiation, chronic ureteral stents and other vascular pathologies.1 Treatment options vary widely with respect to open versus endovascular repair. While ureterovascular fistulae are well-described, to our knowledge there has been only 1 account of a fistula between a neobladder and artery, and endovascular management was used in this setting.2 We are reporting the second case of a neobladder-vascular fistula that was alternatively managed with an open surgical approach.

Case report

A 63-year-old male with a history of muscle invasive bladder cancer was transferred to our medical centre for a suspected ruptured neobladder. Eight months prior to this encounter, he underwent a radical cystectomy, extended bilateral pelvic lymphadenectomy and a Studer ileal neobladder, a urinary diversion using 55 cm of ileum.3 In the month prior to transfer, he had been hospitalized twice at outside hospitals for episodes of gross hematuria. In the first occurrence, he was managed as an outpatient with antibiotics for a suspected neobladder urinary tract infection. The subsequent episode required hospital admission and Foley catheter placement. While continuous bladder irrigation, as well as manual hand irrigation of blood clot, was required initially, this subsided after a few days. The evening prior to arrival at our institution, the patient developed acute abdominal distention, pain and respiratory distress. A peritoneal tap revealed urinary ascites, and he was immediately transferred for further management. Labs on admission included a serum creatinine of 5.6 mg/dL (427 mmol/L), which was presumed to be secondary to reabsorption of urine from the peritoneal cavity. Operative exploration revealed a 2-cm laceration in the afferent limb of the ileal neobladder, likely due to traumatic neobladder catheterization/instrumentation, which was repaired in 2 layers. The afferent limb consists of a 10- to 15-cm ileal chimney used for ureteral anastomosis and serves an anti-reflux role. Endoscopy of the neobladder was performed at that time as well; however, no sources of previous hematuria episodes were identified. During routine postoperative care, the patient’s urine remained clear. On postoperative day 7, nearing hospital discharge, he experienced an acute loss of consciousness and hemorrhagic shock preceded by massive and rapid gross hematuria. The Foley catheter that was in place since surgery was clamped to tamponade active bleeding. This allowed time for transfer to the intensive care unit, central line placement, and volume resuscitation. The patient received 4 units of packed red blood cells through a rapid infuser and follow-up labs revealed a hemoglobin and lactate of 7.2 g/dL and 5.1 mmol/L, respectively. Once hemodynamically stable, he was taken emergently to the operating room for re-exploration. No diagnostic imaging was obtained since the clinical picture strongly suggested a large vascular fistula.
Initially, intraoperative cystoscopy was attempted to localize the bleeding source. However, this was quickly abandoned due to inadequate visualization from a large clot burden. Subsequent laparotomy revealed a tense and distended neobladder, but no evidence of bleeding into the peritoneal cavity. The afferent limb of the neobladder was opened at the same site that was repaired 1 week previously, and no acute source of bleeding was identified immediately in the neobladder. After clot evacuation, pulsatile blood flow was seen emanating from the right side wall of the neobladder, immediately adjacent to the right external iliac artery. The relative location was identified on a CT scan obtained several days prior to this incident (Fig. 1). Proximal and distal vascular control was obtained on the external iliac artery and the fistula tract was dissected. A ruptured atheromatous plaque was identified in contiguity with a hole in the neobladder. The diseased arterial segment was resected and the artery re-anastomosed primarily without graft. The neobladder was closed primarily in 2 layers, and a pedicled flap of omentum was positioning between the right external iliac artery and the neobladder. Intra-operatively, the patient received 3.8 L of packed red blood cells in addition to 1.2 L of other blood products, following our institutional massive transfusion protocol. Despite this, the postoperative labs revealed a hemoglobin of only 8.7 g/dL, emphasizing the degree of acute blood loss that had occurred. The patient recovered well and was discharged 9 days later without any lower extremity abnormalities or compartment syndrome. At the 10-month follow-up, he reported complete continence during the day with mild nighttime leakage, and there have been no untoward sequelae from this encounter.

**Discussion**

A fistula, by definition is an abnormal connection between 2 or more epithelial-lined organs or vessels. Fistulas between the vascular and urinary systems are not common, but have been described in several case series. The exact pathology is not well-understood; however, inflammation, fibrosis, ischemia and necrosis are the final common pathway for a multitude predisposing conditions, such as pelvic surgery, radiation therapy, vascular surgery and chronic ureteral stents. A fistula, by definition is an abnormal connection between 2 or more epithelial-lined organs or vessels. Fistulas between the vascular and urinary systems are not common, but have been described in several case series. The exact pathology is not well-understood; however, inflammation, fibrosis, ischemia and necrosis are the final common pathway for a multitude predisposing conditions, such as pelvic surgery, radiation therapy, vascular surgery and chronic ureteral stents.4

Ureterovascular fistulas represent most of the fistulas reported between the urinary and vascular systems. Still, the most recent review in 2008 reports only 139 cases dating back to 1939.5 Because they are so uncommon, patients presenting with recurrent bouts of gross hematuria often go undiagnosed for several weeks. This likely contributes to mortality rates as high as 58% if untreated.5 Less commonly reported are cases of fistulas between urinary conduits and arteries.6-10 In a review of 80 cases of ureterovascular fistulas, Bergqvist and colleagues reported that 36% occurred in the setting of ileal conduits, suggesting even when a urinary diversion is present, a fistula to the ureter is more likely.11

Certainly, the least reported type of fistula is one between a neobladder and artery. In 2006, Gallucci and colleagues described a case of a 62-year-old male who, following an episode of hematuria and clot urinary retention, was diagnosed with an arterio-neobladder fistula by angiography.2 This was subsequently managed with an endovascular graft under local anaesthesia, and no blood transfusions were required. The long-term outcomes from this case were not reported.

In patients that are hemodynamically stable, endovascular approaches have been successful in several cases of ureterovascular fistulas.12,13 However, a very relevant difference is highlighted in the present case, where the bowel is in direct continuity with the fistula tract. In this situation, placement of a prosthetic graft should be avoided due to a high graft infection risk, as bowel diversions are chronically colonized. In the Mayo Clinic series of ureterovascular fistulae in 7 patients, Krambeck and colleagues recommended that endovascular stenting be used only as a temporizing measure or poor surgical candidate.4 We would agree with this recommendation, and would argue that open surgical repair is the preferred approach in the case of a neobladder-arterial fistula. Cystoscopy may be attempted first to localize the fistula to a particular ureter, but this may prolong definitive treatment and should be avoided in unstable patients. The underlying conditions that lead to fistula formation are often not modifiable, thus consideration should be given to placing a tissue flap over the arterial repair to serve as an interposing layer. This may prevent recurrence of an arterial-conduit fistula as recently reported by Hori and Thiruchelvam.8 Additional surgical treatment options include vessel ligation and subsequent extra-anatomic bypass. This method circumvents many of the predating risk factors altogether, and is optimal for
patients who will require continued chronic ureteral stents to maintain renal function.

As there are few occasions when the urologist is exposed to such a high acuity, stress-provoking and uncontrolled hemorrhage, the importance of having a broad knowledge of possible etiologies cannot be overstated. The presentation of our patient is very similar and even characteristic of this disease process. Gross hematuria is the sine qua non manifestation; however, this can be mild and intermittent (i.e., herald bleeding), often indicating a more common diagnosis. Batter and colleagues reported intermittent gross hematuria for up to 3 weeks prior to massive hematuria in patients ultimately diagnosed with urovascular fistulas.\(^1\) In the setting of a urinary diversion, gross hematuria generally requires some evaluation. Etiologies that are more common than fistulae include cancer recurrence (upper tracts or urethra), stones, and infection. However, these processes rarely cause the massive bleeding that can be encountered with vascular fistulas.

Unfortunately, imaging modalities have not been a reliable strategy for the diagnosis of these rare events, which is likely associated with the intermittent nature of these processes. At the initiation of the tract formation, natural clotting mechanisms may prevent frank exsanguination for a period of time. Therefore, unless active extravasation is occurring, you will not be able to identify an abnormality radiographically. Provoking this temporary clot by manipulating a ureteral stent simultaneous to angiography has been advocated endovascularly. This may be required in the absence of radiographic diagnosis, that culminate in life-threatening hemorrhage. Management requires a multidisciplinary approach, including visits with a urologist, vascular surgeon and interventionist, for successful management and long-term success.

**Conclusion**

Cystectomy patients are at risk for urovascular fistulas. These usually occur between an artery and ureter, but can also involve a direct connection to an orthotopic urinary diversion, as we have reported. As such, episodes of gross hematuria need to be evaluated appropriately, at least to exclude more common causes. This population, which are largely managed with bowel diversions, provides a unique challenge to endovascular approaches, and should be treated with an open repair in most cases. Presentation is often characterized by days to weeks of intermittent gross hematuria that culminate in life-threatening hemorrhage. Management may be required in the absence of radiographic diagnosis, and a high index of suspicion is paramount in this setting. Treatment should be targeted toward patient stabilization, definitive treatment of the lesion, and prevention of recurrence by incorporating omental flaps when indicated. This requires a multidisciplinary approach, including visits with a urologist, vascular surgeon and interventionist, for successful management and long-term success.

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**References**


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