Left Anterior Descending Coronary Artery and Multiple Peripheral Mycotic Aneurysms Due to Mycobacterium Bovis Following Intravesical Bacillus Calmette-Guerin Therapy: A Case Report

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ABSTRACT

The use of live attenuated intravesicular Bacillus Calmette-Guerin (BCG) therapy is a generally accepted safe and effective method for the treatment of superficial transitional cell carcinoma (TCC) of the bladder. Although rare, < 5% of patient's treated with intravesicular BCG therapy may develop potentially serious complications, including localized infections to the genitourinary tract, mycotic aneurysms and osteomyelitis. We present here a case of a 63-year-old male who developed left coronary and multiple peripheral M. Bovis mycotic aneurysms as a late complication of intravesicular BCG therapy for superficial bladder cancer. The patient initially presented with acute onset pain and swelling in the left knee > 2 years following initial therapy, and initial workup revealed a ruptured saccular aneurysm of the left popliteal artery as well as incidental bilateral common femoral artery aneurysms. Following endovascular treatment and additional workup, the patient was discovered to have additional aneurysms in the right popliteal artery and left anterior descending artery (LAD). Surgical pathology and bacterial cultures obtained from the excised femoral aneurysms and surgical groin wounds were positive for Mycobacterium Bovis, and the patient was initiated on a nine-month antimycobacterial course of isoniazid, rifampin and ethambutol. Including the present case, there has been a total of 32 reported cases of mycotic aneurysms as a complication from intravesicular BCG therapy, which we will review here. The majority of reported cases involve the abdominal aorta; however, this represents the first known reported case of a coronary aneurysm.

CASE REPORT

A 63-year-old male with a 45-pack year smoking history presented to his primary care provider with gross hematuria. Subsequent urological referral and cystoscopy revealed a left lateral bladder mass with suspicious cytology. CT urography was negative for upper tract involvement and the patient underwent transurethral resection of the bladder tumor (TURBT) with final pathology consistent with high grade T1 transitional cell carcinoma (TCC). The patient underwent six
cycles of induction BCG therapy from November 2012 through January 2013 and ten cycles of maintenance therapy from March 2013 through August 2014, which he tolerated well during that period.

In March of 2015, the patient presented to the Emergency Department (ED) after the acute onset of left knee pain and swelling. A venous Doppler ultrasound (US) was performed in the ED, demonstrating a complex vascular mass arising from the popliteal artery consistent with a partially thrombosed popliteal aneurysm [Figure 1a, b]. In addition, a partially thrombosed left common femoral aneurysm was identified [Figure 1 c, d]. He received a CTA immediately afterwards, which demonstrated a 5.5 x 6.5 cm ruptured left popliteal aneurysm [Figure 2a] and incidental bilateral common femoral artery (CFA) aneurysms [Figure 2b, c]. Mild scattered calcified atherosclerotic plaque was noted in the inflow vessels, without significant stenosis. The patient was stabilized and subsequently underwent left popliteal artery covered stent placement by vascular surgery and interventional radiology the following day. He had an uneventful post-operative course and was discharged shortly afterwards with future plans for elective repair of the bilateral CFA aneurysms.

In May of 2015, the patient returned to the ED with acute onset pain and swelling behind the right knee. Doppler US demonstrated a ruptured right popliteal aneurysm in addition to enlarging bilateral CFA aneurysms [Figure 3]. He received endovascular repair of the ruptured popliteal aneurysm with a covered stent the following day, followed by open surgical repair of the CFA aneurysms with bilateral interposition grafts the following week [Figure 4, 5]. Pathology results from the excised femoral aneurysms demonstrated the presence of acid bacilli on AFB stains, consistent with mycobacterial infection. The post-operative course was prolonged and complicated by non-healing bilateral surgical groin wounds and progressive weight loss. On post-op day 12, the patient also developed fevers and chills. These developments led to an extensive but unremarkable workup for potential sources of infection and occult malignancy, including a negative CT of the chest, abdomen and pelvis and multiple negative blood cultures. One month post-op, the patient underwent bilateral groin washout with wound VAC placement. Wound cultures obtained at the time of surgery, were positive for Mycobacterium Bovis, consistent with the previous pathology findings. Given the diagnosis of M. Bovis induced mycotic aneurysms, the patient underwent CTA of the head, neck, chest, abdomen and pelvis to screen for additional aneurysms, which were reported as negative for additional aneurysms. Infectious Disease was consulted, who initiated a 9-month course of anti-mycobacterial treatment with Isoniazid, Rifampin and Ethambutol.

Two months following the initial diagnosis, the patient underwent surveillance CTA of the lower extremities to evaluate patency of the surgical interposition grafts. The results demonstrated patent surgical grafts; however, enlarging mycotic aneurysms were discovered in the proximal right common iliac artery and at the distal attachment site of the right surgical interposition graft [Figure 6]. The following week, the patient underwent endovascular repair with a bifurcated distal aortic stent graft and coil embolization of the right common iliac aneurysm [Figure 7]. There was no evidence of new or enlarging aneurysms on 2-month follow-up CTA [Figure 8].

Over the next month, the patient had complaints of increased pain and swelling behind the right knee. CTA of the lower extremities was performed which showed an enlarging hematoma in the right popliteal fossa [Figure 9a, b]. In addition, there appeared to be an enlarging partially thrombosed aneurysm arising from the mid LAD, which, in retrospect, was present on screening CTA performed two months earlier [Figure 9 c, d]. The patient subsequently received a dedicated coronary CTA to further evaluate this new finding. Results demonstrated a large saccular aneurysm with sluggish internal flow arising from the mid LAD with resultant mass effect on the left ventricular outflow tract [Figure 10]. In lieu of the newly discovered LAD aneurysm, cardiothoracic surgery was consulted who deemed that no surgical intervention was warranted at that time due to the poor functional and nutritional status of the patient. Alternatively, a coronary catheter angiogram was performed by interventional cardiology to evaluate the possibility of stent placement; however, stent placement was precluded by occlusion of the mid LAD at the aneurysm site [Figure 11].

Given the rapid interval growth of the LAD mycotic aneurysm and the patient’s poor functional status, his long-term prognosis was deemed poor. As such, the patient was placed on the palliative care service and converted to a DNR status. He continues to receive antimycobacterial therapy in attempt to prevent interval mycotic aneurysm growth and formation. Currently the patient is undergoing subacute rehabilitation with the aims of improving his functional and nutritional status for the future possibility of potential surgical repair.

**DISCUSSION**

**Etiology & Demographics:**

_Bacillus Calmette-Guerin (BCG)_ is a live attenuated strain of Mycobacterium bovis that was initially developed as a vaccination against Mycobacterium tuberculosis infection. In 1976, BCG was introduced as a primary treatment for superficial transitional cell carcinoma (TCC) of the urinary bladder with a reported cure rate of 70% [1]. Intravesical BCG therapy is generally considered safe with the most common reported side effects of low-grade fever, dysuria, urinary frequency and malaise with a typical onset within 2-4 hours following therapy and resolving within 48 hours [2, 3, 24, 29]. In < 5% of cases, more serious localized or disseminated infections due to BCG may occur. The largest reported series followed 2,602 patients who received BCG therapy. In this group, granulomatous prostatitis occurred in 0.9% of patients, hepatitis occurred in 0.7% of patients and pneumonitis in 0.7% of patients [3]. The onset of symptoms following initial treatment is variable and some authors have divided complications into early (< 6 months) and late complications (> 6 months) [4, 5]. A retrospective review of 41 patients who developed treatment complications...
demonstrated that slightly more than half (25/41) of patients were considered early-onset and tended to manifest with non-specific systemic symptoms (fever, chills, malaise and weight loss) due to an inflammatory response. On the contrary, the remaining 16/42 patients who developed late-onset disease tended to develop more localized infections due to BCG reactivation, with the most common site involving the genitourinary tract followed by the vascular system, skeleton and retroperitoneal soft tissues [4, 24].

The incidence of BCG-induced mycotic aneurysms following intravesicular infusion is exceedingly rare. Including the current case, there have only been 32 reported cases, with all but one occurring in middle-aged or elderly men [Table 3]. The majority of reported cases (26/32) involve the thoracic or abdominal aorta [6-33, Table 1]. Approximately one-fourth of reported cases involve the peripheral vascular system of the lower extremities (9/32) and there have been three reported cases involving the carotid arteries [Table 1]. To our knowledge, the current patient marks the first reported case of a Mycobacterium bovis mycotic aneurysm of a coronary artery following intravesicular BCG therapy for bladder cancer.

The exact mechanism for mycotic aneurysm formation is unknown. Potential proposed mechanisms for BCG infection include direct intimal colonization from hematogenous spread, metastatic implantation through the vasa-vasorum or local vascular extension from an adjacent infectious site [5, 22]. Accordingly, the most important known risk factor for BCG infection is disruption of the urogenital mucosa at the time of bladder instillation, including traumatic indwelling catheter insertion, recent biopsy, surgery or active cystitis [2, 22]. Some authors have proposed the use of prophylactic isoniazid at the time of BCG instillation, however this has not been shown to be effective [34]. Most authors now advise delaying BCG therapy for several weeks following cystitis, bladder resection, biopsy or difficult catheterizations [22].

**Clinical & Imaging Findings:**

The diagnosis of M. bovis mycotic aneurysms is difficult and remains elusive owing to the nonspecific clinical findings and relatively low incidence of occurrence. Diagnosis is often delayed, with the mean time of diagnosis occurring 23 months (range 4-69 months) following BCG therapy [5]. Patients usually present with fever of unknown origin, weight loss, a palpable abdominal mass and rarely signs/symptoms of aneurysm rupture. The gold standard for definitive diagnosis is bacterial cultures; however, this requires a minimum incubation period of 6-8 weeks and acid-fast stains require a minimum of 10,000 organisms per gram of tissue to yield positivity [5, 26].

The imaging findings are relatively straightforward but also nonspecific, demonstrating typical findings for mycotic or inflammatory aneurysms such as saccular aneurysms of medium or large-caliber arteries and associated perivascular inflammation [Tables 1, 2]. Catheter angiography remains the gold standard for imaging diagnosis of mycotic aneurysms, but its use is variable based on local expertise and limited by invasiveness. Multidetector CT angiography is the current imaging modality of choice for the evaluation of suspected mycotic aneurysms, with the advantages of rapid image acquisition, high sensitivity, widespread availability, high spatial resolution and multi-planar capabilities [35]. Mycotic aneurysms commonly appear as saccular dilatations of arteries not commonly involved by atherosclerosis (i.e. peripheral arteries), although the infrarenal abdominal aorta is still the most common reported site of mycotic aneurysms. Associated findings of perivascular inflammatory change and relative lack of atheromatous wall calcifications increases the specificity for mycotic aneurysm [Table 2]. MR angiography demonstrates similar findings as CTA but is limited by long acquisition times, availability, motion susceptibility and relatively lower spatial resolution [35]. As in our patient, ultrasound is mainly limited to mycotic aneurysms of the peripheral arteries and is not reliable at diagnosing visceral or abdominal aortic aneurysms.

**Treatment & Prognosis:**

The treatment of severe systemic BCG reactions includes a combination of medical and surgical/interventional therapy. The International Bladder Cancer Group recommends discontinuation of BCG therapy, initiation of high-dose fluoroquinolones, corticosteroids and daily antimycobacterial therapy with isoniazid, rifampin and ethambutol for six months duration [24]. It is well established that this particular strain of Mycobacterium is resistant to pyrazinamide [36]. In regards to treatment of the aneurysm itself, the mainstay of therapy in the reported literature has been surgical stent-grafting, or less commonly surgical bypass procedures. Of the 32 reported cases 68% (22/32) underwent a surgical graft procedure [Table 1]. As endovascular aneurysm repairs are becoming increasingly more popular for the treatment of non-mycotic aneurysms due to decreased peri-operative morbidity and mortality, some have suggested that endovascular repair should be the exclusive means for treating mycotic aneurysms [37]. The data on endovascular repair of mycotic M. bovis aneurysms, however, is scarce. To our knowledge, there have only been two reported cases of primary repair by endovascular stent-graft placement [12, 33]. One patient was treated with EVAR for a mycotic AAA and died due to a MI as a peri-operative complication [12]. The other patient was also treated with an EVAR but lost to follow-up (left the hospital against medical advice, 33). With regards to treatment of peripheral M. bovis mycotic aneurysms, all of the previous reported cases have undergone surgical treatment as the primary means of repair with generally good results [7, 8, 15, 16, 17, 19, 20, 23, 30]. To our knowledge, the current patient marks the first reported case of primary endovascular repair with covered stents for peripheral BCG aneurysms. Furthermore, this also marks the first known reported case of a BCG-induced coronary aneurysm. As such, no established guidelines for treatment exist and the prognosis remains unknown.

**Differential Diagnosis:**

The main differential diagnosis for mycotic aneurysms on imaging is relatively limited and includes atherosclerotic aneurysms, inflammatory aneurysms, contained aneurysm rupture and aortoenteric fistulas [Table 2]. Once mycotic aneurysms are suspected, it is very important to correlate findings with a detailed history and physical. Although rare,
any patient with a recent or prior history of BCG therapy for bladder cancer should yield a lower threshold for the consideration M. bovis mycotic aneurysm in these patients.

Atherosclerotic aneurysm: This is the most common etiology for aneurysm and typically occurs in males older than 50 years of age with atherosclerotic risk factors (i.e. hypertension, smoking, diabetes). The most common location is the infrarenal aorta and manifests on imaging as fusiform dilatation with calcified and fibrofatty wall plaque. Furthermore, if an aneurysm is encountered within a coronary artery, it is most likely to be atherosclerotic in etiology.

Inflammatory aneurysm: Inflammatory aneurysms typically occur in patients with similar risk factors to atherosclerotic aneurysms, but are generally found about a decade earlier and present with systemic inflammatory symptoms (i.e. fever, weight loss, elevated ESR). Similar to atherosclerotic aneurysms, inflammatory aneurysms present with fusiform dilatation of the infrarenal aorta; however, imaging will also demonstrate signs of inflammation (i.e. thickening aortic wall, peri-vascular stranding and/or fluid).

Contained aneurysm rupture: Patients with aneurysms of any etiology may occasionally present with a contained rupture. Clinically, patients will often present with acute onset abdominal or back pain with or without hemodynamic instability. The most reliable sign on imaging is the presence of focal wall discontinuity within a calcified aneurysm wall with associated peri-aneurysmal fluid. Conventional catheter angiography might demonstrate active arterial extravasation if the rupture is ongoing.

Aortoenteric fistula: Defined as an abnormal fistulous connection between small bowel and the aorta, this typically occurs in the setting of prior retroperitoneal surgery, most commonly surgical AAA repair. Patients may present acutely with a brisk lower gastrointestinal bleed. Imaging will demonstrate loss of the normal fat plane between small bowel and abdominal aorta, and if present, gas within the aorta is highly suggestive of aortoenteric fistula.

**TEACHING POINT**

Bacillus Calmette-Guerin (BCG) is a generally safe and accepted method for the treatment of superficial transitional cell carcinoma (TCC) of the bladder. In rare instances, BCG-activation may result in the serious complication of mycotic aneurysm formation, which typically manifests months-to-years following therapy as mycotic AAA, or less commonly mycotic peripheral aneurysms. Therapy consists of cessation of ongoing BCG therapy, antimycobacterial therapy and surgical/endovascular aneurysm repair.

**REFERENCES**


Figures

Figure 1: A 63-year-old male with acute left knee pain and swelling due to a ruptured left popliteal artery mycotic aneurysm. Findings: (a) - Sagittal gray-scale ultrasound (US) of the left popliteal fossa shows a large non-compressible acute hematoma (yellow asterisk). (b) - Transverse color Doppler US of the left medial knee shows a small amount of "ying-yang" flow within a partially thrombosed saccular aneurysm sac (white arrow). (c, d) - Transverse color Doppler US demonstrates an incidental partially thrombosed saccular aneurysm arising from the left common femoral vein (CFV; white asterisk).

Technique: GE Logiq E9 linear 8.4 Mhz transducer.
Cardiac Imaging: Left Anterior Descending Coronary Artery and Multiple Peripheral Mycotic Aneurysms Due to Mycobacterium Bovis Following Intravesical Bacillus Calmette-Guerin Therapy: A Case Report

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Figure 2: A 63 year-old-male with an acute ruptured left popliteal aneurysm.
Findings: (a) - Axial CTA demonstrating a mixed density acute left popliteal hematoma with focal active arterial extravasation (blue arrow). (b, c) - Axial and 3D volume rendered CTA of the lower pelvis demonstrating a left (white arrow) and bilateral (yellow arrow) saccular aneurysms arising from the distal common femoral arteries.
Technique: Siemens SOMATOM Definition Flash, 128 channel, mAs 10,283, kVP 100, slice thickness 3 mm; Contrast material: 125 mL Omnipaque 350 mg/ml injected at a rate of 5 ml/s.

Figure 3: A 63 year-old-male with acute onset right knee pain due to ruptured right popliteal artery mycotic aneurysm.
Findings: (a) - Sagittal grayscale (left) and color Doppler (right) US demonstrating an acute hematoma (asterisk) adjacent to a 5.7 x 2.7 cm irregular-shaped mycotic aneurysm arising from the popliteal artery (arrow). (b) - Sagittal spectral Doppler US demonstrating a "to-and-fro" flow pattern within the aneurysm sac, compatible with pseudoaneurysm (yellow calipers).
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Figure 4: A 63 year-old-male undergoing covered stent placement of a right popliteal mycotic aneurysm.
Findings: (a) - Carbon dioxide digital subtraction angiogram (DSA) demonstrating a saccular aneurysm arising from the above-knee popliteal artery (yellow arrow) with active extravasation (curved black arrow). (b) - A Storq guidewire was used to cross the aneurysm in the mid SFA. (c) - Placement of a 8 mm x 10 cm balloon-expandable Viabahn covered stent. (d) - Post DSA angiogram demonstrating successful exclusion of the aneurysm sac following placement of the covered stent.
Technique: Siemens AXIOM Artis, Fluoroscopy time 8.6 minutes, KAP 605 mGy2, kVP 66, Contrast material: Carbon dioxide and Visipaque, hand-injected.

Figure 5: A 63 year-old-male status post bilateral covered popliteal artery covered stent placement for ruptured mycotic aneurysms.
Findings: (a) - 3D-volume rendered and (b) oblique coronal maximum intensity projection CTA demonstrating patent bilateral covered popliteal stents with successful exclusion of the popliteal aneurysm sacs (arrows).
Technique: GE Lightspeed, 16 channel, mAs 1,066, kVP 120, slice thickness 2.5 mm; Contrast material: 125 mL Omnipaque 350 mg/ml injected at a rate of 5 ml/s.
Figure 6: A 63-year-old-male status undergoing surveillance CTA status post bilateral interposition femoral vein grafts and bilateral popliteal covered stent placement.
Findings: (a, b) - Axial CTA; (c, d) - 3D-volume rendered CTA; (e) - coronal CTA through the right common femoral artery demonstrating incidental saccular pseudoaneurysms arising from the proximal (white arrow) and distal (yellow arrow) attachment sites of the surgical interposition grafts.
Technique: GE Lightspeed, 16 channel, mAs 56,703, kVP 120, slice thickness 2.5 mm; Contrast material: 125 mL Omnipaque 350 mg/ml injected at a rate of 5 ml/s.
Cardiac Imaging: Left Anterior Descending Coronary Artery and Multiple Peripheral Mycotic Aneurysms Due to Mycobacterium Bovis Following Intravesical Bacillus Calmette-Guerin Therapy: A Case Report

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Figure 7: A 63-year-old male undergoing placement of a bifurcated aortoiliac stent graft and coil embolization of a distal right CFA mycotic pseudoaneurysm.
Findings: (a, b, d) - Coronal oblique DSA demonstrating saccular mycotic aneurysms arising from the (a, arrow) proximal and (b and d, arrow) distal attachment sites of a common femoral surgical interposition graft. (c) - Coronal DSA following successful placement of an Endologix AFX bifurcated aortoiliac stent graft. (e) - Coronal DSA showing a deployment of coils into the distal aneurysm sac. (f) - Post-coil DSA demonstrating successful exclusion of the distal mycotic aneurysm sac.
Technique: Siemens AXIOM Artis, Fluoroscopy time 37.8 minutes, KAP 6,036 mGy/m², kVP 66-73, Contrast material: 30 mL Omnipaque 300 mg/mL.
Figure 8: 63 year-old-male undergoing surveillance CTA following coil-embolization of a right CFA mycotic aneurysm. Findings: (a) - Curved planar CTA; (b) - Oblique coronal CTA; (c) - 3D-volume rendered CTA demonstrating patent right common femoral surgical interposition grafts (arrows) with successful coil exclusion of a mycotic aneurysm at the distal graft attachment site (asterisk). Technic: GE Lightspeed, 16 channel, mAs 28,497, kVP 120, slice thickness 2.5 mm; Contrast material: 125 mL Omnipaque 350 mg/ml injected at a rate of 5 ml/s.

Figure 9: A 63 year-old-male undergoing CTA of the lower extremities for increasing pain and swelling behind the right knee. Findings: (a, b) - Axial CTA demonstrating an enlarging 8.1 x 5.9 cm right popliteal hematoma (red arrow) surrounding a patent popliteal stent graft. The hematoma previously measured 7.4 x 5.2 cm three weeks earlier. (c, d) - Axial CTA of the lower chest demonstrates a 5.7 x 4.1 cm thrombosed aneurysm (cyan arrow) which appears to arise from the LAD (yellow arrow). In retrospect, this aneurysm was present on screening CTA of the chest performed 49 days earlier and measured 1.6 x 1.6 cm at that time (white arrow). Technic: GE Lightspeed, 16 channel, mAs 10,400, kVP 120, slice thickness 2.5 mm; Contrast material: 125 mL Omnipaque 350 mg/ml injected at a rate of 5 ml/s.
Figure 10: A 63 year-old-male undergoing coronary CTA for evaluation of a left coronary artery mycotic aneurysm. Findings: (a-d) - Axial pre-contrast (a), arterial-phase (b) and 3D-volume rendered (c, d) coronary CTA demonstrating a large saccular aneurysm with sluggish internal flow (blue arrow) arising from the mid left anterior descending (LAD) artery (yellow arrow). Significant mass effect and compression on the left ventricular outflow tract (LVOT) is noted (purple arrow). Technique: Siemens SOMATOM Definition Flash, 128 channel, mAs 1,683, kVP 120, slice thickness 0.75 mm; Contrast material: 97 mL Omnipaque 350 mg/ml injected at a rate of 5 ml/s.
Figure 11: A 63 year-old-male undergoing left main coronary catheter angiography.
Findings: (a-d) - Dynamic images obtained during injection of the left main coronary artery (white arrow) demonstrates a focal caliber change in the mid LAD (blue arrow). An amorphous collection of contrast is noted adjacent to the mid LAD, which progressively fills a large saccular aneurysm sac arising from the mid LAD (curved red arrow). There is no opacification of the LAD distal to the aneurysm site, indicating occlusion of the distal LAD. The left circumflex artery appears normal (curved yellow arrow).
Technique: Fluoroscopy time 6.2 minutes; DAP 49 mGy-m2; Contrast material: 70 mL Omnipaque 350 mg/ml.
Table 1: Literature review. Reported cases of BCG-induced M. bovis aneurysms

<table>
<thead>
<tr>
<th>First author</th>
<th>Age</th>
<th>Gender</th>
<th>Location of aneurysm(s)</th>
<th>Antibiotics</th>
<th>Intervention</th>
<th>Outcome</th>
</tr>
</thead>
<tbody>
<tr>
<td>Woods⁶</td>
<td>62</td>
<td>F</td>
<td>AAA</td>
<td>+</td>
<td>Surgical graft</td>
<td>Healthy</td>
</tr>
<tr>
<td>Bornet⁷</td>
<td>74</td>
<td>M</td>
<td>Bilateral femoral</td>
<td>+</td>
<td>Surgical graft</td>
<td>Healthy</td>
</tr>
<tr>
<td>Deresiewicz⁸</td>
<td>67</td>
<td>M</td>
<td>AAA, iliac</td>
<td>-</td>
<td>Surgical graft</td>
<td>Died (disseminated infection)</td>
</tr>
<tr>
<td>Izes⁹</td>
<td>69</td>
<td>M</td>
<td>Aortic arch</td>
<td>-</td>
<td>None</td>
<td>Died (palliative care, untreated)</td>
</tr>
<tr>
<td>Wolf¹⁰</td>
<td>80</td>
<td>M</td>
<td>AAA</td>
<td>-</td>
<td>Surgical graft</td>
<td>Alive, developed aortoenteric fistula</td>
</tr>
<tr>
<td>Hellinger¹¹</td>
<td>71</td>
<td>M</td>
<td>AAA</td>
<td>+</td>
<td>Surgical graft</td>
<td>Healthy</td>
</tr>
<tr>
<td>Rozenblit¹²</td>
<td>76</td>
<td>M</td>
<td>AAA</td>
<td>+</td>
<td>Stent graft</td>
<td>Died (myocardial infarction)</td>
</tr>
<tr>
<td>Damm¹³</td>
<td>71</td>
<td>M</td>
<td>AAA</td>
<td>+</td>
<td>Surgical graft</td>
<td>Healthy</td>
</tr>
<tr>
<td>LaBerge¹⁴</td>
<td>75</td>
<td>M</td>
<td>AAA</td>
<td>NS</td>
<td>Surgical graft</td>
<td>NS</td>
</tr>
<tr>
<td>Seelig¹⁵</td>
<td>58</td>
<td>M</td>
<td>AAA</td>
<td>+</td>
<td>Surgical graft</td>
<td>Healthy</td>
</tr>
<tr>
<td>Seelig¹⁵</td>
<td>71</td>
<td>M</td>
<td>AAA, popliteal</td>
<td>+</td>
<td>Surgical graft</td>
<td>Healthy</td>
</tr>
<tr>
<td>Farber¹⁶</td>
<td>74</td>
<td>M</td>
<td>Femoral</td>
<td>+</td>
<td>Prosthesis removal</td>
<td>Healthy</td>
</tr>
<tr>
<td>Geldmacher¹⁷</td>
<td>68</td>
<td>M</td>
<td>Carotid</td>
<td>+</td>
<td>Surgical graft</td>
<td>NS</td>
</tr>
<tr>
<td>Kamphuis¹⁸</td>
<td>65</td>
<td>M</td>
<td>AAA</td>
<td>-</td>
<td>Surgical graft</td>
<td>Died (post-operative shock)</td>
</tr>
<tr>
<td>Wada¹⁹</td>
<td>75</td>
<td>M</td>
<td>AAA, femoral</td>
<td>-</td>
<td>Surgical graft</td>
<td>Healthy</td>
</tr>
<tr>
<td>Witjes²⁰</td>
<td>67</td>
<td>M</td>
<td>Popliteal</td>
<td>+</td>
<td>Surgical graft</td>
<td>Healthy</td>
</tr>
<tr>
<td>Dahl²¹</td>
<td>69</td>
<td>M</td>
<td>AAA</td>
<td>+</td>
<td>Surgical graft</td>
<td>Healthy</td>
</tr>
<tr>
<td>Harding²²</td>
<td>80</td>
<td>M</td>
<td>AAA</td>
<td>+</td>
<td>Surgical graft</td>
<td>Healthy</td>
</tr>
<tr>
<td>Safdar²</td>
<td>79</td>
<td>M</td>
<td>AAA</td>
<td>+</td>
<td>Surgical graft</td>
<td>Healthy</td>
</tr>
<tr>
<td>Costiniuk²³</td>
<td>75</td>
<td>M</td>
<td>AAA, femoral</td>
<td>+</td>
<td>Surgical graft</td>
<td>Healthy</td>
</tr>
<tr>
<td>Coscas⁵</td>
<td>79</td>
<td>M</td>
<td>Thoracic aorta, Iliac, PT, carotids</td>
<td>+</td>
<td>None</td>
<td>Died (cancer progression)</td>
</tr>
<tr>
<td>Maundrell²⁴</td>
<td>75</td>
<td>M</td>
<td>AAA (juxtarenal)</td>
<td>+</td>
<td>Surgical (Ax-bifem bypass)</td>
<td>Died (aneurysm rupture)</td>
</tr>
<tr>
<td>Psoinos²⁵</td>
<td>69</td>
<td>M</td>
<td>AAA</td>
<td>+</td>
<td>Surgical graft</td>
<td>Healthy</td>
</tr>
<tr>
<td>Roylance²⁶</td>
<td>77</td>
<td>M</td>
<td>AAA (aortoenteric fistula)</td>
<td>-</td>
<td>Surgical graft</td>
<td>Died (post-operative AKI, respiratory failure)</td>
</tr>
<tr>
<td>Samadian²⁷</td>
<td>94</td>
<td>M</td>
<td>AAA (osteodiscitis)</td>
<td>+</td>
<td>NS</td>
<td>NS</td>
</tr>
<tr>
<td>Santbergen²⁸</td>
<td>58</td>
<td>M</td>
<td>AAA (infected EVAR stent graft)</td>
<td>+</td>
<td>Surgical graft removal</td>
<td>Healthy</td>
</tr>
<tr>
<td>Mizoguchi²⁹</td>
<td>81</td>
<td>M</td>
<td>AAA (infected EVAR stent graft)</td>
<td>+</td>
<td>Surgical graft removal, surgical bypass</td>
<td>NS</td>
</tr>
<tr>
<td>Kuy³⁰</td>
<td>73</td>
<td>M</td>
<td>Carotid</td>
<td>NS</td>
<td>Surgical graft</td>
<td>NS</td>
</tr>
<tr>
<td>Davis³¹</td>
<td>64</td>
<td>M</td>
<td>Thoracoabdominal aorta (multi)</td>
<td>+</td>
<td>Surgical graft</td>
<td>Healthy</td>
</tr>
<tr>
<td>Seastedt³²</td>
<td>70</td>
<td>M</td>
<td>Thoracic aorta</td>
<td>+</td>
<td>Surgical graft</td>
<td>NS</td>
</tr>
<tr>
<td>Floros³³</td>
<td>57</td>
<td>M</td>
<td>AAA</td>
<td>+</td>
<td>Endovascular stent graft</td>
<td>NS (patient left AMA)</td>
</tr>
<tr>
<td>Present case</td>
<td>63</td>
<td>M</td>
<td>Popliteal (bl), CFA, LAD</td>
<td>+</td>
<td>Stent grafts (popliteal, CFA)</td>
<td>ABX, Palliative care</td>
</tr>
</tbody>
</table>

Total # cases: 32

AAA = Abdominal aortic aneurysm; NS = not specified; EVAR = Endovascular aneurysm repair; PT = posterior tibial; CFA = common femoral artery; LAD = left anterior descending
**Cardiac Imaging:** Left Anterior Descending Coronary Artery and Multiple Peripheral Mycotic Aneurysms Due to Mycobacterium Bovis Following Intravesical Bacillus Calmette-Guerin Therapy: A Case Report

---

<table>
<thead>
<tr>
<th>Clinical history</th>
<th>Mycobacterium bovis mycotic aneurysm</th>
<th>Inflammatory aneurysm</th>
<th>Atherosclerotic aneurysm</th>
<th>Contained aneurysm rupture</th>
<th>Aortoenteric fistula</th>
</tr>
</thead>
<tbody>
<tr>
<td>History of BCG therapy for bladder cancer</td>
<td>• History of BCG therapy for bladder cancer</td>
<td>• Similar risk factors as for atherosclerotic aneurysms</td>
<td>• Elderly (M &gt; F)</td>
<td>• Same as for atherosclerotic aneurysms</td>
<td>• Presents with brisk GI bleed</td>
</tr>
<tr>
<td>Mean age 71 M &gt;&gt; F</td>
<td>• Mean age 71 M &gt;&gt; F</td>
<td>• Typically younger (40-50s)</td>
<td>• Atherosclerotic risk factors</td>
<td>• Acute onset back/abdominal pain</td>
<td>• Risk factors: surgical AAA repair, infectious aortitis, PAU, tumor invasion, XRT</td>
</tr>
<tr>
<td>Weight loss/fever</td>
<td>• Weight loss/fever</td>
<td>• Back pain, fever, weight loss</td>
<td>• Acute - hypotension</td>
<td>• Similar risk factors as for atherosclerotic aneurysms</td>
<td></td>
</tr>
<tr>
<td>+ Cultures for <em>M. bovis</em></td>
<td>• + Cultures for <em>M. bovis</em></td>
<td>• Elevated ESR</td>
<td>• Typically younger (40-50s)</td>
<td>• Typically younger (40-50s)</td>
<td></td>
</tr>
</tbody>
</table>

**X-ray**
- Typically normal
- Typically normal
- Enlarged aorta (with calcified walls)
- Typically normal
- Typically normal

**US**
- Rapidly enlarging, focal saccular
- "yin-yang" color flow
- Perivascular soft tissue
- Fusiform dilatation of the infrarenal aorta and iliacs
- May see thickened aortic wall

**CTA**
- Rapidly enlarging focal saccular aneurysm
- Perivascular soft tissue inflammation (stranding, edema, fluid)
- Thoracoabdominal aorta > peripheral vascular system > carotids > coronaries
- Fusiform infrarenal AAA and iliac aneurysms
- Thickened aortic wall
- May see findings of retroperitoneal fibrosis

**MRI**
- Rapidly enlarging, focal saccular aneurysm
- Perivascular edema (low T1, high T2)
- May see adjacent inflammation or abscess
- Fusiform infrarenal AAA and iliac aneurysms
- Thickened aortic wall
- Perianeurysmal soft tissue (low T1, high T2, enhancing)

**Angiography**
- Focal, saccular aneurysm
- May see active extravasation if acutely ruptured
- Fusiform infrarenal AAA and iliac aneurysms
- Fusiform arterial dilatation
- Active arterial extravasation
- Active arterial extravasation into bowel

**Table 2:** Differential table for Mycobacterium bovis mycotic aneurysms
Etiology | Localized or disseminated infection by live BCG. Exact mechanism for mycotic aneurysm formation unknown.
---|---
Incidence | Very rare (<< 1%); 32 total reported cases
Gender ratio | M >> F (31 M, 1 F in reported literature)
Age predilection | Middle-aged/elderly; Mean age 71 in reported literature
Location | Aorta (infrarenal > suprarenal) > iliac, femoral > popliteal > carotid > coronary
Risk factors | Urogenital trauma (i.e. difficult bladder catheterizations, cystitis, recent biopsy or surgery), immunocompromised
Clinical findings | Fever of unknown origin, unintentional weight loss, abdominal/back pain, abdominal mass, ruptured aneurysm
Treatment | Long course (6-12 months) anti-mycobacterial antibiotics; Surgical or endovascular repair of aneurysms
Prognosis | Variable (highly dependent on location of aneurysm, clinical status of patient, co-morbidities)
Imaging findings | Rapidly enlarging saccular aneurysms in medium-large sized arteries, possible rupture

Table 3: Summary of BCG-induced Mycobacterium bovis mycotic aneurysms

| ABBREVIATIONS | 
|---|---|
| AAA: Abdominal aortic aneurysm | BCG: Bacillus Calmette-Guerin |
| CFA: Common femoral artery | CTA: Computed tomographic angiography |
| DNR: Do not resuscitate | ED: Emergency department |
| EVAR: Endovascular aneurysm repair | LAD: Left anterior descending |
| TCC: Transitional cell carcinoma | TURBT: Transurethral resection of bladder tumor |
| US: Ultrasound | 

| KEYWORDS | 
|---|---|
| Mycotic aneurysm; BCG; Bacillus Calmette-Guerin; Mycobacterium bovis; intravesicular | 

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