Recurrent scrotal Arteriovenous Malformation as a Slowly Increasing Left Testicular Swelling

A case report

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Abstract

Arteriovenous malformations (AVMs) are benign vascular lesions. Although, the majority of AVMs occur in the central nervous system, there are published reports of AVMs involving all systems including the scrotum, kidney, and uterus. Herein we report a case of 37 years old male presented with recurrent gradual scrotal swelling for 4 years attributed to scrotal AVM. Embolization was done but one year later his symptoms reoccurred. As a result, left partial scrotal wall excision was carried out without complications.

Keywords: arteriovenous malformation, AVM, scrotal swelling, scrotal malformation.
Introduction
Arteriovenous malformations (AVMs) are benign vascular lesions. They are described as abnormal vessels fed by arteries and drained by veins without intervening capillaries.\textsuperscript{1,2} Although the majority of AVMs occur in the central nervous system, there are published reports of AVMs involving the scrotum, kidney, and uterus.\textsuperscript{2,3,4} Few cases of scrotal AVMs have been described in the literature. Based on the published reports, the clinical presentation of scrotal AVMs is highly variable, ranging from an incidental finding on imaging for infertility to a bleeding mass.\textsuperscript{1,5} Since scrotal AVMs have variable presentations and is rarely described in the literature, we are reporting a case of a 37-year-old male presented with a slowly increasing left testicular swelling attributed to scrotal AVM.

Case Report
37 years old male smoker presented to the urology clinic with a gradual scrotal swelling that started four years ago. He complained of on and off scrotal pain, occasional feeling of scrotal warmth, and scrotal discomfort. The patient was diagnosed in another hospital with a testicular artery aneurysm and left testicular varicocele. The patient denied any history of trauma, urinary tract infection, voiding symptoms, previous surgeries, and his past medical history was unremarkable. Upon physical examination, the testes were intra-scrotal. There were no signs of inflammation, and the cremasteric reflex was intact bilaterally. Both epididymides were palpable and non-tender. However, pampiniform plexus at the neck of the scrotum was very pulsatile (figure A). Moreover, multiple skin varices over the left scrotum were seen. Urinalysis was normal and urine culture was negative. Routine laboratory tests were unremarkable. Abdominal and pelvis computed tomography (CT) showed left scrotal arteriovenous malformation with enlarged small and medium-sized serpiginous structures with a feeder artery arising from the proximal superficial artery. Two months later, the patient was referred to interventional radiology for embolization, which was successfully done utilizing Onyx 18\% (figure B). After one year, on follow-up, the pampiniform plexus were pulsatile again which necessitated a CT angiogram. CT angiogram confirmed the recurrence of arteriovenous malformation. The patient was counselled about the available treatment options and given time to decide. Due to the risk of recurrence as well as the
possibility of technical failure with embolization, he decided to go with the surgical
treatment. The patient was booked for surgery, and partial scrotal wall excision was
done through an elliptical incision (figure C). Three arteries that feed into the
arteriovenous malformation were identified and controlled with vicryl ties. The
malformation and the skin that covering it were removed and sent to the pathology
lab. Dartos muscle was closed in a multi-fashion layer. The skin was closed by vicryl
rapide 4-0 in a vertical mattress. The histopathological study confirmed the diagnosis
by detecting vascular structures extending from fibrofatty tissues measuring 8x0.5 cm
grossly, and prominent subcutaneous large congested vascular spaces
microscopically. The patient was discharged one day after the surgery with no
complications. Two months postoperatively, the patient was doing fine with no active
complaint, and the wound healed properly.

The consent was obtained orally as the images were taken from the patient in the
clinic. We explained to him the importance of reporting and publishing his case for
educational purposes, and he agreed.

Discussion
AVMs are malformations in the circulatory system characterized by arteries and veins
that are not connected by capillaries leading to various degrees of ischemia and
pain.\textsuperscript{1,2} Even though Central nervous system cases represent the majority of AVMs,
there are published reports of AVMs involving the kidney, uterus, and scrotum.\textsuperscript{2,3,4}
AVMs are rarely present in the urinary tract.\textsuperscript{6} We reviewed four previously published
scrotal AVM cases (Table 1). All revealed ages ranging from 19 to 31 years while our
patient was 37 years old. Scrotal AVM embraces wide-ranging presentations
including infertility, acute recurrent pain in the hemiscrotum, pain and swelling on the
testicle, and progressive diffused swelling in the scrotum with flashing skin and local
warmth. Our case presented with gradually increasing left testicular swelling with on
and off scrotal pain, and occasional feeling of scrotal warmth and discomfort. Of the
four cases we have reviewed, three denied any history of trauma, and one had a
positive trauma history which was a severe pelvic fracture, and the patient indicated
that there is difficulty in maintaining erection since. Our patient denied any trauma
history. Varicocele was found in two of the cases and was seen by sonography
whereas our patient had multiple skin varices above the left scrotum that was seen
during physical examination. Each one of the four cases we reviewed diagnosed
crotal AVM with a different modality. Some were challenging and required
orchietomy for a diagnosis while others were simple and detected by pelvic
angiography. Our patient was diagnosed by abdominal and pelvic CT. Two studies
were able to find and embolize the feeding arteries. Our patient underwent
embolization but had recurrence one-year later. Similar to our case, surgical
intervention was eventually done in all four cases, and it varied from left scrotal AVM
excision, orchiectomy, ileo-femoral bypass, and resection of the whole left side of the
scrotum. In our case, partial scrotal wall excision was done. After the surgical
intervention, all patients were symptom-free.\textsuperscript{5,6,7,8} We believe that the difference in
the presentation could be attributed to the location of the AVM, onset, duration, and if
there is a history of trauma. A possible explanation for the differences in imaging
modalities used to diagnose scrotal AVMs is the availability of imaging techniques in
the hospitals that encountered those cases. The decision of surgical intervention is
mainly based on the symptoms and how symptoms negatively affect the patient’s
quality of life.

Conclusion
Our case calls attention to a rare and challenging diagnosis that is scrotal AVM.
Recurrent scrotal pain, swelling, and warmth together with varicocele should raise
suspicion for scrotal AVM. Treatment varies depending on the symptoms present. We
believe that embolization of the feeding arteries is a possible option to start with, and
surgery should be preserved for recurrent cases.

Authors’ Contribution
FMA, MSA, and SIA were responsible for conceiving the idea, literature search, data
acquisition, and manuscript writing and revision. AA, SA, YA, and AB were
primarily involved in the management of the case and critically reviewed the final
version of the manuscript. All the authors have read and approved the final version of
this manuscript.
References


**Figure 1:** a picture of the left scrotal swelling with clear multiple skin varices.

**Figure 2:** the left image (A) showing a feeder artery supplying scrotal AVM. the right image (B) angiogram following Onyx embolization through the AVM is almost occluded.

**Figure 3:** partial left scrotal wall with AVM excision.
<table>
<thead>
<tr>
<th>Author</th>
<th>Age</th>
<th>Trauma History</th>
<th>Presentation</th>
<th>Semen analysis</th>
<th>Varicocele</th>
<th>Diagnosis method</th>
<th>Embolization</th>
<th>Treatment</th>
<th>Follow-up</th>
</tr>
</thead>
<tbody>
<tr>
<td>Monoski et al.</td>
<td>31</td>
<td>No history of trauma</td>
<td>Infertility</td>
<td>Severe oligospermia</td>
<td>A left varicocele</td>
<td>Pelvic angiography</td>
<td>Performed</td>
<td>Surgical left scrotal AVM excision</td>
<td>3 years post-surgery, successful spontaneous pregnancy</td>
</tr>
<tr>
<td>Agrawal et al.</td>
<td>25</td>
<td>Positive - Severe pelvic fracture 4 years ago</td>
<td>Pain associated with a soft swelling on his right testicle</td>
<td>Not Performed</td>
<td>No evidence of varicocele</td>
<td>Histopathological examination</td>
<td>Not performed</td>
<td>Ileo-femoral bypass surgery</td>
<td>Not mentioned</td>
</tr>
<tr>
<td>Sountoulides et al.</td>
<td>22</td>
<td>No history of trauma</td>
<td>Acute recurrent pain in the right hemiscrotum</td>
<td>Not Performed</td>
<td>No evidence of varicocele</td>
<td>Post-orchiectomy specimen</td>
<td>Not performed</td>
<td>Orchiectomy</td>
<td>2 years post-surgery, there was no complain</td>
</tr>
<tr>
<td>Mohammed et al.</td>
<td>19</td>
<td>No history of trauma</td>
<td>Progressive diffused swelling in the scrotum with flashing skin and local warmth</td>
<td>Not Performed</td>
<td>Varicocele with 1 cm dilated veins</td>
<td>CT arteriography</td>
<td>Performed</td>
<td>The whole left side of the scrotum was removed, and the left testicle was fixed to the right side</td>
<td>12 months post-surgery, there was no complain</td>
</tr>
<tr>
<td>Current case</td>
<td>37</td>
<td>No history of trauma</td>
<td>Gradually increasing scrotal mass with on and off scrotal pain and discomfort</td>
<td>Not performed</td>
<td>Multiple skin varices over the left scrotum were seen</td>
<td>Abdominal and pelvis CT</td>
<td>Performed</td>
<td>Partial scrotal wall excision</td>
<td>Two months post-surgery, there was no complain</td>
</tr>
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