A ureteral mass presenting as deep vein thrombosis: a very rare presentation

Abstract

Venous Thromboembolism is a dreaded condition. Many conditions and systemic diseases are known to predispose to its occurrence including malignancy, pancreatitis, burns, clotting disorders and direct compression. All of these have been known to affect one or more factors in Virchows Triad proposed roughly a century ago. It consists of Hypercoagulability, Vascular Endothelial Dysfunction, Stasis. Stasis, one of the factors in Virchows triad has been studied to a very small extent and it is this feature that is mostly responsible for the development of Deep Vein Thrombosis (DVT) due to direct compression. Iliofemoral thrombosis in malignancy patients can be caused due to vein compression by pelvic malignancy and usually presents as unilateral lower limb swelling. In patients with sudden onset of unilateral lower limb swelling without any perceived or diagnosed medical condition, detailed evaluation may lead to early identification, appropriate management and possibly cure.

We present a case of acute unilateral ilio-femoral DVT caused by external compression by a mid-ureteric mass.

Keywords: deep vein thrombosis, iliofemoral thrombosis, ureteric malignancy, upper tract urothelial cancers

Case presentation

A 45 year old post-menopausal female presented to her primary physician with complaints of left foot swelling which gradually progressed to involve the entire left lower limb. She did not give any antecedent history of trauma, ulceration on lower limb, haematuria, lower urinary tract symptoms, bowel complaints, vaginal bleeding, loss of weight, loss of appetite, previous history of DVT. There was no history of previous surgeries and co-existing morbidities. Her left lower limb was reddish in color, tense and tender to touch. Per vaginial examination revealed induration with in the left fornix. Rest of the general and systemic examination, were within normal limits. A venous Doppler revealed a partial thrombus of the Left Proximal external iliac vein and Left Common Iliac vein. Thrombophillic screen (PT-INR, aPTT, Protein C, Protein S, D, and Factor V Leiden Mutation) was within normal limits. Patient was started on Warfarin and put on total bed rest. Due to absence of any predisposing factors, further evaluation was considered and an ultrasound of the abdomen was done. It revealed a grossly dilated left renal pelvis with thinned out parenchyma & a 1.3*1.1cm lesion near the left vesico-ureteric junction. Contrast-Enhanced CT of Abdomen & Pelvis Figure 1 was done which shows poorly functioning left kidney with gross hydronephrosis & thinned out parenchyma, large 6.6*4.5cm heterogeneously enhancing solid cystic mass in the left ureter at the level of the iliac vessel crossing and 3.5*2.1cm heterogeneously enhancing solid cystic lesion in left distal ureter extending up to VUJ. Her hemoglobin, serum creatinine, liver function tests, urinalysis were within normal limits. Urine for malignant cytology was negative. Cystoscopy showed a bulge just lateral to the left ureteric orifice and no other visible mass. The left ureteric orifice could not be cannulated. Considering the diagnosis of a ureteric mass, a Left Nephroureterectomy with bladder cuff excision was planned. Warfarin was stopped 5 days before surgery and patient was put on bridging therapy with low molecular weight heparin. A cardiology consult was taken and it was decided to insert an IVC filter 24 hours before surgery as prophylaxis to decrease risk of intra-operative and post-operative pulmonary embolism. Intra-operatively DVT pump was not applied. A Radical Nephroureterectomy along with Hysterectomy and Left Salpingo-oophorectomy (Figure 2) (Figure 3) was done with gynecological assistance in view of local invasion of cervix. Mid ureteric mass was dissected off the iliac vessels with CTVS assistance and final specimen consisted of the Left kidney, ureter, Uterus, Left Ovary along with Lymph Node dissection. Total intra-operative blood loss was 600ml. Post-operatively patient was kept in the ICU before shifting to the ward. Warfarin was re-started on day 3. Patient had a relatively uneventful post-operative period.

Abbreviations: DVT, Deep Vein Thrombosis; VTE, Venous Thromboembolism

Figure 1 Pre-op CECT Abdomen + Pelvis( Upper half – Coronal Section, Lower half- Sagittal section.)
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Discussion

Upper tract urothelial carcinomas are a rare entity comprising only 5−10% of all urothelial tumors but recent literature reveals an increasing incidence.5 Ureteral tumors occur more commonly in the lower than in the upper ureter. Overall, about 70% of ureteral tumors occur in the distal ureter, 25% in the mid-ureter, and 5% in the proximal ureter.5,14 The most common clinical feature of urothelial carcinoma of the ureter is haematuria, either gross or microscopic, present in roughly 56−98% cases,5,12 followed by flank pain in 30% cases. Urothelial carcinoma of ureter presenting as unilateral lower limb swelling secondary to DVT is very rare. Venous thromboembolism (VTE) risk factors have been classified into- Modifiable, Non- modifiable and Temporary.8 Modifiable risk factors include obesity, non modifiable risk factors include genetic and familial disorders resulting in a hypercoagulable state. Temporary risk factors include hospitalization, malignancy, long distance travel, prolonged immobilization, hormonal therapy for various diseases. But most of the VTE is still labeled as ‘Idiopathic’ without any known antecedent cause.

Malignant compression of iliofemoral veins in the pelvis by malignancies is one such case in which patients may be mistakenly labeled as having idiopathic VTE with potentially disastrous outcomes. Ureteric masses causing direct iliofemoral venous compression with resultant DVT and unilateral lower limb swelling have seldom been reported. Most of the reported cases of iliofemoral venous compression with resultant DVT have been due to compression by enlarged lymph nodes secondary to Urothelial or gynecological malignancies. Liao et al.,9 analyzed 893 cancer patients over a 10 year period. The inclusion criteria for their study were:

1. presence of unilateral lower limb swelling,
2. a CT showing malignancy causing external compression on iliac or femoral vein and
3. a duplex ultrasound revealing vein thrombosis or venous flow insufficiency over a femoral vein or saphenous vein.

Patients with bilateral lower limb edema suggestive of IVC involvement were excluded. 63 (8%) of these patients were diagnosed with iliofemoral venous obstruction. 21 of these 63 patients developed iliofemoral venous thrombosis. Most of the compression was by lymph nodal metastasis secondary to urothelial cancers (46). Bladder was the most common primary site (29) followed by ureter (n = 8), renal pelvis (n = 5), prostate (n = 2), or penis (n = 2) but none of these were due to direct malignant compression by Ureteric Mass. Other causes of iliofemoral venous compression included the most commonly described syndrome called the Iliac Vein Compression Syndrome or the May-Thurner Syndrome involving compression of Left Iliac Vein by Right Iliac Artery.10

Many other rare causes of iliofemoral compression and thrombosis have been reported in literature including uterine fibroids,12 psoas abscess,12 retroperitoneal fibrosis,15 penile prosthesis reservoir,14 synovial cyst of hip.15 Other urological entities have also been reported to cause iliofemoral venous thrombosis. Alliota et al.,16 reported a giant hydrenephrosis as a cause of iliofemoral thrombosis. Bladder diverticulae have also been known to cause iliofemoral thrombosis according to various reports.17−19 Through our case report, we would like to emphasize the need for thorough evaluation of patients presenting with unilateral lower limb swelling due to DVT without any predisposing factors. A simple ultrasound of the abdomen or contrast enhanced CT might shed light on a sinister underlying malignancy. Proper diagnosis will guide therapy and will lead to early management of malignancy resulting in better prognosis.

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Conflict of interest

Author declares there is no conflict of interest in publishing the article.

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