Case Report

Bladder cancer invasion along a tension-free vaginal mesh

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Abbreviations & Acronyms BCG = Bacillus Calmette– Guérin CIS = carcinoma *in situ* CT = computed tomography LRC = laparoscopic radical cystectomy LSC = laparoscopic sacrocolpopexy MRI = magnetic resonance imaging POP = pelvic organ prolapse TVM = tension-free vaginal mesh

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Received 28 September 2020; accepted 25 December 2020. Online publication 21 January 2021 **Introduction:** The effect of synthetic mesh after pelvic organ prolapse surgery on the progression of bladder cancer remains unclear.

Case presentation: A 79-year-old woman who underwent a tension-free vaginal mesh procedure 8 years prior was diagnosed with carcinoma *in situ* of the bladder. Although intravesical Bacillus Calmette–Guérin therapy was started, the tumor rapidly became muscle invasive. Laparoscopic radical cystectomy was performed following radiochemotherapy; however, the tumor extended to the left internal obturator muscle along the mesh arm. Pathological findings showed desmoplastic high-grade urothelial carcinoma infiltrating around the mesh. Finally, cancer recurred rapidly in the left internal obturator muscle.

Conclusion: Synthetic mesh can become an abnormal anatomical pathway for tumor infiltration. Therefore, in high-risk bladder cancer patients who underwent a tension-free vaginal mesh procedure, radical cystectomy should be performed without delay before the tumor invades the perivesical tissue.

Key words: bladder cancer, carcinoma *in situ*, laparoscopic radical cystectomy, pelvic organ prolapse, tension-free vaginal mesh.

Keynote message

We report a rare case of bladder cancer invading the pelvic wall muscle along the TVM arm. LRC was performed, and pathological findings showed desmoplastic high-grade urothelial carcinoma infiltrating around the mesh. Synthetic mesh can become an abnormal anatomical pathway for tumor infiltration.

Introduction

The TVM procedure is commonly used for POP.^{1,2} A synthetic mesh is placed on the anterior and/or posterior aspect of the vagina, suspending the pelvic organ via a transvaginal procedure.³ Although LSC has recently become the gold standard,^{4,5} the TVM procedure remains a valid technique in cases contraindicated for LSC or in recurrent cases of native tissue repair for cystocele.

CIS of the bladder is a flat non-muscle invasive urothelial carcinoma usually treated with intravesical BCG therapy; however, it is a poorly differentiated tumor and can progress aggressively.⁶ CIS of the bladder progress to muscle-invasive cancer in approximately 54% of patients without any treatment and approximately 10–66% if intravesical treatment with BCG or chemotherapy is performed.⁷

The effect of synthetic mesh after surgery for POP on bladder cancer progression has not been described. We report a rare case of muscle-invasive bladder cancer that progressed from CIS and invaded the pelvic wall along the mesh arm of the TVM procedure.

Case presentation

A 79-year-old woman, complaining of pain on urination, was referred to our department. She had a surgical history of left nephrectomy for renal tuberculosis 64 years earlier and the anterior and posterior TVM procedure for POP 8 years earlier. Cystoscopy showed diffuse



Fig. 1 MRI findings of bladder cancer after radiochemotherapy. (a) Axial view: bladder cancer extends to the perivesical tissue (arrow). (b) Coronal view: bladder cancer attaches to the left internal obturator muscle (arrow-head). B, bladder; O, internal obturator muscle; U, uterus.

erythematous bladder mucosa, and urinary cytology was positive for cancerous cells. MRI and CT findings showed neither bladder tumors nor metastatic lesions. Cold-cup biopsies from the bladder mucosa were taken; the pathological analysis indicated a high-grade, flat, non-muscle invasive urothelial carcinoma. CIS of the bladder was diagnosed, and weekly intravesical BCG therapy was administered. After five injections, intravesical BCG therapy was discontinued because of severe pain. Follow-up MRI showed a bladder tumor (4.5 cm in diameter) in the trigone area, which had infiltrated the perivesical tissue. Upon vaginal examination, a hard mass was palpable at the anterior vaginal wall. Right ureterostomy was established before the right ureteral orifice obstruction. Despite radiochemotherapy (nedaplatin 30 mg/m² weekly and external beam radiotherapy 39.6 Gy/22 Fr), the tumor only decreased from 4.5 cm to 3.5 cm in size on MRI (Fig. 1a,b). Chronic cystitis persisted from admission and antibacterial treatment was necessary on occasion; however, the degree of inflammation by blood test recovered to an almost normal level.

LRC was performed after a month of radiochemotherapy because of persistent severe bladder pain. In the right



Fig. 2 Intraoperative view of LRC. (a) The right side of the retropubic space. The anterior mesh arm (arrowhead) penetrating the right internal obturator muscle is observed. (b) The left side of the retropubic space. Extensive and strong adhesion between the bladder wall and internal obturator muscle was indicated around the left anterior mesh arm (arrowhead). (c) The anterior vaginal fornix. Sutures that fixed the anterior mesh to the uterine cervix were observed (arrow). B, bladder; O, internal obturator muscle.

retropubic space, we identified the right anterior mesh arm of the TVM procedure penetrating the right internal obturator muscle (Fig. 2a). There was an extensive and strong adhesion between the bladder wall and left internal obturator muscle around the left anterior mesh arm (Fig. 2b). Although intraoperative rapid pathological diagnosis did not be performed, the left internal obturator muscle around the mesh arm was widely resected to excise the tumor. At the anterior vaginal fornix, sutures fixing the anterior mesh to the uterine cervix were visualized (Fig. 2c). From the



Fig. 3 Findings of pathology. (a) Cut surface of the resected urinary bladder. Elastic firm tumor surrounding the blue mesh (arrow). Scale bar: 10 mm. (b, c) Histopathology of the urinary bladder tumor (hematoxylin-eosin stain). (b) Bird's eye view shows carcinoma cells (arrows) infiltrating the polypropylene mesh (arrowheads). Scale bar: 100 μm. (c) High magnification of the red square section in (b), showing high-grade urothelial carcinoma cells (arrow). Scale bar: 100 μm.

anterior vaginal fornix, the anterior vaginal wall was incised, and the bladder was removed. The uterus was resected at the cervix, and the anterior vaginal wall defect was closed. The posterior mesh was preserved without exposure. Bilateral obturator lymph node dissection was also performed. The operation time was 9 h 59 min,



Fig. 4 Axial view of CT image at 6 weeks postoperatively. Tumor has recurred in the left internal obturator muscle (arrow).

estimated blood loss was 100 mL, and no perioperative complications occurred. The cut surface of the resected specimen showed a firm fibrotic tumor measuring $5 \text{ cm} \times 3 \text{ cm}$, which contained the mesh (Fig. 3a). Histologic sections revealed high-grade urothelial carcinoma involving the entire bladder wall and infiltrating the perivesical tissue and vagina; additionally, the mesh was embedded within the firm cancer tissue (Fig. 3b,c). No obturator lymph node metastasis was indicated. Pelvic pain initially improved but recurred 6 weeks later. CT revealed a tumor recurrence in the left internal obturator muscle (Fig. 4) and sacral metastasis. Consequently, the patient died 5 months postoperatively.

Discussion

We report a rare case of bladder cancer rapidly invading the pelvic wall muscle in a patient after a TVM procedure. The carcinogenicity of synthetic mesh in humans has been debated. Two cases of squamous cell carcinoma have been reported in postoperative polyester mesh surgeries of inguinal hernias that might have been caused by long-term chronic infection.8 Although chronic inflammation due to implanted material was an etiological factor of carcinogenicity, there is no solid evidence of the carcinogenicity of polypropylene,9-11 which was the material used in the TVM procedure. In our case, the bladder cancer was a urothelial carcinoma and initially occurred in the bladder mucosa. Therefore, it is unlikely that the bladder cancer was caused by the mesh. However, pathological findings showed desmoplastic high-grade urothelial carcinoma infiltrating around the mesh. The effect of mesh on the progression of existing cancers is not well known. There is only one case report of ovarian serous carcinoma recurrence in a synthetic mesh; the authors speculated that chronic inflammation caused by the insertion of polypropylene mesh affected tumor progression.¹² We speculate that the scar tissues around the mesh may have become an abnormal anatomical pathway, and newly developed bladder cancer may have spread along the mesh from the left internal obturator muscle.

The use of LRC in a patient who has undergone a TVM procedure remains unclear. Usually, radical cystectomy in female patients is performed by incising the vaginal wall at the posterior vaginal fornix and resecting the uterus *en bloc* with the bladder.¹³ By incising the vaginal wall at the anterior vaginal fornix, we could resect the bladder and the anterior mesh without exposing the posterior mesh.

Our case suggests that mesh can become an abnormal anatomical pathway for tumor invasion. Both presurgical screening and long-term postoperative follow-up are needed in patients after a TVM procedure for early detection and treatment of bladder cancer before the tumor invades along the mesh.

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Conflict of interest

The authors declare no conflict of interest.

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