INTRODUCTION
Ectopic prostatic tissue may occasionally be found in the posterior urethra or bladder wall, in accordance with embryonic development.\textsuperscript{1,2} However, to date, ectopic prostatic tissues detected outside the genitourinary tract in the pelvic cavity is very rarely reported, not to mention giant mass lesions.\textsuperscript{3} Since imaging studies are inconclusive with a variety of differential diagnoses, a giant pelvic mass often requires a well-planned open surgery and intraoperative fast frozen section analysis as a diagnostic and therapeutic approach.

CASE REPORT
A 71-year Chinese male was admitted to the hospital due to detection of a lower abdominal mass for eight months, and recurrent acute urinary retention for one week. This patient already had mild lower urinary tract symptoms, such as frequency, nocturia (2-3 times), and dysuria (for three years). He had taken medicines (tamsulosin etc.) irregularly for the treatment of benign prostatic hypertrophy, without any auxiliary examination at a local hospital. This patient suffered from first episode of acute urinary retention approximately eight months ago, and a lower abdominal mass was found by prostatic ultrasonography for the first time. However, this patient refused surgical treatment for fear of high medical risks at that time. After receiving an indwelling catheter for one week, this patient succeeded for avoiding trial with mild urinary symptoms, as previously. In the patient's past medical history, he only had mild hypertension, which was well-controlled for five years.

On physical examination, the blunt round upper margin of the solid mass could be touched in the right lower quadrant of the abdomen, while on digital rectal examination, Grade II prostatic enlargement was detected, and a giant pelvic mass was attached to it. The laboratory study revealed elevated PSA (44.48 ng/ml, normal: 0-4 ng/ml, CA19-9 (48.91 U/ml, normal: 0-39.00 U/ml), and CA242 (21.42 IU/ml, normal: 0-15.00 IU/ml).

The previous study with ultrasonography showed a heterogeneous mass located under the bladder, which was approximately 12.0 x 10.6 cm in size, but the renal collecting system was not affected. On computed tomography (CT) study, the tumor was well-circumscribed and heterogenous, and no sign of lymphadenopathy or osseous lesions were noted (Figure 1). MRI revealed a large mass lesion (approximately 12.0 x 10.7 x 9.5 cm) in the pelvic cavity surrounded by a capsule-like structure, which was posterior to the cranially displacing bladder, and closely correlated to the left lobe of the prostate and seminal vesicle (Figure 2). The tissue had a weak signal only on T2 scans (Figure 2C). A tentative diagnosis of pelvic low malignant potential sarcoma was made, and the possibility of adenoma with prostate or seminal vesicle origin could not be ruled out.

A suprapubic incision for pelvic exploration and tumorectomy was chosen. Intraoperative frozen section analysis suggested the pelvic mass lesion as an adenoma of unknown origin with moderate dysplasia. Hence, complete tumor resection and right ureteral re-implantation (in order to avoid rupture of right ureter and ureterovesical junction) were performed under general anesthesia (Figure 3). During the procedure, it was found that the encapsulated tumor mass, posterior to the bladder, occupied the whole pelvic space, with its pedicle-like upper part located slightly left to the seminal vesicle and prostate. The pathological study suggested no signs of infiltrating growth and no carcinomatous
areas were present in the whole specimen, and the tumor had an intact fibrous capsule surrounding hyperplastic glands and mesenchymal tissues, with some glands having a cystic appearance. The glands were lined by double-layered epithelial cells and exhibited no signs of mitoses and prominent nucleoli (Figure 4A). Immunohistochemically, these tumor cells were positive for PSA (Figure 4B), 34BE12, CD10 and cytokeratin (CK), but negative for CK7, CK20, CEA, P63 and P504S. The pathological diagnosis was ectopic prostatic adenoma.

FIGURE 1: Computed tomography (CT) scan showing the 12 × 11 cm pelvic mass. The mass adjacent to the prostate was well-circumscribed with a surrounding clear boundary, and was slightly enhanced on contrast scan.

FIGURE 2: Magnetic resonance imaging (MRI): (A) the T1-weighted SE sequence of the MRI; (B) the sagittal plane of the pelvic imaging; (C) the T2-weighted SE sequence of the MRI; (D) the coronal plane of the pelvic imaging. The arrows show the tumor mass. MRI reveals a moderate hypersignal similar to benign prostatic hyperplasia on the T2-weighted SE sequence.

FIGURE 3: The specimen of the giant adenoma was obtained from the tumorectomy. It was a large mass lesion of approximately 12.0 × 10.6 × 9.0 cm.

FIGURE 4: Hematoxylin-eosin (H&E) staining (magnification, ×400) and immunohistochemistry results of the prostate specific antigen (PSA) detection. (A) Glands with light-pink secretions in the glandular lumens were formed by double layered epithelial cells, which had a flat and columnar shape, and a pale cytoplasm, but presented no signs of mitoses and prominent nucleoli. (B) The immunohistochemistry of the tumor cells were positive for PSA.

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DISCUSSION

Ectopic prostatic tissue, which is an uncommon aberration, may be incidentally found in the urinary tract, such as the posterior urethra and bladder. However, occasional cases may be found in testis, epididymis, penis and seminal vesicle, and even in the anal canal.4-6 This suggests that the embryonic prostatic tissue can migrate and was be found in ectopic sites. In the present case, authors thought that this ectopic prostatic lesion may have originated from seminal vesicle due to the anatomic position of the mass.

Ectopic prostatic tissue outside the urinary tract seldom causes low urinary tract symptoms, unless the lesion has reached large size to cause compression in the tract. Hence, it remains difficult to detect this at an earlier stage without an imaging study. When pelvic giant mass lesions develop and are found, it is crucial to obtain a clear working diagnosis of the tumor entity prior to planning the therapeutic procedure. A single imaging test (ultrasound, CT, or MRI) is often not sufficient, and it cannot reveal the exact anatomic relations, although MRI appears to be the best imaging modality to determine the nature of the pelvic mass. For the present case, MRI revealed a moderate hypersignal similar to benign prostatic hyperplasia on the T2-weighted SE sequence, and a low-signal ring was attributed to the capsular fibrous tissue at pathology. In addition, the mass was in contact with the left seminal vesicle and base of the prostate and beneath the bladder. This may suggest its possible origin from ectopically situated prostatic tissue in the seminal vesicle.

Tumors in adult pelvis of male patients may have diverse origins, such as teratoma, lipoma, fibroids, lymphoma, adenocarcinoma, sarcoma, etc. Hence, it is often difficult to arrive at a precise diagnosis before surgery. Since the present differential diagnoses included sarcoma of the pelvis, the investigators deliberately preferred intra-operative frozen section to preoperative transrectal biopsy due to fear of potential tumor spreading. During the surgical procedure, prophylactic isolation and re-establishment of the pelvic organs may have to be taken for the complete resection of such giant pelvic tumor.

In conclusion, the recognition and awareness of this unusual lesion is important, in order not to confuse this particular lesion with other pelvic tumors.
REFERENCES


