

Primary Urothelial Carcinoma of the Anterior Urethra

Anterior Üretranın Primer Ürotelyal Karsinomu

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Abstract

We report the case of an 89-year-old male with an isolated anterior urethral invasive urothelial carcinoma. This is a rare primary tumour of the anterior urethra, as this area is not lined by urothelium. It was managed with radical penectomy and perineal urethrostomy, and the patient has no recurrence to date.

Keywords: Urothelial carcinoma, Urethra, Penectomy

Öz

İzole ön üretral invaziv ürotelyal karsinomu olan 89 yaşında bir erkek olguyu sunuyoruz. Bu, anterior üretranın nadir görülen primer tümörüdür, çünkü bu bölge ürothelium ile kaplı değildir. Bu radikal penektomi ve perineal üretrastomi ile tedavi edildi ve hastanın bugüne kadar nüksü olmadı.

Anahtar Kelimeler: Ürotelyal karsinom, İdrar yolu, Penektomi

Introduction

Primary urethral carcinoma is very rare, being less than one per cent of urological malignancy, and only a small proportion of this is urothelial carcinoma, which is predominantly seen in the posterior urethra (1,2,3). Primary urothelial carcinoma of the anterior urethra is unusual because this area is not normally lined by urothelium. Postulated mechanisms include the presence of foci of ectopic urothelium or metaplastic change (4). One case in 2006 detected human papillomavirus (HPV) type 16 in a grade 3 urothelial carcinoma of the fossa navicularis, suggesting that HPV may play a role in development of urothelial carcinoma particularly in immunosuppressed patients and that this may also be influenced by dissemination via urethral instrumentation (3). However, this role is likely a minor one (5).

Case Presentation

An 89-year-old man was referred for painless macroscopic haematuria and obstructive lower urinary tract symptoms. He was a lifelong non-smoker, with no significant family history

or risk factors for urothelial carcinoma. Abdominal examination was unremarkable, external genitalia were normal and the prostate was small and firm. There was no palpable inguinal lymphadenopathy.

Urine cytology showed small clusters of highly atypical urothelial cells with large numbers of atypical spindled cells with dense orangeophilic cytoplasm, suspicious for high-grade urothelial carcinoma with squamous differentiation. Computed tomography (CT) urography showed no upper tract abnormalities.

Rigid cystourethroscopy found a solid pale tumour in the penile urethra, almost entirely occluding the lumen (Figure 1). A guidewire was passed beyond the tumour, which was then debulked, however, poor visibility prevented adequate examination of the bladder and a catheter was left in situ. Histology showed high-grade papillary urothelial carcinoma with no invasion seen.

Three weeks later, repeat rigid cystourethroscopy showed circumferential polypoid tumour involving a five centimetres

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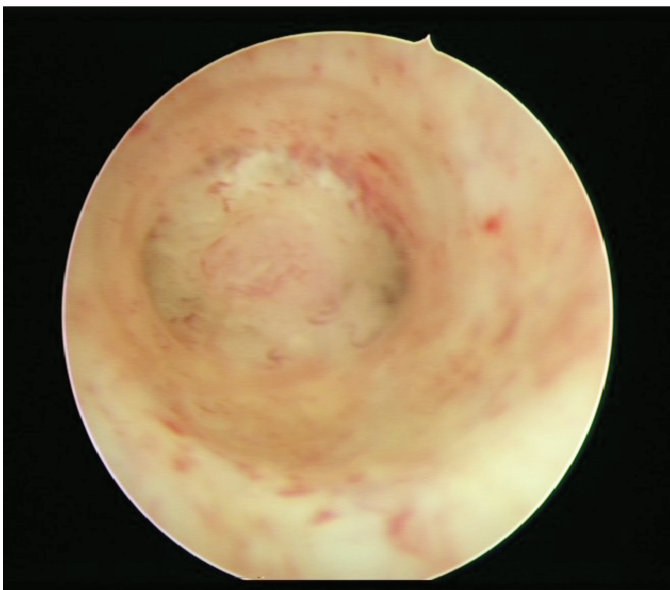


Figure 1. Rigid cystourethroscopy showing urethral lumen occluded by tumour

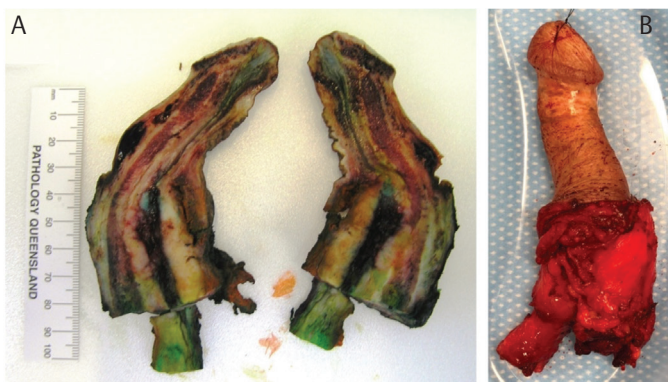


Figure 2. A) Macroscopic appearance of radical penectomy and urethrectomy specimen, with tumour invading corpus spongiosum and corpus cavernosum. B) Haematoxylin and eosin-stained section showing urothelial carcinoma with squamous differentiation

segment from penoscrotal junction into mid-bulbar urethra. Erythematous regions within the bladder were biopsied at the posterior, left and right walls. The urethral tumour was further debulked and diathermied, and the catheter was replaced.

Discussion

Histology confirmed papillary and endophytic high-grade urothelial carcinoma with some squamous differentiation, invading muscularis propria. The bladder biopsies contained only

cystitis cystica and mixed inflammation in the lamina propria. Repeat voided urine cytology yielded atypical urothelial cells. CT abdomen/pelvis had no suspicious lymphadenopathy.

With confirmation of invasive urothelial carcinoma, the patient then proceeded to radical penectomy, urethrectomy and perineal urethrostomy. He recovered well and a catheter was left in situ for six weeks. Histology showed pT3 high-grade urothelial carcinoma, arising at the penoscrotal junction 65 millimetres from the urethral meatus, invading into the corpus spongiosum and corpus cavernosum (Figures 2A, 2B). Margins were clear.

Ethics

Informed Consent: Consent form was filled out by all participants.

Peer-review: Externally peer-reviewed.

Authorship Contributions

Concept: M.S., D.D., **Design:** M.S., D.D., **Data Collection or Processing:** M.S., D.D., **Analysis or Interpretation:** M.S., D.D., S.M., **Literature Search:** M.S., **Writing:** M.S.

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