# CASE REPORT

# Clear cell adenocarcinoma of bladder expressing prostate-specific antigen, alpha-methylacyl-CoA racemase and CD10

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#### ABSTRACT

Backgrounds: Clear cell adenocarcinoma (CCA) of urinary bladder (UB) is rare.

Aims: To report a case of CCA of UB immunohistochemically expressing prostate-specific antigen (PSA), alpha-methylacyl-CoA racemase (AMACR) and CD10; thus this case was difficult to differentiate from metastasis from prostatic adenocarcinoma or renal cell carcinoma (RCC).

**Case:** An 84-year-old man presented with dysuria and hematuria, and cystoscopy revealed a bladder tumor. The biopsy showed proliferation and invasion (more than pT2) of high-grade carcinoma cells focally forming glandular structures. Lymphovascular permeations were recognized. Focally, usual urothelial carcinoma (UC) was seen and there were gradual direct transitions between the clear cell carcinoma and UC. Immunohistochemically, the carcinoma cells were positive for PSA, AMACR, CD10, p53, Ki67 (labeling = 10%), uroplakin II, thrombomodulin, and cytokeratin (CK) 20, but negative for Hepatocyte-paraffin1 (HepPar1), CK7, PAX8, and TTF-1.

**Conclusions:** A rare case of CCA of UB with immunoexpressions of PSA, AMACR and CD10 is reported. The aberrant expressions of these three antigens are only speculative. The gradual transitions between UC and CCA and the presence of urothelial antigens in this tumor suggest that the tumor might have originated from UC.

Key Words: Urinary bladder, Clear cell adenocarcinoma, Prostate-specific antigen, Alpha-methylacyl-CoA racemase, CD10

## **1. INTRODUCTION**

Carcinoma of the urinary bladder (UB) is common and characterized by frequent multicentricity and high rate of recurrence. Most of UB carcinoma is urothelial carcinoma (UC) of various grade, followed in order by ordinary adenocarcinoma and squamous cell carcinoma. Clear cell adenocarcinoma (CCA) of UB, first coined by Young and Scully<sup>[1,2]</sup> is rare; to the author' knowledge, 29 case reports or case series have been reported.<sup>[1–29]</sup> It can be present in focal foci in other types of bladder carcinomas. CCA of UB is different from much more common clear cell carcinoma which is defined as carcinoma having hobnail cells and glycogen-rich clear cells. High specificity and high sensitivity are most important in immunohistochemical studies. Various tumorspecific proteins with high sensitivity and high specificity were discovered in recent years. For example, TTF-1 is specific for thyroidal and lung alveolar epithelium and their tumors, CDX-2 for gastrointestinal and pancreatobiliary ep-

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ithelium and their neoplastic counterparts, surfactant apoprotein A (SA) for pulmonary alveolar epithelium and its tumors, prostate-specific antigen (PSA) for prostatic glands and their tumors, alpha-methylacyl-CoA racemase (AMACR) for prostatic adenocarcinoma, CD10 for renal cell carcinoma (RCC) and B-cell lymphoma and Hepatocyte-paraffin1 (HepPar1) for hepatocytes and hepatocellular carcinoma (HCC), and so many proteins. These so called "specific" antigens or proteins have been used to determine cell types or even malignant potentials for many tumors. Only a little is known about immunoexpression in CCA of UB, while ordinary urothelial carcinomas of UB are known to express EMA, high-molecular-weight cytokeratins, p63, p40, thrombomodulin, uroplakin II, invulcurin, and other molecules. The author herein reports a case of CCA of UB expressing PSA, AMACR and CD10.

## 2. CASE REPORT

An 84-year-old man presented with dysuria and hematuria. The patient received cystoscopy which revealed a polypoid bladder tumor measuring circa  $3 \text{ cm} \times 3 \text{ cm}$  at the triangle. Digital rectal examination of prostate was normal and serum PSA level was normal (3.8 ng/ml). Radiologic imaging studies showed the UB tumor, but no other tumors in the body including kidney, ureter and renal pelvis. Many biopsies were taken from the UB tumor, like transurethral resection of bladder tumor (TUR-BT), and they showed proliferation and invasion (positive muscular invasion, pT2 or more) of high-grade (Grade 3) carcinoma cells with very clear cytoplasmas (see Figure 1A-B). Glandular structures were focally noted (see Figure 1A-C). Lymphovascular permeations were evident. Focally, usual UC of Grade 2 was seen in the surface and there were gradual direct transitions between the clear cell carcinoma and UC (see Figure 1C).

A histochemical stains using periodic acid-Schiff after diastase digestion (d-PAS), mucicarmine and alcian blue at pH2.5 showed a little mucin in the tumor cell cytoplasmas. No significant glycogen was seen in the tumor cell cytoplasmas by PAS and d-PAS stainings. Fat stains using formalin-fixed tissue revealed no fat. An Immunohistochemical (IHC) study was carried out with the use of Envision method. In the much dominant adenocarcinoma areas, it was as follows: PSA + (see Figure 1D), AMACR + (see Figure 1E), CD10 ++ (see Figure 1F), Uroplakin II + (see Figure 1G), thrombomodulin +, p53 + (see Figure 1H), Ki67 (labeling = 20%) cytokeratin (CK) 20 +++, CKAE1/3 ++, CKCAM5.2 +, CK34BE12 +. CK5 +, CK6 +, CK14 -, CK18 +, p63 -, HepPar1 -, CK7 -, CDX-2 -, PAX 8 -, RCC marker -, and TTF-1 -. In the very tiny UC areas, it was as follows: PSA -, AMACR -, CD10 -, Uroplakin II +, thrombomodulin +, p53 -, Ki67

(labeling = 4%) cytokeratin (CK) 20 +, CKAE1/3 +, CK-CAM5.2 -, CK34BE12 +. CK5 +, CK6 +, CK14 -, CK18 -, p63 -, HepPar1 -, CK7 -, CDX-2 -, PAX 8 -, RCC marker -, and TTF-1 -.

The patient is now treated by chemotherapy (cisplatin-based), focal radiation (40Gy) without operation because of his high age. No obvious tumor reduction was recognized. The patient is still alive with the bladder tumor 5 months after the histological diagnosis.

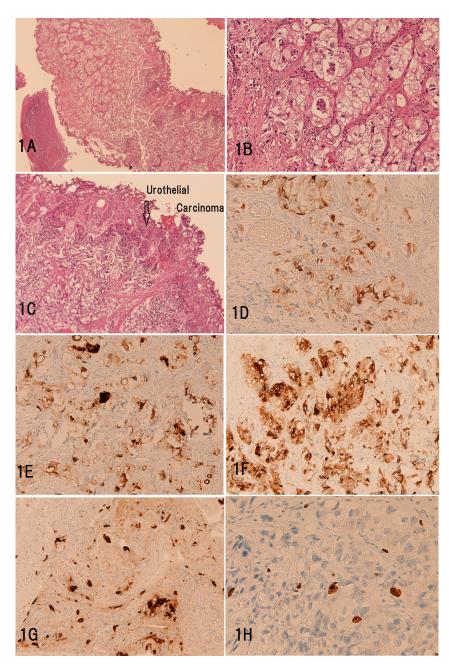
## 3. DISCUSSION

CCA of UB should be differentiated from much more common clear cell carcinoma which is characterized by glycogenrich clear cells and hobnail cells. A review of CCA of UB in English literature showed 42 cases (SE) with mean age of 54 years (9 years) and with female predominance (M:F=12:29).<sup>[1-29]</sup> Most patient presented with hematuria or dysuria. The treatment was variable, because no clinical tests of this tumor for treatments are available; it was almost the same as treatments of ordinary UC (endoscopic, operation, irradiation and anti-cancer drugs). The most interesting was the histogenesis of this tumor; this tumor seems to be heterogenous. Suspected origins are Mullerian ducts, endometriosis, and nephrogenic metaplasia.<sup>[1-29]</sup> In the present study, the CCA was histologically in contact with UC cells and immunohistochemically expressed urothelial antigens (thrombomodulin and uroplakin II), suggesting that the present CCA of bladder developed from UC. The present case is male patient; therefore the former two hypotheses are unlikely. The author thought that the present CCA of UB have arose frpm preexisting UC of UB.

The present UB tumor is apparently a carcinoma. The present tumor is not usual UC. The present tumor is highly likely to be adenocarcinoma because of the presence of focal gland formations and mucin.<sup>[1–3]</sup> However, the histology resembles RCC, HCC, prostatic adenocarcinoma, and clear cell malignancies of other primary sites. The present case is apparently different from papillary urothelial neoplasma and inverted papilloma. The tumor somewhat resemble nephrogenic adenoma but is different from it because of positive malignant features. The tumor should be differentiated from urachal carcinoma, but it usually shows colonic-type adenocarcinoma and most often arise from tip of bladder.

Therefore IHC investigation was performed, and it showed positive reactions to PSA, AMACR, and CD10. In general, AMACR is known to be relatively specific to prostatic adenocarcinoma, but it is shown that non-prostatic carcinomas infrequently are positive for AMACR, a pitfall of diagnostic pathology of prostatic carcinoma. CD10 is relatively specific for B-cell neoplasma, myoepithelial cells, mesenchymal cells, in addition to RCC. The present case shows that CCA of UB can have positive immunoproducts of PSA, AMACR and CD10. The problem in this case is positive for PSA immunoreaction. PSA is now though to be highly specific for prostatic gland and its malignant counterparts. However, ectopic production of PSA by non-prostatic tissues has been reported.<sup>[30,31]</sup> Ectopic prostate is also reported. In Dennis report,<sup>[32]</sup> the sensitivity and specificity of PSA for prostatic adenocarcinoma is 100% and 99%, respectively. However, it was shown that PSA is expressed in vast kinds of carcinomas. From these findings, it is highly conceivable that the present tumor is primary CCA of UB ectopically producing PSA,

AMACR and CD10. Positive expression of AMACR in CCA of UB was reported only once,<sup>[14]</sup> and positive expression of CD10 may support the nephrogenic origin of the current tumor. The possibility that the present tumor is UB extension from occult prostatic adenocarcinoma is still remains now. The findings that the present tumor cells showed mucins and the observation that the present CCA cells are in continuity with overlying UC component, are strongly suggestive that the present tumor is primary CCA of UB. It is quite unlikely that the present tumor are also not compatible with prostatic adenocarcinoma, but are consistent with primary UB carcinoma.



#### **Figure 1.** Histological (A-C) and Immunohistochemical (D-H) features of the bladder tumor

A: The biopsies shows proliferation and invasion of high-grade carcinoma cells with clear cytoplasms. HE,  $\times 20$ . B: High power view. The tumor cells had high grade atypia and very clear cytoplasms. Tubular formations are seen in focal areas. HE,  $\times 200$ . C: This field shows ordinary urothelial carcinoma (upper right) and the clear cell carcinoma cells (lower) with gradual merges between the two. HE,  $\times 100$ . D-F: Immunohistochemistry. The tumor cells are positive for PSA (D), AMACR (E), CD10 (F), uroplakin II (G) and p53 (H). D-G:  $\times 100$ . H:  $\times 300$ .

#### 4. CONCLUSIONS

A rare case of primary CCA of UB with immunoexpressions of PAS, AMACR, and CD10 was reported. The carcinoma

**CONFLICTS OF INTEREST DISCLOSURE** 

The author declares no conflict of interest.

appeared to originate from ordinary UC.

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