Case Report: Immunohistochemical analysis of a case of urothelial carcinoma of the renal pelvis showing gland-like lumina

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Abstract

A Japanese woman in her mid-seventies presented with a case of urothelial carcinoma (UC) of the renal pelvis showing a peculiar gland-like lumina (GLL) pattern and exhibiting no evident glandular structure or mucin production. She showed macroscopic hematuria and experienced left lumbar pain for 2 months. After detailed examination, renal pelvic tumor and bilateral cortical adenoma of the adrenal glands were diagnosed, followed by nephrectomy and adrenalectomy of the left kidney by using the retroperitoneal laparoscopic method. Histopathological examination of the pelvic tumor revealed a grade 2 UC with GLL, which exhibited a smooth transition from ordinary UC to GLL with respect to its morphology. No mucinophilic materials were detected in the GLL areas, which showed distinct trabecular, circular, or meshwork arrangement of tumor cells without a defined glandular structure. Immunohistochemical findings showed that both ordinary UC and peculiar GLL areas were similarly stained when cytokeratin antibodies such as AE1/AE3, CAM5.2, CK7, CK20, and p63, were used. No neuroendocrine differentiation was detected in the GLL. Additionally, the average MIB-1 labeling index of the UC was 5%, but was almost negligible in the GLL areas. Furthermore, the GLL possessed only weak proliferative activity. Thus, prognosis of such a UC with GLL is probably similar to that of an ordinary UC without the GLL. Because of its rarity and the presence of a peculiar histopathological pattern of tumor cells, the histogenesis of such a tumor should be further investigated in further accumulated cases. (to be the same character with the others, please)

Keywords: Renal pelvis, Urothelial carcinoma, Gland-like lumina, Immunohistochemistry, MIB-1

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Introduction

As a variant of the ordinary urothelial carcinoma (UC), UC with peculiar gland-like lumina (GLL) has been rarely reported [1-4]. In some cases, adenocarcinoma was also detected in patients with ordinary UC [5, 6]. The UC cancer cells at times have been found to secrete mucin in the intracytoplasmic or the intercellular areas [4, 7-10]. However, no secretory materials have been observed in the GLL areas [1].

We studied a case of UC with a peculiar GLL, exhibiting a smooth morphological transition from one to the other. Tumor cells were arranged in a circular or a meshwork pattern, with no observable glandular spaces. Moreover, immunohistochemical staining demonstrated similar staining patterns of both the UC and the GLL areas, which

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implied that the GLL areas were the morphological variants of the ordinary UC. We found no differences in the patient's prognosis between ordinary UC and the UC with GLL [1].

Here, we present the immunohistochemical comparisons of both the UC and the GLL components of the reported case.

Case Report

The patient was a Japanese woman in her mid-seventies who showed macroscopic hematuria and had undergone laparoscopic resection of the left kidney and the left adrenal gland. The renal pelvic tumor was found with surface papillary growth, and the tumor cells had proliferated within the pelvic cavity showing a clear border (Fig.1). The excised left kidney and the adrenal gland were fixed in 10% formalin solution and were then embedded in paraffin. Dewaxed sections were obtained and stained with hematoxylin and eosin (HE), mucicarmine and alcian blue stains, and diastase-PAS reaction. Immunohistochemical staining was performed as described previously [11,12] by using labeled streptavidin-biotin complex (LSABC) 2 kit/horseradish peroxidase (HRP) (DakoCytomation, Kyoto, Japan) with diaminobenzidine as the substrate; the antibodies are listed in Table 1.

A written informed consent was obtained from the patient, and her identity has been protected.

Pathological Findings

The tumor proliferated within the thick fibrous stalk attached to the renal pelvic wall, with the cancer cells proliferating in the form of a solid sheet, showing ordinary papillary UC associated with peculiar trabecular or meshwork pattern in the GLL area (Fig. 2 arrows). The GLL area made up almost 30–40% of the entire cancer tissue. In the part of the UC, which was grade 2 in nature, the cancer cells proliferated in the form of a solid sheet and/or peculiar meshwork or trabecular pattern corresponding to the GLL (Fig. 3), with no mitotic figures. At a

Antibodies	Dilution	Pretreatment	Producers	
AE1/AE3	1:400	P ^c	Boehringer-Mannheim Biochimica	
CAM5.2	1:20	Р	Becton-Dickinson	
CK7	1:50	Р	Dako, Kyoto, Japan	
CK20	1:25	AC^{d}	Dako, Kyoto, Japan	
P63	1:50	AC	Dako, Kyoto, Japan	
MA902	1:25	Р	Enzo-Diagnostics	
MA903	1:25	Р	Enzo-Diagnostics	
KL-6	1:2560	NP ^e	Provided by Eisai Co., Ltd.	
CEA ^a	1:25	Р	Dako, Kyoto, Japan	
Chromogranin A	1:100	NP	Dako, Kyoto, Japan	
Synaptophysin	1:25	MW^{f}	Dako, Kyoto, Japan	
NSE ^b	1:100	NP	Dako, Kyoto, Japan	
MIB-1	1:50	AC	Dako, Kyoto, Japan	
HER 2	1:200	MW	Dako, Kyoto, Japan	
Collagen Type IV	1:50	Р	Dako, Kvoto, Japan	

Table 1. Antibodies used in this study (Be careful on red characters in the Table)

^aCEA: Carcinoembryonic antigen, ^bNSE: Neuron-specific enolase, ^cP: Pronase ^dAC: Autoclave, ^eNP: Non-pretreatment, ^fMW: Microwave

Table 2. Results of the immunohistochemical staining

Antibodies	UC ¹	GLL^2	Ep/p^3
AE1/AE3	+	+	+
CAM5.2	+	(+)	+
CK7	+	+	+
CK20	-~(+)	(=)	-
P63	+	+	+
MA902	-	-	_
MA903	-~(+)	-~(+)	+
KL-6	-~(+)	-~(+)	-~(+)
CEA^4	-	-	-
Chromogranin A	-	-	-
Synaptophysin	-	-	-
NSE ⁵	-	-	-
MIB-1	5%	-	-
HER 2	-	-	-

 UC^1 : Urothelial carcinoma; GLL^2 : Gland-like lumina Ep/p^3 : Epithelium/pelvis; CEA^4 : Carcinoembryonic antigen NSE^5 : Neuron-specific enolase

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Figure 1. Paraffin section of the tumor following fixation. Note the intrapelvic tumor with a clear margin (arrows).

Figure 2. General view of the central part of the renal pelvic tumor, revealing the stalk region. Note the meshwork pattern, suggesting a gland-like lumina (GLL) (arrows). HE stain, magnification $20 \times$ *Figure 3.* GLL arrangement of the tumor cells, exhibiting a smooth transition from solid sheet. HE stain, magnification $100 \times$

Figure 4. *Higher magnification of the transitional part, showing the change from solid urothelial carcinoma (UC) to the GLL pattern. HE stain, magnification* $400 \times$

Figure 5. Diffuse positive immune reaction of CK7 for both UC and GLL areas. Labeled streptavidinbiotin complex (LSABC) method, magnification $100 \times$

Figure 6. Weak positive or negative reaction of tumor cells with CK20 immune stain. LSABC method, magnification $200 \times$

Figure 7. Diffuse positive results of GLL with p63. LSABC method, magnification $200 \times$

Figure 8. *Scattered positive nuclei in the UC (lower part), and negative in the GLL (upper part). MIB-1 stain, LSABC method. Magnification* 200×

higher magnification, the GLL areas did not show real lumina, but some contained small blood vessels or edematous connective tissues (Fig. 4). Morphologically, both UC and GLL areas often transitioned smoothly across each other (Figs. 3 and 4). Mucicarmine and alcian blue staining and PAS diastase staining showed no positively stained materials in the tumor cells. As shown in Table 2, immunohistochemically, the cancer cells of both UC and GLL areas were diffusely positive for AE1/AE3 and CK7 (Fig. 5), but not CK20 (Fig. 6). Some of the cancer cells were stained for p63 (Fig. 7), CAM5.2, MA903, and KL-6, albeit weakly. MA902, CEA, and HER 2 staining was negative. Markers of neuroendocrine differentiation, such as chromogranin A, synaptophysin, and NSE, were all negative, including for the GLL areas. Type IV collagen failed to demonstrate linear positivity around the basement membranes in the GLL areas, but showed circular positivity at regions surrounding the blood vessels. Interestingly, proliferative activity of the cancer cells was much higher in UC than in the GLL areas, as revealed by the MIB-1 stain (Fig. 8). Specifically, the UC cancer cells showed approximately 5% positivity with MIB-1 (Fig. 8, lower part), but those in the GLL areas were mostly negative (Fig. 8, upper part).

Discussion

UC with GLL has been rarely described as a variant of

ordinary UC [1-4, 6], but the evidence of divergent differentiation of UC has been reported previously (4, 6). In addition to the adenocarcinoma of the urinary tract, some cases have shown glandular lumina, which contained mucinophilic materials [7-9]. In fact, mucin secretion has been observed in the intracytoplasmic and the intercellular lumina in UC [1, 7]. In the present case, GLL areas showed trabecular meshwork pattern with smooth transition to solid UC, without producing any mucinous materials. Immunohistochemically, both ordinary UC and the one with GLL demonstrated similar staining with various types of cytokeratins and other antibodies as shown in Table 2. Diffuse positive staining for CK7, an important marker of adenocarcinoma in differentiating the metastasizing colon cancer from urinary tract cancer [2], was observed, as shown in Fig 5, in keeping with the findings of previous studies [3, 4], while uneven and weak staining results were obtained for CK20, including at the negative parts, as indicated in Fig. 6. Regarding CK20, positive [3, 4] and negative [2] staining for UC with GLL were observed as previously reported. It has also been reported that HER 2 was positive in both UC and GLL in 1 case [3], but these findings were not constant in GLL [3]. In contrast, our case was completely negative for HER2 in both UC and GLL components.

Interestingly, the proliferative activity of the GLL areas was much lower than that of ordinary UC, as shown in Fig. 8. These results indicate that the prognosis of a case of UC with GLL is similar to that of an ordinary UC, with no relationship with the presence of GLL areas [1]. We suggest that additional immunohistochemical data should be collected by studying cases of UC with GLL in the future.

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