<table>
<thead>
<tr>
<th>Title</th>
<th>A case of papillary renal cell carcinoma suggestive of Bellini duct origin</th>
</tr>
</thead>
<tbody>
<tr>
<td>Author(s)</td>
<td>Gohji, Kazuo; Minayoshi, Kenji; Higuchi, Akihiro; Fujii, Akio; Gotoh, Akinobu</td>
</tr>
<tr>
<td>Citation</td>
<td>泌尿器科紀要 (1994), 40(4): 329-332</td>
</tr>
<tr>
<td>Issue Date</td>
<td>1994-04</td>
</tr>
<tr>
<td>URL</td>
<td><a href="http://hdl.handle.net/2433/115245">http://hdl.handle.net/2433/115245</a></td>
</tr>
<tr>
<td>Right</td>
<td></td>
</tr>
<tr>
<td>Type</td>
<td>Departmental Bulletin Paper</td>
</tr>
<tr>
<td>Textversion</td>
<td>publisher</td>
</tr>
<tr>
<td></td>
<td>Kyoto University</td>
</tr>
</tbody>
</table>
A CASE OF PAPILLARY RENAL CELL CARCINOMA SUGGESTIVE OF BELLINI DUCT ORIGIN

Kazuo Gohji, Kenji Minayoshi, Akihiro Higuchi and Akio Fujii

From the Department of Urology, Hyogo Medical Center for Adults

Akinobu Gotoh

From the Department of Urology, Kobe National Hospital

A case of Bellini duct carcinoma is reported. A left renal tumor was detected by abdominal computerized tomography in a 76-year-old male, although he had no symptoms, such as hematuria, weight loss or flank pain. Radical nephrectomy was performed under the diagnosis of renal cell carcinoma in the left kidney.

Macroscopic examination of the resected kidney revealed a tumor 2.0 cm in diameter, with a yellow-brown cut surface, located in the renal medulla. Histological examinations showed malignant tumor cells with eosinophilic cytoplasm with a papillary growth pattern. Immunohistostaining examinations using Lectin and two kinds of monoclonal antibodies demonstrated no significant staining with soybean agglutinin, peanut agglutinin, Dolichos biflorus agglutinin, Lotus tetragonolobus agglutinin or cytokeratin, and negative staining with Tamm-Horsfall protein. Although the results of immunohistostaining did not provide support, both macroscopic and microscopic findings strongly suggested that this tumor originated from Bellini duct epithelium (Bellini duct carcinoma). The patient is alive with no evidence of disease 1 year after surgery.

Bellini duct carcinoma is a rare malignant condition and the prognosis is usually poor. Differential diagnosis from other renal or pelvic tumors is difficult and long-term careful follow-up is necessary.

Key words: Bellini duct, Carcinoma, Kidney

INTRODUCTION

Primary epithelial tumors of the kidney are generally recognized as either of transitional cell origin arising from the renal pelvis or as renal cell carcinoma arising from the proximal convoluted tubule.

Recently, Cromie et al. described a kidney tumor with cellular elements of both renal and transitional cells and suggested that its origin was the epithelium of the collecting ducts of Bellini. This tumor is an exclusively malignant condition.

The present case shows features suggesting that it originated from the cells of the collecting duct epithelium (Bellini duct carcinoma).

CASE REPORT

A 76-year-old man visited our hospital because an abdominal computerized tomograph (CT) taken by a local physician showed a tumor in the left kidney. There was no evidence of macroscopic hematuria, flank pain or weight loss. Laboratory examination including cytologic study of urine did not reveal any abnormalities. Chest X-rays did not show any metastases. Excretory urography showed a deformity of the left renal pelvis. Abdominal CT showed an isodense mass located near the renal pelvis of the left kidney, but there was no enhancement by contrast material (Fig. 1). Retrograde pyelography showed compression of the renal pelvis by the
Abdominal CT demonstrated an isodense mass located near the renal pelvis of the left kidney. The tumor was not enhanced by contrast media.

**Fig. 1.**

Macroscopic findings of the resected kidney. A small white-yellow-brown tumor (2.0 cm in diameter) was located in the middle part of the renal medulla.

**Fig. 2.**

Histological findings of the tumor. Malignant tumor cells with cuboidal, eosinophilic cytoplasm, nuclear polymorphism grew in a papillary pattern (H & E stain ×200).

**Fig. 3.**

Table 1. Antibodies, their dilutions, and the results of immunostaining

<table>
<thead>
<tr>
<th>Markers</th>
<th>Sources</th>
<th>Dilutions</th>
<th>Results</th>
</tr>
</thead>
<tbody>
<tr>
<td>Lotus tetragonolobus agglutinin (LTA)</td>
<td>E. Y. Lab</td>
<td>×20</td>
<td>+</td>
</tr>
<tr>
<td>Peanut agglutinin (PNA)</td>
<td>E. Y. Lab</td>
<td>×100</td>
<td>+</td>
</tr>
<tr>
<td>Dolichos biflorus agglutinin (DBA)</td>
<td>E. Y. Lab</td>
<td>×100</td>
<td>+</td>
</tr>
<tr>
<td>Soybean agglutinin (SBA)</td>
<td>E. Y. Lab</td>
<td>×100</td>
<td>+</td>
</tr>
<tr>
<td>Cytokeratin PKK1 (CK)</td>
<td>Lab. systems</td>
<td>×100</td>
<td>+</td>
</tr>
<tr>
<td>Tamm-Horstall Protein (THP)</td>
<td>Cedarlane</td>
<td>×2000</td>
<td>−</td>
</tr>
</tbody>
</table>

+ : weakly positive staining, − : negative staining
staining in our case were not clear, the tumor was strongly suspected of being Bellini duct carcinoma by macroscopic and microscopic findings. The patient is alive with no evidence of disease 1 year after surgery.

**DISCUSSION**

Bellini duct carcinoma is a rare exclusively malignant condition. Only 16 cases have been reported in the English and Japanese literature, since Cromie et al. reported the first case in 1979. Recently, tumors originating from the renal distal collecting tubules were divided into two types. The "mixed-type" is gray-white macroscopically, and histologically contains renal cell carcinoma and transitional cell carcinoma. The "papillary type" is yellow or brown-red macroscopically, with some small, pale yellow areas, and histologically, tumor cells demonstrated eosinophilic cytoplasm, nuclear polymorphism, distinct nucleoli, and are similar to distal collecting duct epithelium with a papillary growth pattern accompanied by an invasion of macrophage and fibrous connective tissue. Such tumors develop from the renal medulla and grow to reach the cortical tissue. Therefore, if the tumor is small, it would be located in the renal medulla only. In our case, the tumor was located in the medulla tissue of kidney, had a papillary growth pattern and tumor cells demonstrated eosinophilic cytoplasm. Therefore, this tumor was diagnosed as "Bellini duct carcinoma" and subclassified as "papillary type".

Recently, several investigators demonstrated immunohistochemical analysis using various antibodies against several portions of normal renal tubules. Kikuchi et al. demonstrated differential diagnosis from common papillary renal carcinoma using 8 monoclonal antibodies. Rumpelt et al. demonstrated 6 cases diagnosed by immunohistochemical studies using monoclonal antibodies. However, the results of immunohistochemical analysis are not clear; the results reported by several investigators differ. Aizawa et al. reported that LTA reacted with the normal epithelium of proximal tubule and Bellini duct; THP reacted with the normal epithelium of distal tubule; and SBA, PNA, DBA, CK and VM reacted with the distal portion of the tubule. Our findings show that PNA, SBA, DBA, LTA and CK reacted weakly, but THP did not react with tumor cells. Results of immunohistostaining in our case did not confirm the results by Kikuchi. A possible reason for weak staining with lectin was that the tumor was kept in formalin solution for a long time therefore the antigen was lost. However, in our case, the tumor was small and located in the renal medulla. Therefore, this tumor is strongly suspected to be "Bellini duct carcinoma".

Usually Bellini duct carcinoma is a high grade malignancy. In their series, Rumpelt et al. demonstrated a very pronounced infiltrative pattern of tumor growth, resulting in replacement of the original renal and hilar tissue by tumorous tissue without signs of compression; they suspected the tumor cells spread via interstitial, intratubular and intravenous pathways. In our case, the tumor size was 2.0 cm in diameter, and there was no evidence of lymph node or distant metastases, but microscopically tumor cells had invaded the surrounding parenchymal tissue of the affected kidney. Although the numbers of cases is small, the prognosis of Bellini duct carcinoma is usually very poor. Therefore, long-term and careful follow-up is necessary in patients with Bellini duct carcinoma.

In our case, although electron microscopic examinations were not done, this tumor was strongly suspected to be "Bellini duct carcinoma" because the histological features were typical of Bellini duct carcinoma and the tumor was located in the medulla of the left kidney.

In conclusion, Bellini duct carcinoma is a rare tumor with a high malignant potential. This tumor should be considered in the differential diagnosis of common renal cell carcinoma or urothelial cancer, and long-term, careful follow up is necessary after radical nephrectomy.
REFERENCES


和文抄録

ペリニ管由来が示唆された乳頭状腫瘍の1例

兵庫県立成人病センター泌尿器科（部長：藤井昭男）

直司 和男，源吉 頭治，橋口 彰宏，藤井 昭男

佩里尼管癌の1例を報告した。患者は76歳男性で血尿、体重減少、側腹痛などの症状を有していたが、腫瘤のため近医で施行された腹部CTで左腎腫瘤が疑われて治療目的で当科へ紹介入院となった。腹部CTでは左腎腫瘤に腫瘍陰影をみらみ左腎腫瘤の診断の下に根治的左腎摘出術が施行された。肉眼的に腫瘤の被膜に剝離黄褐色で直径約2cmの腫瘤を認めめた。病理組織学的に円柱状で好酸性胞体を有する悪性細胞が乳頭状に増殖していた。レクチンおよび2種のモノクローナル抗体を用いた検討では、Soybean agglutinin, Peanut agglutinin, Dolichos biflorus agglutinin, Lotus tetragonolobus agglutininおよびCytokeratinは弱陽性でTamm-Horsfall proteinが陰性であった。われわれの免疫組織学的検討の結果は他の報告と必ずしも一致するものではないが、腫瘤の肉眼的および組織学的所見は、本腫瘤がペリニ管上皮由来であることを強く示唆した。患者は術後1年を経たが腫瘤の再発を認めず健在である。ペリニ管癌は非常に稀な腎腫瘤であり一般に予後は不良である。特に他の腎腫瘤、腎盂上皮癌との鑑別は困難であり術後長期にわたる注意深い經過観察が必要である。

（泌尿器科 40：329-332，1994）