Case Report

Thyroid-like follicular carcinoma of the kidney: A Case report

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ABSTRACT

Primary thyroid-like follicular carcinoma of the kidney (TLFCK) is a least-frequent and least-studied variant of renal tumor which is rich in follicular structures of colloidal substances, otherwise, it is typically negative for the thyroid immunohistochemical markers. We report a 44-year-old patient with mild anemia, who was found a right kidney occupying lesion incidentally during the medical examination. The microfollicular and macro-follicular structures under the microscope and typical negative performance of thyroid immunohistochemical markers is in line with the diagnosis of TLFCK. To our knowledge this is the first report of TLFCK with anemia. Brief review of previously published TLFCKs, we discuss the clinical and histopathological features of limited cases.

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Introduction

TLFCK is a kind rare pathological type of renal tumor, which is potentially considered originating in the renal parenchymal urinary tubule epithelial system [1]. This unique histopathological type of RCC (renal cell carcinoma), described firstly in 1996 by Angell et al, had not been recorded in the 2004 world health organization literature to date [2-15]. TLFCK has a similar feature with the follicular carcinoma of the kidney in morphology. It is rich in follicular structures of colloidal substances, conversely, it is typically negative for the thyroid immunohistochemical markers [1]. Here we report a case of TLFCK with mild anemia and discuss the clinical presentation and histopathological features.

Case report

A 44-year-old woman who was found a right kidney occupying lesion incidentally during the physical examination was admitted to the hospital on April 21, 2017, no relevant history of related family disease noted. She had no gross hematuria, no flank pain, no weight loss, no obvious abnormality in the abdominal physical examination and no palpable lesion in the thyroid gland. Routine laboratory tests revealed that HGB was 87 g/L, HCT 32.10 %, MCV 63.2 fL. The results of the other routine laboratory tests, including blood routine, urine routine, liver and kidney function, blood biochemistry, coagulation series, PCT, were all within normal reference range. Electrocardiogram, chest radiography, lung function test was all negative. ECT prompted a delay of the right renal pelvis drainage. CFR suggested that bilateral renal function was normal. Computer tomography (CT) demonstrated that a heterogeneous mass with obvious calcification and well-defined border, low contrast enhancement, measuring 20mm, in the right kidney (Figure 1). There was no evidence indicated the visible invasion of the renal vascular and regional enlarged lymph nodes. The patient underwent a laparoscopic partial right nephrectomy. Microscopically, the carcinoma is composed of microfollicular and macro-follicular structures similar to thyroid follicles (Figure 2). Immunohistochemically, the tumor was positive for CK7, and negative for TG, TTF1, CD10, P504S, CD117 (Figure 3).

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Discussion

Primary thyroid-like follicular carcinoma of the kidney (TLFCK) is a least-frequent and least-studied variant of renal tumor, named purely from the light microscopic appearance, with well-circumscribed pseudocapsules resembling to thyroid follicular carcinoma microscopically but negative for the thyroid-specific markers [2, 3, 15]. In all reported cases, there were no significant distinguish from other types of renal tumor on location and clinical performance. The age of most patients ranges 19 to 83 years old, and the female is more effected than the male [1, 5, 11, 12]. Most of them were incidentally found with or without other clinically positive performance [14]. The macroscopic hematuria and abdominal pain are thought as the common characteristics around TLFCK [10]. It is also difficult to distinguish between TLFCK and other pathological types of renal cancer through imaging performance, as there has been no specific clinical characteristics of TLFCK. In particular, the patient, in this case, was suffering from mild anemia, and the condition gradually recovers after surgery. But, the association between TLFCK and anemia remains unclear.

The diagnosis of TLFCK mainly depends on the microscopic appearance and immunohistochemical experience. Morphologically, TLFCK consists of different sized follicular structures containing abundant eosinophilic colloid resembling to the well-differentiated follicular thyroid carcinoma, as a comparison, TLFCK show a obviously different morphological features from other types of renal carcinoma[6]. Immunohistocemically, TLFCK has a typical negative performance of thyroid immunohistochemical markers, like thyroid transcription factor-1 (TTF1) and thyroglobulin (TG), respectively, the staining of CD10 and CK17 is variable [1, 3, 5, 7]. In this case, Immunohistochemical results show a positive result of CK7, and negative result of TTF1, TG, P504S, CD117, meeting the diagnostic criteria of TLFCK.

The differential diagnosis of TLFCK includes renal metastases of thyroid carcinoma and thyroidization of renal tubules. Thyroidization of renal tubules is a well-known phenomenon, of which the general structure of atrophic distal tubules or collective tubules is similar to that of colloidal transparent tubes of thyroid [14]. In general, the process of glandularization occurs after the chronic pyelonephritis or obstructive nephropathy or the end-stage renal disease [7]. Whereas, TLFCK, which appears as a clear-cut mass, occurs in the patients without suffering from kidney disease attack. TLFCK is composed almost exclusively of dense follicular structures and colloidal substances. The renal tumors described herein are histologically similar to follicular thyroid cancer but still should be distinguished from metastatic thyroid cancer. Thyroid follicular carcinoma usually metastasizes to the lungs and bone, rarely to the kidney. In addition, the tumor shows a positive reaction of thyroid-specific markers TTF-1 and TG [8]. The available data indicated that TLFRCC immunohistochemical staining shows a consistent expression of the transcription factor PAX8, and a typical negative performance of thyroid immunohistochemical markers, which is significant to the identification of thyroid metastatic tumors. Therefore, on the basis of morphology and the result of f immunohistochemistry, the diagnosis of this tumor as TLFCK was made.

The surgery is still the referred treatment [11]. Among the reported TLFCK cases, all of them were low grade and indolent, with only two cases metastases, and no reports of death related to the disease [10].
Figure 3: Immunohistochemical stains. The tumor cells are positive for CK7, and negative for TG, TTF1, CD10, CD117.

Disclosure

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