Renal myxoma: a case report and review of the literature

Renal miksoma: Olgu sunumu ve literaturün gözden geçirilmesi

Sibel Hakverdi¹, Sadık Görür², Mehmet Yaldız¹, Ahmet Namık Kiper²

¹Mustafa Kemal University Faculty of Medicine, Department of Pathology, Hatay, Turkey
²Mustafa Kemal University Faculty of Medicine, Department of Urology, Hatay, Turkey

Abstract
Myxomas are uncommon soft-tissue neoplasms, which are extremely rare in the kidney. In the present case, a right nephrectomy was performed because of suspected presence of a malignant renal tumor. In the macroscopic examination, the kidney contained a well-circumscribed, yellowish, and gelatinous tumor. The tumor was measured 6 cm in diameter. It was composed of fibroblast-like, slender spindle cells with myxoid stroma. With these histopathological findings, the case was diagnosed as renal myxoma. The case was discussed in the light of literature.

Key words: Kidney; myxoma; neoplasm.

Case report
A 59-year-old male was admitted to our hospital with a lower urinary tract symptoms. Ultrasonography of the abdomen demonstrated a well-defined mass occupying the upper portion of the right kidney. The computed tomography (CT) scan showed a 6 cm low-density tumor in the upper portion of the right kidney (Fig. 1). The routine chemistry and the hematologic assessments yielded normal results. A right nephrectomy was performed. The nephrectomy specimen weighed 300 g except the perirenal fat. On sagittal section, a 6x5x4 cm tumor, which was easily distinguished from the adjacent renal parenchyma, was found. The mass was gray-white, semitranslucent, and gelatinous. Mucinous liquid was dripped from the cut surface (Fig. 2). No calculi were identified within the renal pelvis and ureter.

Histopathologic examination revealed a paucicellular and hypovascular tumor consisting of basophilic interstitial mucoid material containing sparse slender, oval, spindle, and stellate cells (Fig. 3). The myxoid material was stained positively with alcian blue and colloidal iron. Lipoblast, giant cells, mitoses, and necrosis were absent. Surrounding renal parenchyma was atrophic.

Immunohistochemicaly, the tumor cells were stained positively as diffuse for vimentin. Negative reactivity was for S-100 protein, epithelial membran antigen, pancytokeratin, and smooth-muscle actin. With these macroscopic, microscopic, and immuno-
histochemical findings, the final histopathological diagnosis was primary renal myxoma.

Discussion

Myxomas were originally described by Virchow in 1863 with structures similar to the mucous tissue of the umbilical cord.[1] Myxomas can be identified in various anatomic locations and the currently recognized subtypes are restricted to the skin, soft tissues, bone, juxta-articular space, sinonasal cavity, maxillary antrum, and in organs or viscera. Organ myxomas may be observed in eye, heart, ovary, and kidney.[2-9]

Primary true intrarenal myxomas are extremely rare. Until 1994, seven case reports of renal myxoma were published. In 1994, Melamed et al.[3] presented two cases of renal myxoma and reinvestigated seven cases previously reported (including his two cases) as renal myxoma, and assumed that only three of them were pure renal myxomas. The remaining five cases exhibited features of sarcoma, fibroepithelial polip, or myxolipoma. They stressed the importance of distinguishing this benign tumor from other benign or malignant mesenchymal tumors.[3] There have been three additional case reports on this entity recently.[2,7,9] Our case might be the eleventh case among the renal myxoma cases reported so far.

Age and gender in all renal myxoma cases were variable. Patients' ages range from 27 to 68 years (mean 47.5 years). The tumors are found more commonly in renal pole, and they vary in size from 4 to 28 cm (mean 10.5 cm) (Table 1).

The radiological appearance of the renal myxoma is various in CT and magnetic resonance imaging (MRI) investigations. In previous reported cases, it was found out that although CT and MRI of the abdomen demonstrated a well-defined mass occupying the middle of the right kidney, and a 8 cm low density solid tumor in the lower pole of the left kidney, in another study CT and MRI of the abdomen demonstrated a well-defined semisolid/semicystic mass occupying the lower calyces of the left kidney.[2,7,9] The findings of the first two studies share similarities.
with our CT result which showed a 6 cm low density solid tumor in the upper pole of the right kidney (Fig. 1). However, in the last study, the appearance of the mass was different from our case in that it has semisolid/semicystic appearance in imaging methods. In the light of these data from literature, it can be concluded that renal myxoma has not typical CT and MRI appearance.

On sections, these tumors can easily be enucleated from the surrounding parenchyma. On histological examination, it was a tumor composed of fibroblast-like spindle cells and abundant myxoid stroma. No giant cells, mitoses, or necrosis were present. Immunohistochemically, the tumor cells were stained positive for vimentin, but negative for S-100 protein, epithelial membran antigen, pancytokeratin, and smooth-muscle actin. In ultrastructural examination, spindle-shaped, fibroblast like cells with long thin cytoplasmic processes, abundant and often dilated rough endoplasmic reticulum, a well-developed golgi complex, and secretory vesicles were disclosed. The nuclei were found to be oval or elongated, and a few with irregular nucleoli. There found to be no basal lamina around the tumor cells. These cells were extensively distinguished by abundant myxoid matrix containing scattered collagen fibrils.[8]

It is essential to distinguish renal myxoma from other tumors exhibiting prominent secondary myxoid feature. These are composed of perineurioma, myxoid neurofibroma, myxoid leiomyoma, myxolipoma, and myxoid variants of malignant fibrous histiocytoma, leiomyosarcoma, rhabdomyosarcoma, and extraskel-etal chondrosarcoma. Grossly, all these neoplasm have an incostinent gelatinous property. However, each tumor has completely different ultrastructural, immunophenotypic, and genotypic features. In the particular case of renal sarcomas, increased cellularity, pleomorphism, high mitoses, well-developed vascularity, and necrosis are usually apparent.[3,7,8]

It has been reported that the patient with renal myxomas has no invasion, no metastasis, or no recurrence. As was in our case, the removal of the affected kidney was conducted in ten patients. Only in one patient with renal sinus myxoma, enucleation of the tumor was applied.[4]

In conclusion, renal myxoma is a benign mesenchymal tumor. Since this tumor has a good prognosis, simple tumor enucleation might be a preferable option for the treatment instead of the complete removal of the kidney. It is essential to distinguish renal myxoma from other highly malignant neoplasms with secondary myxoid features.

**Conflict of interest**

No conflict of interest was declared by the authors.

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<table>
<thead>
<tr>
<th>Reference</th>
<th>Age (year)/gender</th>
<th>Presenting symptom</th>
<th>Side/size (cm)</th>
<th>Situation</th>
<th>Treatment</th>
</tr>
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<tbody>
<tr>
<td>Bolat et al. [5]</td>
<td>27/female</td>
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<td>Left/15</td>
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<td>Right renal colic</td>
<td>Left/7</td>
<td>Lower pole</td>
<td>Nephrectomy</td>
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<td>Asymptomatic</td>
<td>Right/10</td>
<td>Upper pole</td>
<td>Nephrectomy</td>
</tr>
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<td>Appel and Schoenberg [4]</td>
<td>NS</td>
<td>Hematuria for 2 months</td>
<td>Right/8</td>
<td>Parapelvic</td>
<td>Enucleation of mass</td>
</tr>
<tr>
<td>Kundu et al. [5]</td>
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<td>Hypochondrium mass for 2 months</td>
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<tr>
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<td>Lower urinary tract infection</td>
<td>Right/6</td>
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</tr>
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</table>

Table 1. Clinicopathologic data in ten cases of renal myxoma
References


Correspondence (Yazışma): Doç. Dr. Sadık Görür. Urgen Paşa Mah. 75. Yil Bulvan, Cem Apt: No: 10/17, Antakya 31100 Hatay, Turkey. Phone: +90 326 229 10 00 e-mail: sadikgorur@yahoo.com doi:10.5152/tud.2010.037