Solitary fibrous tumor of the kidney
Magnetic resonance imaging characteristics in 4 patients

Zongyu Xie, MD\textsuperscript{a}, Guanghui Zhu, MD\textsuperscript{a}, Liuquan Cheng, MD\textsuperscript{b}, Jinhong Liu, MD\textsuperscript{c}, Huiyi Ye, MD\textsuperscript{b}, Haiyi Wang, MD\textsuperscript{a,}\textsuperscript{b}

	extbf{Abstract}
To investigate the characteristics of magnetic resonance image (MRI) in solitary fibrous tumor (SFT) of the kidney.

MRI findings and clinical features of SFT of the kidney in 4 patients (2 men and 2 women with a mean age of 37.8 ± 8.7 years) were reviewed retrospectively. All patients were scanned by a 3.0-T magnetic resonance (MR) imaging system and the lesions were detected with emphasis on size, shape, location, margin, presence of pseudocapsule, signal intensity, degree of MR enhancement, and apparent diffusion coefficient (ADC).

The 6 tumors from 4 patients were located in renal parenchyma (n = 1), renal pelvis (n = 3), and renal capsule (n = 1). On MRIs, the tumors were round (n = 1) or oval-shaped (n = 4), and presented pseudocapsule (n = 2) and well-circumscribed margins (n = 3) can be found. On T2-weighted images (T2WIs), solid components of the tumor presented homogeneously mild hypointensity or isointensity (n = 4) compared with the renal cortex. On diffusion-weighted images (DWIs), the lesions showed normal or mild hyperintensity (n = 4) with mean ADC of $1.687 \times 10^{-3}$ mm$^2$/s. On dynamic contrast-enhanced MRIs, all lesions showed progressively mild enhancement. In the follow-up of 24 to 36 months after the surgery, 3 patients survived and 1 deceased.

The SFT of the kidney appeared as a circle or oval and presented homogeneously mild hypointensity or isointensity on T2WIs, hyperintensity on DWIs, and progressively mild enhancement on DCE MRIs.

**Abbreviations:** ADC = apparent diffusion coefficient, DCE-MRI = dynamic contrast-enhanced magnetic resonance image, DWI = dynamic weighted image, FOV = field of view, MRI = magnetic resonance image, SFT = solitary fibrous tumor, T1WI = T1-weighted dual-echo image, T2WI = T2-weighted images, TE = echo time, TR = repetition time.

**Keywords:** kidney, MRI, neoplasm, solitary fibrous tumor, spindle cells

1. Introduction
Solitary fibrous tumors (SFT) are the neoplasms of mesenchyma\textsuperscript{1} (or uncommon spindle cell neoplasms), which may occur in any part of the human body but mostly appear in the pleura.\textsuperscript{2} In a literature review, it is reported that SFT can be found in the head and neck regions,\textsuperscript{3,4} sinonasal cavity,\textsuperscript{5} orbit,\textsuperscript{6,7} extracranial region,\textsuperscript{8} larynx,\textsuperscript{9} pleura,\textsuperscript{10} buccal cavity,\textsuperscript{11} thigh,\textsuperscript{12} and paranasal sinuses.\textsuperscript{13,14}

SFT of the kidney is a rare case but usually displays favorable prognosis.\textsuperscript{15} Only 55 cases of kidney SFT have been reported worldwide to date and most of them concentrated in the tumor size, location, and immunohistochemistry without description on the features of magnetic resonance images (MRIs) and follow-up situation.

Magnetic resonance (MR) imaging, based on its excellent soft tissue contrast, has been widely applied to detect and characterize diseases. Besides, a literature review on the adoption of MR scanner has reported its effectiveness on examining the SFT in different parts of the human body such as in the left thigh, head, and neck regions.\textsuperscript{16}

Since inadequate cases reported the disease, SFT of kidney remains a medical mystery, leading to more difficulties in its diagnosis and treatment. Furthermore, images on SFT of kidney were also rare. In light of this, MRI now shoulders the responsibility to figure out the questions upon SFT of the kidney. Therefore, this article aims to investigate the MRI characteristics of SFT of the kidney, based on the cases from 4 patients.

2. Materials and methods

2.1. Patient characteristics
Four patients with pathologically confirmed soft fibrous tumor of the kidney were enrolled in this study from October 2013 to October 2015. Of these 4 participants, 2 were males and 2 were females with a mean age of 37.8 ± 8.7 years (range, 27–47 years). All patients received MRI examination on the tumor with emphasis on size, location, margin, signal intensity, and apparent diffusion coefficient (ADC). Three patients underwent DCE-MR imaging with enhanced hyperintensity, except the last one who was not suitable due to the large tumors in both kidneys.

This study was approved by the ethics committee of Chinese PLA General Hospital and the informed consent was obtained from each patient.
2.2. MRI examination

MRI examinations were performed on a 3.0-T MR imaging system (Signa; GE Medical Systems, Milwaukee, WI) with a surface phased-array coil (gradient magnet strength of 23 mT/m). Patients were scanned in supine position. Morphologic evaluation of the tumors was performed with the following MR sequences: T1-weighted dual-echo image (T1WI) (repetition time [TR] = 260 milliseconds; echo time [TE] = 2.2–2.5 milliseconds, 5.5–5.8 milliseconds; FOV = 36–44 cm; matrix 256 × 192; section thickness 5 mm; gap 1 mm), T2-weighted image (T2WI) (TR = infinite; TE = 90–105 milliseconds; FOV = 36–44 cm; matrix 320 × 224; section thickness 5 mm; gap 1 mm), and 3-dimensional fat-saturated T1-weighted dynamic contrast-enhanced (DCE) image (TR = 3.0–3.9 milliseconds; TE = 1.2–1.6 milliseconds; FOV = 34–40 cm; matrix 288 × 224; section thickness 5 mm; interpolated section thickness = 2.5 mm). Gadobenate dimeglumine (Magnevist; Schering, Berlin, Germany) was intravenously injected with a dose of 0.1 mmol (per kilogram of body weight) at a flow rate of 2 mL/s by a power injector (Spectris Solaris; Medrad, Indianola, PA), followed by a 20-mL flush of normal saline. DCE-MRI was performed in transverse plane at arterial phase, venous phase, and delayed phase. Diffusion-weighted imaging (DWI) was performed before DCE-MRI with b values of 0.80 s/mm².

2.3. Pathologic analysis

All specimens were retrospectively examined by 2 pathologists with an experience in urine pathology for more than 10 years, neither of whom was blinded to the MRI findings.

3. Results

The SFTs of the kidney were located in the renal parenchyma (n = 1), renal pelvis (n = 3), and renal capsule (n = 1). On the MR images, the tumors presented a round shape (n = 1) or an ovoid shape (n = 4). Well-circumscribed margins (n = 3) and pseudocapsule (n = 2) can be detected. On DWI findings, the lesions showed hyperintensity or mild hyperintensity (n = 5) (Fig. 1). The sizes of tumors on renal pelvis from the 3 survived patients were 4.5 × 3.5 × 2.8, 4 × 4 × 3.8, and 4 × 3.9 × 3.7 cm, respectively; and the sizes of 2 tumors on renal parenchyma and renal capsule from the deceased patient was 11.4 × 10.2 × 11.9 and 10.1 × 7.6 × 12.1 cm. The ADC of the 5 lesions were 1.56 × 10⁻³, 1.69 × 10⁻³, 1.58 × 10⁻³, 1.63 × 10⁻³, and 1.97 × 10⁻³ mm²/s, respectively, with a mean ADC of 1.687 × 10⁻³ mm²/s. Solid components of the tumors presented homogeneously mild hypointensity or isointensity (n = 4) compared with the renal cortex (Fig. 2) on T2WIs as well as homogeneous hyperintensity (n = 5) compared with the renal cortex (Fig. 3) on T1WIs. On
DCE MR images (DCE-MRIs) (Fig. 4), all lesions presented progressively mild enhancement, which were confirmed by venous phase images (Fig. 5) and delay phase images (Fig. 6). In the follow-up of 24 to 36 months after the surgery, 3 patients survived and 1 deceased. Three patients underwent transurethral resection of the SFTs, and 1 patient with 2 tumors underwent posterior laparoscopic renal tumorectomy. Pathology examination revealed that the tumors were composed of short spindle cells and appeared with rich vessels, visible coarse collagen fibers, and branching veins. Immunochemical studies showed focal positivity for CD34 (Fig. 7A) and Bcl-2 (Fig. 7B), as well as weak focal positivity for Ki-67 (Fig. 7C).

4. Discussion

As a rare entity, SFT is a spindle cell neoplasm of mesenchymal, which may arise in any part of the human body but uncommon in the kidney.[17] SFTs in the kidney have similar morphologic and immunologic features and biologic behaviors as the SFTs found elsewhere.[18] The disease usually displays favorable prognosis and extremely infrequent retroperitoneal recurrence. Histologically 91% of the SFTs were benign and only 9% were malignant, which presented aggressive behavior.[19] SFT of the kidney can demonstrate an aggressive character, while histopathological features and clinical results are still unknown. Furthermore, more difficulties are found in diagnosing SFT of the kidney, which may lead to unnecessary treatment without obvious malignant findings by current medical facilities.

Among the in hand cases, MRI examination on SFT showed high signal on T1WIs with undetermined characteristics. On T2WIs, the focal region presented high or other signals (for the lower part of the pleura signal), and DCE-MRIs revealed homogenous enhancement features. Besides, fewer reports on SFT by MR scanner, mostly 1 or 2 cases, may lead to inaccurate conclusions. Among the 5 renal masses we examined, T1WIs of 2 lesions presented high or other signals, and 3 lesions showed low signal, which suggested unclear pattern. All the T2WIs showed low or isointense signal on SFT of the kidney. In some literatures, T2WIs also exhibited low or other signals in chest, which are consistent with our findings but different to other parts of the human body.

Despite the fact that MRI can expose the tumor from different perspectives thus reduces the prevalence of misdiagnosis, the technique has been scarcely applied to SFT of the kidney in particular. To the best of our knowledge, only 1 case from the National Center for Biotechnology Information reported the
application of MRI on examining the SFT in the kidney, in which diffuse hypointense was revealed on T2WIs without central necrosis despite the large size of the renal mass.[20] In our research, the MR images show that the SFTs were located at renal parenchyma, renal pelvis, or renal capsule, and the tumors are round shape or ovoid shape. On T2WIs, the solid components of the 4 tumors demonstrated homogeneously mild hypointensity or isointensity relative to the renal cortex. On DWIs, the 4 lesions showed normal or mild hyperintensity with mean ADC of $1.687 \times 10^{-3}\text{mm}^2/\text{s}$.

In an MRI review,[18] patients with kidney SFT basically treated by nephrectomy were followed up between 6 months to 4 years, and as a result most of these patients were healthy without recurrence. Only 2 cases observed metastasis: one is a 50-year-old man with cancer metastasis and recurrence 3 years after kidney removal. Another is a 48-year-old man with cancer metastasis after kidney removal for 8 years. In our study, the follow-up time for 4 patients was between 18 and 30 months. Three patients with only 1 lesion became healthy, and the last 1 with 2 large tumors lost contact, indicating that the patient may depart. Conventional MRI found that the 2 tumors in both kidneys were so large (11.4 x 10.2 x 11.9 and 10.1 x 7.6 x 12.1 cm) that this patient was not suitable for kidney surgery. As time goes by, the renal function gradually degenerated, thus indirectly resulting in fatigue and death finally. It was the first case that both the kidneys have such large SFTs at the same time.

To sum up, we found that the SFT of the kidney appears with the shape of round or oval; homogeneously mild hypointensity or isointensity on the focal masses was detected on T2WIs, hyperintensity or mild hyperintensity was observed on DWIs, and all lesions presented progressively mild enhancement on DCE-MRIs. Considering the results from our study and previous literatures, patient with kidney removal should regularly receive MRI to early address any possibility of relapse, which may contribute to death. Besides, the lack of research in the application of this promising technique on detecting SFT of the kidney request further studies for verification.

Author contributions
Conceptualization: Guanghui Zhu.
Data curation: Haiyi Wang.
Formal analysis: Liuquan Cheng, Jinhong Liu.
Investigation: Liuquan Cheng, Jinhong Liu.
Methodology: Zongyu Xie.
Project administration: Guanghui Zhu.
Software: Jinhong Liu.
Supervision: Zongyu Xie.
Validation: Haiyi Wang.
Writing – original draft: Huiyi Ye.
Writing – review & editing: Zongyu Xie, Huiyi Ye.

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Figure 7. Immunostaining of tumorous cells deliver positive results. (A) Immunostaining of tumorous cells deliver positive results for CD34; (B) Immunostaining of tumorous cells deliver positive results for bcl-2; (C) and Immunostaining of tumorous cells deliver negative results for Ki-67.
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