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<td>Nakatani, Tatsuya; Tamada, Satoshi; Iwai, Yoshihito; Tanimoto, Yoshiaki</td>
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Kyoto University
SOLITARY FIBROUS TUMOR IN THE RETROPERITONEUM: A CASE WITH INFILTRATIVE GROWTH

Tatsuya Nakatani and Satoshi Tamada
From the Department of Urology, Osaka City University Graduate School of Medicine
Yoshihito Iwai and Yoshiaki Tanimoto
From the Department of Urology, Isuzu City Hospital

Solitary fibrous retroperitoneal tumor is rare. We present a case with infiltrative growth in a 56-year-old female patient whose initial symptom was palpable tumor in the lower abdomen. Computed tomography and magnetic resonance imaging indicated a mass in the retroperitoneum under the left kidney with a poorly demarcated infiltrative growth. Surgical findings revealed a gelatinous tumor in the retroperitoneum, which had invaded up to the fatty tissue surrounding the Gerota's fascia and to the fatty tissue surrounding the descending colon. However, as there was no invasion into the Gerota's fascia, it was possible to preserve the left kidney. Pathohistological examination revealed increased cellularity in the tumor tissues as well as tissues with atypical nuclei of the tumor cells with some cell division. Due to these findings, it was diagnosed as malignant solitary fibrous tumor. Only surgical treatment was performed and the patient is alive without recurrence 2 years and 4 months after surgery.

Key words: Solitary fibrous tumor, Retroperitoneum, Malignant tumor, Infiltrative growth

INTRODUCTION

Solitary fibrous tumor (SFT) was first reported in 1931 as a primary neoplasm of the pleura. It is a rare spindle cell tumor that occurs in adults and is known as a benign tumor of the pleura, often described as encapsulated or sharply circumscribed. It rarely occurs in the retroperitoneum, and only 24 cases have been reported in the English-language literature. Among them, only 2 cases were malignant SFT, and our case is the first case of retroperitoneal SFT with infiltrative growth in the surrounding tissues.

CASE REPORT

The patient was a 56-year-old female who noticed a mass in the left lower abdomen and visited our hospital. On physical examination, a hard mobile mass was palpable in the left lower abdomen. Values of carcinoembryonic antigen, carbohydrate antigen 19-9, and cancer antigen 125 were normal. Excretory urography indicated no abnormality in the urinary tract. Barium enema revealed a compression figure in the descending colon. Abdominal ultrasonography demonstrated a somewhat poorly demarcated tumor with heterogenous internal echo. Computed tomography (CT) revealed a slightly heterogenous 9X6X6 cm tumor in the retroperitoneum under the left kidney. Enhanced CT showed a tumor with some deep staining. Magnetic resonance imaging (MRI) T1-weighted image were low intensity similar to the muscles, but T2-weighted images were high intensity and internally partly heterogenous. Angiography showed an internally hypovascular tumor with an indistinct outline, but no feeding vessels. Malignant teratoma or sarcoma occurring in the retroperitoneum was suspected, and laparotomy was performed. On laparotomy, a partly encapsulated tumor was observed and in the retroperitoneum, a poorly demarcated gelatinous substance had spread diffusively up to the fatty tissue surrounding the Gerota's fascia and the fatty tissue surrounding the descending colon. The tumor was removed along with the fatty tissue in the retroperitoneal SFT with infiltrative growth in the surrounding tissues.
Fig. 1. Abdominal MRI. (a) T1-weighted transverse image reveals a tumor (black arrow heads) with a beak-like component extending posteriorly into the fatty tissue and closely bordering the intestines; (b) T2-weighted coronal oblique image shows the high intensity tumor extending up along the Gerota’s fascia.

observed (Fig. 2b). There were both tissues with weak and strong nuclear atypia, and strongly magnified images of the tissue with strong nuclear atypia revealed cells with spindle shaped nuclei and nuclear atypia with some cell division (1 mitoses/10 HPFs) (Fig. 2c). Immunohistological staining indicated the presence of vimentin and CD34 and absence of S-100, actin and CD68. Accordingly, a diagnosis of retroperitoneal malignant SFT was made. No further treatment was performed, and the patient is alive without recurrence 2 years 4 months after surgery.

DISCUSSION

SFT originating in the pleura has been well documented, and cases of extrapleural SFT have also been reported. When the 25 reported cases of retroperitoneal SFT including our present case (Table 1) were examined, there were 11 male and 13 female patients aged 17 to 82 (mean 49.0), with tumor sizes from 2 to 26 cm (mean 10.6 cm). The symptoms in 8 cases were incidental, followed by pain and hypoglycemia in 3 cases. Histologically, 21 were benign and 3 were malignant. Excluding our case, all others were well circumscribed or encapsulated, and ours was the first case of retroperitoneal SFT with infiltrative growth. Surgical excision was performed...
Solitary fibrous tumor of the retroperitoneum is rare, and in particular, no reports have been published on retroperitoneal solitary fibrous tumors with infiltrative growth. In our case, although the tumor was histologically malignant and infiltrative, the outcome of surgical treatment without further treatment was good. The patient is without recurrence at 2 years 4 months after surgery. However, due to reported cases of recurrence after surgical excision, this case is being carefully followed on a long-term basis.

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和文抄録

浸潤性発育を示した後腹膜孤立性線維性腫瘍の1例

大阪市立大学大学院医学研究科泌尿器病態学教室（主任：仲谷達也教授）

仲谷 達也，王田 聡

和泉市民病院泌尿器科（部長：岩井謙仁）

岩井 謙仁，谷本 義明

孤立性線維性腫瘍は稀な疾患である。今回われわれは56歳の女性に発生した悪性後腹膜孤立性線維性腫瘍の1例を報告した。主訴は腹部腫瘤触知で，腹部CTおよびMRI検査にて左腎下方に接して，Gerota筋膜に沿うように発育する境界不明瞭な腫瘤を認めた。手術所見では，後腹膜腔にゼラチン状の腫瘍を認め，下行結腸周囲の脂肪組織やGerota筋膜周囲の脂肪組織に浸潤していた。Gerota筋膜内への浸潤は認めなかったので左腎は温存した。病理組織像では同質の線維線維は著明に増殖し，hemangiopericytoma様のくちばし状を呈する血管も豊富に認められた。腫瘍組織内に腫瘍細胞密度の増加しているところや核異型度が強く分裂像を認める組織が混在しており，また周囲脂肪組織内に浸潤している所見を認めた。以上より，悪性後腹膜孤立性線維性腫瘍と診断した。治療は外科的切除のみを行い，2年4か月を経過しても再発無く生存中である。

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