Intravenous leiomyomatosis with right atrium extension in two patients: A case report

LIANXUE ZHANG¹, YANG DUAN², FULIN SONG³, JUNRUI XIAO² and BENQIANG YANG²

¹Jinzhou Medical University, General Hospital of Shenyang Military Command Training Base for Graduate; Departments of ²Radiology and ³Pathology, General Hospital of Shenyang Military Command, Shenyang, Liaoning 110016, P.R. China

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Abstract. Intravenous leiomyomatosis (IVL) is a rare benign tumor of the uterus mesoderm, which can spread via the vein, from the micro-vein to the inferior vena cava, and even to the right atrium. IVL was first described this disease in 1896 and the first presented case of IVL with right atrium extension in the autopsy was described in 1907. On rare occasions, the tumors may extend into the regional and systemic veins, thus reaching the heart. This may subsequently cause intracardiac leiomyomatosis (ICL), which may lead to congestive heart failure and on occasion, sudden fatalities. Due to its rarity and diffuse symptoms, the misdiagnosis of ICL is common and as a result, the condition may be under-reported. The present study reported two separate cases of IVL resected at the General Hospital of Shenyang Military Command.

Case report

Case 1. A 60-year-old female was admitted to General Hospital of Shenyang Military Command due to a right atrium neoplasm identified during examination. A hysteromyomectomy for uterine fibroids was performed on each patient (Case 1, 41 years ago; Case 2, 3 years ago). One patient presented with chest pain following a period of activity and the other presented with heart neoplasm during a routine health examination.

Introduction

Intravenous leiomyomatosis (IVL) is a rare tumor, which is usually of uterine origin. Birch-Hirschfeld (1) first described this disease in 1896, and following its discovery, Dürck (2) reported IVL with right atrium extension in the autopsy. This tumor type is characterized by intravascular nodular masses of histologically benign smooth muscle that may extend variable distances, including into the inferior vena cava, right atrium and pulmonary arteries. The tumor may arise from uterine leiomyoma, walls of uterine vessels, or myometrium. The age of presentation is usually between 20 and 70 years, with a median age of 45 years. The most commonly affected women are pre-menopausal and multiparous. Intracardiac extension may represent a diagnostic challenge, as it is usually misdiagnosed as a right atrial myxoma and may cause multiple symptoms, including shortness of breath, tachycardia, chest pain, syncope and even mortality. The present study reported two separate cases of IVL resected at the General Hospital of Shenyang Military Command.

Correspondence to: Professor Benqiang Yang, Department of Radiology, General Hospital of Shenyang Military Command, 83 Wenhua Road, Shenyang, Liaoning 110016, P.R. China
E-mail: bqyang888@sina.com

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cava, vena iliaca interna and vena iliaca externa (Fig. 2A). Following surgery, gross pathology was shown in sections of smooth grey white and grey red masses. One mass measured 9.0x4.5x3.0 cm and the others measured 4.0x3.5x1.5 cm in total (Fig. 2B). Histology revealed muscle-like cell arranged in a cord-like formation and shaped as a long fusiform and nucleus shaped long rod (Fig. 2C). The diagnosis was IVL with right atrium and inferior vena cava extension. The patient recovered 28 days later. Regrettably, no follow up was performed.
Discussion

The symptoms of IVL are atypical and have no specialty; therefore, it is often clinically misdiagnosed. Clinical features included pelvic mass, irregular uterus enlargement and bleeding in the vagina, menstruation variation, abdominal distension, and pelvic cavity compression. Before the tumor spreading via the inferior vena cava, all clinical symptoms were hidden and led to misdiagnosis by physicians. However, when the tumor extended to the right atrium, this caused palpitation, chest pain, discontinuous syncope, sudden mortality and even pulmonary transfer (3).

IVL is a rare disease and the CT features of IVL were unique, particularly in cases of mass with heart extension. At an early stage, contrast-enhanced CT and CT vein (CTV) imaging can distinguish IVL. In non-contrast-enhanced CT images, a mass consisting of an isodense lesion was revealed in the vein and heart lumen; and an off-centered filling-defect was revealed in the vein and heart lumen. The mass was dissociated in the lumen with low density and medium heterogeneous enhancement in the contrast-enhanced CT image. The involved the thickening and extension of the vein system in the lumen (4,5). CTV imaging revealed the mass, which performed vascular reconstruction more accurately and directly. The CTV image confirmed the correlation between the tumor and other viscera vessels, collateral vessels, which was significant for pre-operation evaluation and operation strategy.

For IVL with right atrium extension, differential diagnosis includes vascular thrombus, as well as primary and metastatic tumor types. Vascular thrombus was always without enhancement in the contrasted CT scanning. Metastatic differential diagnosis includes metastasis with IVC invasion (e.g. renal cell carcinoma, adrenal cortical carcinoma, hepatocellular carcinoma and lymphoma), atrial myxoma, right-sided heart thrombus or embolus, and leiomyosarcoma. Atrial myxoma is usually the initial diagnosis in these patients given that it is the most common primary heart tumor. However, 60-75% of atrial myxomas are located in the left atrium attached to atrial septum, usually at the fossa ovalis, and no IVC extension exists unless it arises from the IVC proper (in extremely rare cases) (6,7). Another diagnosis to consider is benign metastasizing leiomyomas (BML). This disease is characterized by uterine leiomyoma in young adulthood, with pulmonary metastasis occurring in the pre-menopausal period (8).

Treatment of IVL is almost always surgical (9-11), which includes excision of the extra-uterine tumor and myomectomy, or total hysterectomy, as necessary. The tumor usually spreads via the veins. Therefore, once intravenous leiomyomatosis was suspected, the vein system must be assessed completely and the tumor was excised as completely as possible. As for patients where resection is not possible or where the tumor remains, anti-estrogen treatment was a good choice. Anti-estrogen existed in uterine vascular leiomyoma. The level of estrogen in the uterine vascular leiomyoma was 10-times higher compared with healthy tissue. Kokawa et al (12) suggested that higher estrogen and estrogen receptor levels may influence the growth and invasion of the tumor.

In the present study, IVL, a rare diagnosis that merits consideration in young pre-menopausal or old post-menopausal female patients with cardiac symptoms associated with a right atrial mass. From the two-presented cases, it was clear that radiologists serve a vital role in the diagnosis and follow-up of patients with the diagnosis of IVL. Once patients present with a uterine or peritumeral mass, and deep vein thrombus at the same time, IVL must be considered. Early detection and correct diagnosis is imperative for appropriate treatment and surgical planning.

References