Case 7530

Broad Ligament Leiomyoma
Published on 01.06.2009

DOI: 10.1594/EURORAD/CASE.7530
ISSN: 1563-4086
Section: Genital (female) imaging
Case Type: Clinical Cases
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Patient: 22 years, female

Clinical History:
A 22 year-old female patient presenting with pelvic pain.

Imaging Findings:
A 22 year old woman presented with a 3 months history of persistent pelvic pain. Menstrual cycles were regular and there was no abnormal bleeding. She had menarche at 11, gravida 1, para 1 and using oral contraceptives for 3 years. Physical examination evidenced a firm pelvic mass in the right lateral vaginal recess. Transvaginal ultrasound revealed a solid right adnexal lesion with a diameter of 6 cm, hypoechoic and well circumscribed (Fig 1). To better characterize the lesion and to determine the relationship between the nodule and the surrounding structures, magnetic resonance imaging was performed. MRI demonstrated a low signal intensity lesion on T1- and T2-weighted images (Fig 2-3, respectively), with no claw of myometrium surrounding tumour and no interface vessels, that after endovenous contrast administration enhanced less than myometrium (Fig 4). The findings were consistent with a fibrotic leiomyoma with extruterine and extraovaric origin. The patient underwent surgical resection of this lesion and a tumour of 6.5x5x5cm attached by a stalk to right broad ligament was found. Pathological examination confirmed the diagnosis of a right broad ligament fibrotic leiomyoma weighing 99gr (Fig 5), and showed very good correlation with imaging findings.

Discussion:
Extrauterine fibroids are not as common as uterine fibroids, accounting for less than 3% of all pelvic leiomyomas. These arise from the supporting structures of the uterus, such as the broad ligament, or are free in the pelvis. The leiomyoma of the broad ligament is one of the most common mesenchymal tumours of a uterine ligament and is benign. As a diagnostic criterion, broad ligament leiomyoma must be completely separated from and in no way connected with either the uterus or the ovary. These tumours usually are attached by a stalk to the broad ligament, which is also responsible for their blood supply. Although in most cases broad ligament leiomyomas are asymptomatic, they may present pelvic pain or a palpable pelvic/ abdominal mass. Pelvic pain may be due to pressure effects on adjacent organs, such as bladder or rectum, or torsion.
On US, broad ligament leiomyomas have the same appearance as any other uterine fibroma and present as an hypoechoic solid mass, well-circumscribed, that can be heterogeneous when large. Usually there is no interface between tumour and uterus and no straight relation to the homolateral ovary. As sonograms have false negatives and eventually pedunculated leiomyomas may be mistaken for solid ovarian masses, MRI is indicated whenever ultrasound results are limited, to demonstrate the exact location of the lesion and to characterize it for differential diagnosis. Broad ligament leiomyomas have the typical appearance of leiomyomas on MRI and are depicted as sharply emarginated lesions of low signal intensity on both T1W and T2W sequences due to their high fibrotic content. They are not surrounded by a pseudocapsule of compressed neighbouring tissue and do not show interface vessels imaged as perilesional rim enhancement. Most enhance similarly to the myometrium, whereas larger leiomyomas tend to enhance less and heterogeneously. Postgadolinium images may evidence the presence of a stalk. In addition to standard MR sagittal and axial views, coronal or oblique views may be indicated for accurate
localization and for establishing the most probable origin of a lesion. The differential diagnosis is with parasitic leiomyoma of the broad ligament, subserosal leiomyomas, fibrotic adnexal lesions and other solid tumours from the broad ligament. Parasitic leiomyoma of the broad ligament originates from the uterus and invades the broad ligament, additionally recruiting arterial supply, maintaining or losing its original uterine attachment. Subserosal leiomyomas, if pedunculated, may be hard to distinguish from broad ligament fibroid; identification of the site of attachment is crucial to establish the uterine origin. Ovarian neoplasms include fibroma, adenofibroma, fibrothecoma, Brenner's tumour and metastatic ovarian tumours with highly fibrous component, particularly those from the gastrointestinal tract; usually they originate, and are inseparable, from the ovary, may appear as a complex mass and may have enhancement of solid portions tumour and septa. Other ligamentous mesenchymal tumours are lipoma, neurofibroma, schwannoma and leiomyosarcoma. Leiomyosarcoma is differentiated from leiomyoma microscopically.

Surgical removal is needed for symptomatic relief and for impingement on nearby structures, and because differential diagnosis may include malignancy and pedunculated tumours with risk of torsion.

**Differential Diagnosis List:** Right broad ligament leiomyoma.

**Final Diagnosis:** Right broad ligament leiomyoma.

**References:**

Figure 1

Description: Ultrasonography shows a hypoechoic right adnexal mass, well circumscribed, slightly heterogeneous, with 6.5 x 5 cm. Origin:
Description: Axial T1-weighted TSE image shows a right adnexal mass that is hypointense and slightly heterogeneous. Origin:
**Description:** Axial T2-weighted TSE image shows a right adnexal mass that is markedly hypointense.

**Origin:**
Description: Gadolinium-enhanced fat-suppressed T1-weighted gradient-echo image shows the right adnexal mass with mild and homogeneous enhancement. Origin:
Description: Gross specimen of the broad ligament fibroid. Origin: