

Benign Metastasizing Leiomyoma: Rare Manifestation of a Frequent Pathology



differentiation, without

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Problem Statement:

Benign metastasizing leiomyoma (BML) is a rare variant of uterine leiomyoma, characterized by multiple leiomyomatous lesions in distant locations, most commonly the lungs. Patients are usually asymptomatic and the disease is discovered incidentally. Histopathological confirmation is required for definitive diagnosis. Treatment of BML is controversial. The course of this disease is usually indolent, but requires close surveillance.

Methods: Overview of two clinical cases of BML diagnosed in Portuguese Oncology Institute of Porto in the first semester of 2017 and review of the literature. **Case Reports: Case Report 1 Case Report 2** 48-year-old premenopausal woman Identification 49-year-old premenopausal woman Total hysterectomy 10 years previously for uterine leiomyoma Total hysterectomy 13 years earlier for uterine leiomyoma Medical History Asymptomatic. Incidental diagnosis. **Clinical presentation** Persistent cough. Imaging findings: Chest radiography and computed tomography (CT) Multiple pulmonary bilateral nodules Miliary pattern PET Positron emission tomography: Weak fluorodeoxyglucose (FDG) uptake in lung nodules. Spindle cells consistent **Diagnosis:** with smooth muscle

pulmonary nodule and review of uterine	CAdminution			cellular atypia, necrosis nor mitotic figures.
specimen previously resected. Both specimens showed identical	Immunohistochemical staining	A	C	A. Smooth Muscle Actin;B. Desmin;C. Estrogen and Progesterone Receptors.
immunohistochemical and cytogenetic characteristics.	Cytogenetic study, using "fluorescence in situ hybridization" (FISH)		B	A.22q12 deletion; B.19q13 deletion.
Treatment	Bilateral salpingo-oophorectomy followed by Letrozole.		Bilateral salpingo-oophorectomy.	
Follow-up	6 months: No further development of the disease occured.		3 months: The remaining lesions remained stable.	

Conclusion:

SML diagnosis is challenging and should be based on similar histopathological and immunohistochemical pattern between lung nodules and uterine leiomyoma. When the uterine specimen is not available for retrospective review, the presence of consistent chromosomal abnormalities (deletions of 19q and 22q) is highly suggestive of BML, allowing its individualization as a genetically distinct entity.

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Histopathologic

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