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# Benign splenosis mimicking peritoneal seeding in a bladder cancer patient: a case report

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#### **Abstract**

Introduction: Splenosis is a post-trainal autor asplantation and proliferation of splenic tissue in ectopic sites. These implants may nimic may nancy in healthy patients or peritoneal metastases in cancer patients. When a previous tory of splenic injury is known, the finding of soft tissue nodules in many thoracic and abdominal cations might raise the suspicion of the benign condition of splenosis, in order to avoid unnecessary surgery or chemotherapy.

Case presentation: A 56- r-old man with history of persistent hematuria from bladder cancer was referred to out institution for suspected peritoneal carcinosis. For staging purposes he underwent abdominal compared tomography and ultrasound. The integration of patient's history and imaging . Its led to the diagnosis of peritoneal splenosis. The patient therefore underwent regular T. s U otheral Resection of Bladder for the known malignancy; while no treatment was necessary for plenosis. Two years follow-up was negative for metastases.

C inc. sion: Spienosis is a benign condition after traumatic splenectomy which should be taken to account in the differential diagnosis with peritoneal seeding of malignancy because its appearance may resemble malignancy.

#### uction

Trauma ac disruption of the splenic capsule causes fragments of splenic tissue to be seeded throughout the peritoneal cavity [1]. It has been demonstrated that splenic implants can survive and grow when transplanted to ectopic sites, such as the splenic fossa, any site of the peritoneal cavity, gastro-intestinal tract, liver, kidney, thorax, subcutaneous tissues and even in the head [2].

Since these implants may mimic malignancy or peritoneal metastases, in cancer patients with previous history of splenic injury, the finding of soft tissue nodules in the abovementioned locations might raise the suspicion of splenosis in order to avoid unnecessary surgery or chemotherapy.

This report describes the case of a bladder cancer patient with peritoneal splenosis mimicking metastases.

### Case presentation

A 56-year-old white man native from Italy, was admitted to our Institution for a diagnosis of bladder cancer and a previous CT scan suggesting peritoneal carcinosis. His history started about 4 months in advance for the occurrence of hematuria. His previous personal history was unremarkable, other than a car accident about 20 years in advance, which requested a long hospital staying and a splenectomy.

At our hospital he underwent a Trans-Urethral Resection of Bladder (TURB) which led to the diagnosis of spinocellular carcinoma of the bladder. Usual patient's blood tests, including complete blood cell count and tumour markers, were unremarkable.

The patient underwent a second computed tomography (CT) scan for staging purposes, showing the presence of many peritoneal hypodense masses, with heterogeneous contrast-enhancement during the arterial phase and almost isodense to the hepatic parenchyma during the portal and equilibrium phases (Figure 1).

The spleen was not visualized on the CT so n and lenosis was thereafter suspected as a possible diagnosis and a second-look ultrasound (US) exam confirmed the pres-

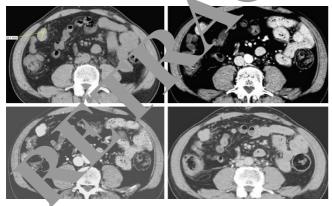


Figure 1
Abdominal CT scan of a splenic nodule before and after contrast medium injection. This 28 mm peritoneal nodule located in the right paracolic space showed slight hypodensity during the pre-contrast phase (a), inhomogeneous contrast-enhancement during the arterial phase (b), slightly iso-hypodenity during the portal (c) and equilibrium phase (d).

ence of peritoneal solid splenic-like masses on the sites indicated by CT (Figure 2).

Diagnosis of post-splenectomy splenosis was confirmed and the patient underwent usual TURV as treatment for bladder cancer, with no chemotherapy, or further treatment for splenosis.

#### **Discussion**

Splenosis is a relatively common finding in clinically silent patients, frequently misdiagnosed accause of the lack of symptoms. According to Normand eacht involves 16% to 67% of patients with pall splenic trauma and/or past splenectomy [3]. On thother hand, Khosravi et all found only five cases of splene as in a retrospective review of a 29-year period of the pathological registry [1], while Lin et all identified eight pathological swith splenosis in a period of 8 years [2].

Based on the location of the splenic nodules, differential diagnoses to be considered may be endometriosis, in presence of personneal seeding [4], peritoneal mesothelioma in case of personneal seeding [5], renal cancer [6] in case of renal implants, abdominal lymphomas in case of retropersonneal locations mimicking lymph nodes [7], hepatic denomas in case of intra-hepatic implants [8] and periton denomas in case of intra-hepatic implants [8] and periton denomas in case of intra-hepatic implants [8].

Thus, patient's history is substantial; secondarily imaging modalities to consider are: CT, US, magnetic resonance (MR), scintigraphy.

If the splenic implant is intra-hepatic, CT imaging may show hypodense masses with strong enhancement at the early phase and pooling enhancement at delayed phase [9]. Grande et al described multiple well-demarcated nod-

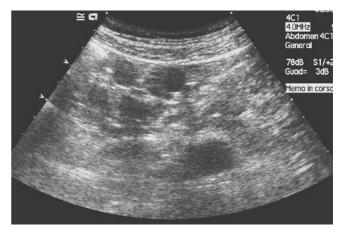


Figure 2
Second-look ultrasound of the nodule in figure 1. US shows homogeneous splenic-like echogenicity of the nodule in the paracolic space, thus confirming its splenic origin.

ules without calcifications, slightly hypodense compared to the liver at non-contrast CT, showing as lobular or oval well-circumscribed structures, hyperdense in the arterial phase and isodense in the portal phase after contrast administration [10]. Accordingly, Imbriaco described a well-demarcated intra-hepatic 3 cm mass, hypodense compared with the surrounding liver parenchyma at unenhanced CT, heterogeneously enhanced in the arterial phase, becoming hypodense during the portal end equilibrium phases [5].

In our case study, according to previous studies, CT scan showed multiple hypodense peritoneal masses with a maximum diameter of 15-20 mm, which characteristically had slightly hypodensity in the pre-contrast phase, inhomogeneous contrast-enhancement during the arterial phase and iso - slightly hypodensity to the liver parenchyma during the portal and equilibrium phase.

Ultrasound may incidentally show hypoechoic masses on the surface of solid abdominal organs, such as the liver [9] or the kidney [5]. It may also show multiple solid masses with a smooth round or ovoid shape, with a homogeneous echo-texture and a hyperehoic peripheral rim, without specific arterial or venous Doppler signals [10]. In our case, a second-look ultrasound, confirmed the presence of multiple solid nodules with homogeneous splenic like echogenicity in the splenic fossa and in the peritone.

MRI may be considered as an alternative moda. If for the identification of splenosis, in case of unce tainty diagnosis with other exams. Splenic implants have been described as hypointense on T1-wei hted images and hyperintense on T2-wieghted images, therefore similar to normal splenic tissue [11,12].

Nowadays scintigraphy (p. formed with heat-demaged 99Tc-labelled red blood ells in still considered the more sensitive and specific imaging modality for the diagnosis of splenosis [10,1,14], being sensitive even in early splenosis, in cases where plenic tissue is minimally present, in functional hyposplenism and in case of poor splenic uptake. The boat se splenic tissue takes up more than 90% of dame od red blood cells [15].

One opic. Is is diagnosed with anyone of the previous modal 25, no more treatment is usually required unless the patient is symptomatic.

#### **Conclusion**

In conclusion this report describes a case of a bladder cancer patient with peritoneal lesions related to personal history of post-trauma splenectomy (splenosis). Since this benign condition may mimic metastases, it should be kept in mind in managing cancer patients with personal

history of post-traumatic splenectomy, in order to avoid unnecessary surgery or chemotherapy.

#### Consent

Written informed consent was obtained from the patient for publication of this case report. A copy of the written informed consent is available for review by the Editor-in-Chief of this journal.

## **Competing interests**

The authors declare that they have no con. ting interests.

# **Authors' contributions**

BR analyzed and interpreted be point data regarding the oncological disease and the recreated treatment. SR performed the radiological examinations (CT and ultrasound) of the abdoruen and was the major contributor in writing the manufactor. LM and MB contributed to the literature search and approve the manufactor. All authors read and approve the manufactor.

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