

## **Chronic Abdominal Pain from Disseminated Splenosis**

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 $K\!E\!Y$   $W\!OR\!D\!S$ : clinical image; abdominal mass; disseminated splenosis; nuclear scintigraphy.

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A 60-year-old woman presented with several years of dull left upper quadrant, mid-epigastric, and right flank pain. Abdominal CT showed multiple soft tissue densities of unclear etiology. She reported having a post-traumatic splenectomy in the 1960s. A technetium-99m-labeled heat-damaged RBC scan (Tc<sup>99</sup>m-dRBC) demonstrated radiotracer uptake in the left upper quadrant, posterior to the right kidney, and along the right diaphragm, confirming ectopic splenic deposits (Fig. 1).

Disseminated splenosis (DS) is a benign condition caused by metastatic deposits of splenic tissue following trauma or surgery. DS is usually asymptomatic and diagnosed incidentally by ultrasound or computed tomography, but can cause site-specific discomfort mimicking endometriosis or peritoneal metastases. Other reported complications include gastrointestinal bleeding, bowel obstruction, and hydronephrosis. Nuclear scintigraphy with Tc<sup>99</sup>m-dRBC localizes ectopic splenic tissue based on the increased uptake of damaged erythrocytes within the reticuloendothelial system. In the setting of previous splenic trauma, this noninvasive technique is a sensitive and specific tool to establish the diagnosis, potentially avoiding invasive tissue sampling or diagnostic laparoscopy.

Patients with DS may retain partial immunoprotection from encapsulated organisms, <sup>6</sup> but no studies have shown an optimal approach to assessing residual splenic function. <sup>7</sup>

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## Compliance with ethical standards:

**Prior presentations:** None.

**Conflict of Interest:** The author declares that he does not have a conflict of interest.

## **REFERENCES**

- Short NJ, Hayes TG, Bhargava P. Intra-abdominal splenosis mimicking metastatic cancer. Am J Med Sci 2011;341:246–9.
- Connell NT, Brunner AM, Kerr CA, Schiffman FJ. Splenosis and sepsis: the born-again spleen provides poor protection. Virulence 2011;2:4–11.
- Schiff RG, Leonidas J, Shende A, Lanzkowski P. The noninvasive diagnosis
  of intrathoracic splenosis using technetium-99m heat-damaged red blood
  cells. Clin Nucl Med 1987;12:785–7.
- Horger M, Eschmann SM, Lengerke C, Claussen CD, Pfannenberg C, Bares R. Improved detection of splenosis in patients with haematological disorders: the role of combined transmission-emission tomography. Eur J Nucl Med Mol Imaging 2003;30:316-9.
- Lake ST, Johnson PT, Kawamoto S, Hruban RH, Fishman EK. CT of splenosis: patterns and pitfalls. AJR Am J Roentgenol 2012;199:W686–93.
- Pearson HA, Johnston D, Smith KA, Touloukian RJ. The born-again spleen. Return of splenic function after splenectomy for trauma. N Engl J Med 1978;298:1389–92.
- de Porto AP, Lammers AJ, Bennink RJ, ten Berge IJ, Speelman P, Hoekstra JB. Assessment of splenic function. Eur J Clin Microbiol Infect Dis 2010;29:1465–73

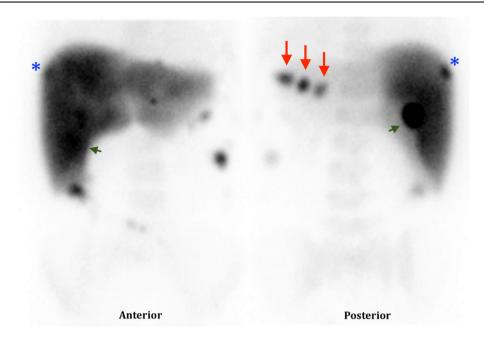


Fig. 1 Maximum intensity projection (MIP) anterior and posterior images of  $Tc^{99}$ m scan showing increased radiotracer uptake in three separate splenic tissue deposits in the left upper quadrant measuring 2.3, 2, and 1.5 cm (red arrows), a 1.7 cm subphrenic deposit beneath the right diaphragm (blue star), and a 3 cm deposit posterior to the right kidney (green arrowhead), all correlating with her sites of discomfort.