Disagreement between ultrasound and magnetic resonance imaging in the identification of schistosomal perportal fibrosis

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Abdominal ultrasound (US) has been widely used in the evaluation of patients with schistosomiasis mansoni. It represents an important indirect method of diagnosis and classification of the disease, and it has also been used as a tool in the evaluation of therapeutic response and regression of fibrosis. We describe the case of a man in whom US showed solid evidence of schistosomal perportal fibrosis and magnetic resonance imaging revealed that perportal signal alteration corresponded to adipose tissue which entered the liver together with the portal vein.

Key words: schistosomiasis - Schistosoma mansoni - ultrasound - magnetic resonance imaging


Magnetic resonance imaging (MRI) has been described in four cases of hepatosplenic schistosomiasis mansoni (Patel et al. 1993, Willemsen et al. 1995, Lambertucci et al. 2002, 2004). The first two studies used US, computed tomography (CT) and MRI in the evaluation of a patient with hepatosplenic schistosomiasis mansoni. The findings of the different methods were in agreement in both studies. Perportal echogenic thickening on US, suggestive of fibrosis, and low attenuation bands about the portal vessels, which markedly enhanced with contrast CT were described. MRI demonstrated the same perportal bands that were seen on liver US and CT scans. These bands showed hiperintense signal on T2-weighted sequences, and were isointense in T1-weighted sequences, with enhancement after contrast administration. The studies of Lambertucci et al. (2002, 2004) evaluated patients with advanced forms of hepatosplenic schistosomiasis by US and MRI. Both methods were again in agreement with respect to the presence of intense perportal fibrosis. Once more, MRI demonstrated broad perportal bands hypointense to the liver on T1-weighted sequences, while they had increased signal on T2-weighted images. Thickening of the gallbladder wall, enlargement of spleen, splenic and portal veins and collateral vessels were detected. After contrast administration, T1-weighted images revealed enhancement of the gallbladder wall and perportal space.

We describe the case of a man in whom the results of US and MRI were not in agreement with respect to the identification of perportal fibrosis.

PATIENTS AND METHODS

A 59-year-old man was referred to the Hospital of the Federal University of Minas Gerais, in Brazil, for the evaluation of schistosomal perportal fibrosis, suggested by an abdominal US performed elsewhere. He was a resident of Belo Horizonte, a city where schistosomiasis is not endemic, but informed previous contact with stream waters in areas were transmission of the disease is known to occur. Examination of nine stool samples did not disclose eggs of Schistosoma mansoni, and no eggs or granulomas were detected by a rectal biopsy. The patient denied previous episodes of digestive bleeding, ascitis or jaundice, as well as chronic prescribed drugs use or alcohol abuse. His clinical examination revealed no abnormalities: liver and spleen were non palpable, and no signs of
portal hypertension of hepatic dysfunction were found. Blood counts also revealed no alterations: $6.28 \times 10^6$ red cells/mm$^3$, hemoglobin 16.7 g/dl, $149 \times 10^3$ platelets/mm$^3$, $6.1 \times 10^3$ white cells/mm$^3$. Serological investigation for hepatitis B and C were negative and blood chemistry and coagulation were unremarkable (albumin 4.5 g/dl, ASAT 29 g/dl, ALAT 39 g/dl, GGT 31 g/dl).

A new US examination of the abdomen was performed using real-time ALOKA SSD 1700 device (Japan) with electronic linear 3.5 MHz transducers. Additional imaging was obtained using a GE 1.5 T Sigma unit (General Electric, New Jersey, US). Axial and coronal 5 mm slice thickness images were performed in T1 and T2-weighted sequences, before and after contrast administration.

**RESULTS**

US of the abdomen showed enlarged central periportal bands with increased echogenicity, suggesting intense central fibrosis (Fig. 1). Echogenic thickening of the gallbladder wall was also detected (Fig. 2). The image pattern was characteristic of hepatic fibrosis produced by *S. mansoni*. No evidence of portal hypertension, such as portal vein or spleen enlargement and collateral veins, was noticed.

MRI demonstrated broad central periportal bands hypointense to the liver on T1-weighted sequences, and with increased signal on T2-weighted images. However, after suppression of fat tissue signal, these periportal bands disappeared (Fig. 3). This finding shows that US periportal thickening suggestive of schistosomal fibrosis corresponded in fact to adipose tissue. Like US, MRI did not detect any evidence of portal hypertension.

**DISCUSSION**

MRI was decisive to put away the diagnosis of schistosomal periportal fibrosis suggested by US in the patient described above. All clinical and laboratory aspects

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**Fig. 1:** ultrasound of the abdomen showing enlarged central periportal bands with increased echogenicity (arrows), suggestive of intense central schistosomal fibrosis.

**Fig. 2:** ultrasound of the abdomen showing echogenic thickening of the gallbladder wall (arrows).

**Fig. 3:** magnetic resonance imaging axial scan demonstrating broad central periportal bands hyperintense to the liver on T2-weighted sequences (A). Scan obtained after suppression of fat tissue signal reveals disappearance of these periportal bands (B). Yellow arrows indicate periportal signal alteration; red arrow points to the portal vein.
of the case reported suggested absence of schistosomiasis, and there were no evidences of the hepatosplenic form of the disease. However, US images in this patient showed solid evidence of schistosomal periportal fibrosis, since characteristic periportal thickening was observed. The hypothesis of schistosomiasis was then made, since periportal fibrosis can occur without enlarged liver or spleen, or without portal hypertension. Surprisingly MRI revealed that periportal signal alteration corresponded to adipose tissue which entered the liver surrounding the portal vein.

Up to 20 years ago, field studies would distinguish hepatosplenic schistosomiasis from other chronic forms of the disease by detecting a palpable spleen on physical examination (Kloetzel 1962, Barreto et al. 1985, Lambertucci 1988a). However, US images in this patient can lead to splenomegaly, and subjects with severe forms of the disease could have not been included in previous studies evaluating morbidity. Symmers fibrosis in patients without spleen enlargement has also been described previously by one US study in Africa (Homeida et al. 1988b, Abdel-Wahab et al. 1992, Lambertucci et al. 1996, Gerspacher-Lara et al. 1998).

Lambertucci et al. (2001) have studied 741 subjects with schistosomiasis in an endemic area of Minas Gerais. US examination identified 15 patients (2%) with intense periportal thickening without spleen enlargement, showing that intense periportal fibrosis can occur without splenomegaly, and that subjects with severe forms of the disease could have not been included in previous studies evaluating morbidity. Symmers fibrosis in patients without spleen enlargement has also been described previously by one US study in Africa (Homeida et al. 1988b) and by one autopsy study in Brazil (Prata & Andrade 1963).

These are examples of how new techniques allow us to better access and comprehend the morbidity in schistosomiasis (Barbosa et al. 1996). MRI has been established as a very sensitive image method in various diseases and, different from US, is not a dynamic study and can be less examiner-dependent. Disagreement between US and MRI (in favor of the second) in the identification of schistosomal periportal fibrosis was found in this case. Studies of series of cases are necessary to compare these image methods and to define the role of MRI in the diagnosis and evaluation of morbidity in schistosomiasis mansoni.

REFERENCES


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