Case Report

Extrahepatic Textiloma Long Misdiagnosed as Calcified Echinococcal Cyst

Federico Cattaneo, 1 Massimo Graffeo, 2 and Enrico Brunetti 1

1 Division of Infectious and Tropical Diseases, University of Pavia, IRCCS S. Matteo Hospital Foundation, WHO Collaborating Center for Clinical Management of Cystic Echinococcosis, 27100 Pavia, Italy
2 Unit of Gastroenterology and Digestive Endoscopy, Poliambulanza Hospital Foundation, 25124 Brescia, Italy

Correspondence should be addressed to Federico Cattaneo; cattomail@alice.it

Received 8 January 2013; Accepted 29 January 2013

1. Introduction

The term textiloma or gossypiboma indicates a gauze pad that is left behind in a body cavity during a surgical operation. This type of complication is uncommon but may cause significant morbidity (close to 50%) and a high mortality rate (11–35%) [1–3]; furthermore, it may represent a diagnostic dilemma with important legal implications [3].

The incidence of textiloma is between 1 in 100 and 1 in 3000 for all surgical procedures [4–7] and 1 case in every 1000–1500 abdominal operations (most commonly complicated by its occurrence) per year [4, 6, 8]. The real incidence, however, may be higher because case numbers are calculated only based on malpractice claims and because of fear of legal repercussions [9].

Therapy consists of the removal of the textiloma on laparoscopy or laparotomy with treatment of complications. Only reoperation allows a definitive diagnosis [1, 10, 11].

We report a case of abdominal textiloma that was initially misdiagnosed as echinococcal cyst and discuss the differential diagnosis based on sonographic features and the WHO-IWGE classification.

2. Case Report

A 55-year-old Italian woman was referred to our clinic for a suspected echinococcal cyst of the liver.

She had type 2 diabetes mellitus, β thalassemia trait, dyslipidemia, and cervical arthrosis and had undergone cholecystectomy in 1973.

In April 2006 an abdominal ultrasound performed at another hospital showed an enlarged liver with regular edges, steatosis, and a focal lesion 9.5 × 7.5 cm in diameter described as a cyst (suspected parasitic) partially solid within segments VI and VII. A hyperechoic area consistent with calcification was also found. The patient reported that she was aware of the cyst but could not provide any documentation. She was asymptomatic and stated that she had always refused to undergo further clinical investigations.

The patient had three further hospitalizations in the same hospital for poorly controlled diabetes in October 2007, May 2009, and March 2010 during which repeated ultrasound scans showed no changes in the lesion, diagnosed as a morphologically stable parasitic cyst.
Figure 1: Gossypiboma seen at TC scan. There is an extra hepatic cystic mass with central calcifications.

During the third hospital stay she underwent a CT that showed a “mass with expansive growth with regular edges 11 cm in diameter, with liquid and calcified content external to the liver parenchyma.” Serology for cystic echinococcosis (complement fixation test) tested negative.

On April 28, 2010, the patient was evaluated again in a hepatology clinic in a different town, with a new serology returning borderline result (1:80 with indirect hemagglutination - IHA). The CT scan done on 03/12/2010 during the third hospitalization was reviewed and considered not suggestive of a parasitic cyst (Figure 1) so the patient was referred to the Division of Infectious Diseases of the Policlinico San Matteo in Pavia for a second opinion.

On May 5, 2010 the patient had a new serology for CE tested in Pavia with IHA (Cellognost™-Echinococcosis;
A textiloma can cause two types of reactions: a fibroblastic reaction, as with a foreign body reaction, when an aseptic process begins (asymptomatic/palpable mass), or an exudative reaction which often leads to abscess (pain, fever, weight loss, fistula, intestinal obstruction or perforation, ileus caused by surgical adhesions, granulomatous peritonitis, and sepsis) [1, 13]. Therefore, clinical presentation of gossypiboma is variable and depends on the location of the retained swab and on the type of biological reaction.

Textiloma can be discovered in the first days after surgery or can remain asymptomatic (hence undiagnosed) for many years and be discovered accidentally.

3. Discussion
In our case the gauze pads left in the patient’s body produced no symptoms for an exceptionally long time (37 years), whereas in the available literature gauze pads have remained undetected on average for 6 to 9 years [14–16]. The longest reported interval between the probable causative operation and the diagnosis of retained surgical spoon is 43 years [3, 4, 14, 17].

A thorough medical history, which can reveal previous surgery, and lab tests together with imaging (ultrasonography, computed tomography, or magnetic resonance) are crucial elements for diagnosis of textiloma [11].

Differential diagnosis includes tumor; cysts, parasitic and otherwise; hematoma, and inflammatory tumor [4, 13, 18, 19], we ruled out CE based on the knowledge of the echinococcal cyst structure (Figure 4). The origin of the patient from an endemic area (Southern Italy) and the negative serologic tests, that would have been in accordance with the cyst being inactive (although calcifications can be found at virtually any stage of the cyst history [20, 21]), were two confounding factors.

In our case, the correct diagnosis was made in two steps: The “calcifications”, however, were not seen at the periphery of the “cyst” as seen in CE, but at the center (Figures 1 and 2).

As seen in the WHO-IWGE ultrasound classification images (IWGE-WHO 2003) [22] (Figure 5) cysts are either fluid filled (CE1, CE3a) or filled with daughter cysts (CE2) or matrix with (CE3b) or without (CE4, CE5) daughter cysts and calcifications are seen around the cyst but not inside [20].

To our knowledge, the only focal infectious lesion that has a central calcification is brucellar abscess (Figure 6) [23, 24].

On closer inspection, though, parallel, wavy hyperechoic lines strongly reminiscent of a gauze pad were seen (Figure 2), which were in line with the patient’s previous surgical intervention on the liver and helped exclude CE.

These two elements were enough to make a diagnosis with CT scan not adding much besides confirming the intrahepatic location of the lesion. In addition, US performs better than CT in staging the cyst [21] and stages as defined by US have been shown to match cyst activity [25].

Differential diagnosis should take into account the different CE stages, understood not simply as different types of cysts, but as different phases in the natural history of a chronic disease (Figure 7) [12].

Although there are reports of gossypibomas taken for CE because of serpiginous lines mimicking the waterlily sign of CE3a produced by the detached endocyst [11], our case has none of this and was in the end rather easily diagnosed based on US findings alone [21].

This case underlines that knowledge of the main sono-graphic features of echinococcal cyst should be part of the differential diagnosis and, in case of suspected echinococcal cyst, contacting a specialist from a referral center may shorten the time to diagnosis.

References

[1] S. Yildirim, A. Tarim, T. Z. Nursal et al., “Retained surgical sponge (gossypiboma) after intraabdominal or retroperitoneal surgery:14 cases treated at a single center,” Langenbecks Archives of Surgery, vol. 391, no. 4, pp. 390–395, 2006.

[2] P. R. Lauwers and R. H. Van Hee, “Intrapерitoneal gossypibomas: the need to count sponges,” World Journal of Surgery, vol. 24, no. 5, pp. 521–527, 2000.

[3] M. Garg and A. D. Aggarwal, “A review of medicolegal consequences of gossypiboma,” 2010.

[4] S. Akbulut, Z. Arikanoglu, Y. Yagmur, and M. Basbug, “Gossypibomas mimicking a splenic hydatid cyst and ileal tumor: a case report and literature review,” Journal of Gastrointestinal Surgery, vol. 15, no. 11, pp. 2101–2107, 2011.
[5] T. Archer and A. Macario, “The drive for operating room efficiency will increase quality of patient care,” Current Opinion in Anaesthesiology, vol. 19, no. 2, pp. 171–176, 2006.

[6] H. Alis, A. Soylu, K. Dolay, M. Kalayci, and A. Ciltas, “Surgical intervention may not always be required in gossypiboma with intraluminal migration,” World Journal of Gastroenterology, vol. 13, no. 48, pp. 6605–6607, 2007.

[7] S. Disu, A. Wijesiriwardana, H. Mukhtar, and F. Eben, “An ileal migration of a retained surgical swab (gossypiboma): a rare cause of an epigastric mass,” Journal of Obstetrics and Gynaecology, vol. 27, no. 2, pp. 212–213, 2007.

[8] A. A. Gawande, D. M. Studdert, E. J. Orav, T. A. Brennan, and M. J. Zinner, “Risk factors for retained instruments and sponges after surgery,” The New England Journal of Medicine, vol. 348, no. 3, pp. 229–235, 2003.

[9] M. K. Moslemi and M. Abedinzadeh, “Retained intraabdominal gossypiboma, five years after bilateral orchiopexy,” Case Reports in Medicine, vol. 2010, Article ID 420357, 4 pages, 2010.

[10] I. Taçyildiz and M. Aldemir, “The mistakes of surgeons: ‘Gossypiboma’,” Acta Chirurgica Belgica, vol. 104, no. 1, pp. 71–75, 2004.

[11] E. Marchiori, G. Zanetti, B. Hochhegger, and D. M. Machado, “Hydatid disease versus textiloma: a diagnostic challenge,” Thorax, vol. 66, no. 7, p. 635, 2011.

[12] E. Brunetti, H. H. Garcia, and T. Junghanss, “Cystic echinococcosis: chronic, complex, and still neglected,” PLoS Neglected Tropical Diseases, vol. 5, no. 7, Article ID e1146, 2011.

[13] A. Kalovidouris, D. Kehagias, L. Moulopoulos, A. Gouliamos, S. Pentea, and L. Vlahos, “Abdominal retained surgical sponges: CT appearance,” European Radiology, vol. 9, no. 7, pp. 1407–1410, 1999.

[14] W. Wan, T. Le, L. Riskin, and A. Macario, “Improving safety in the operating room: a systematic literature review of retained surgical sponges,” Current Opinion in Anaesthesiology, vol. 22, no. 2, pp. 207–214, 2009.

[15] J. C. Le Néel, J. B. De Cussac, B. Dupas et al., “Textilomas. A series of 25 cases and a review of the literature,” Chirurgie; Mémoires de l’Académie de Chirurgie, vol. 120, no. 5, pp. 272–277, 1994.

[16] S. Gupta and A. K. Mathur, “Spontaneous transmural migration of surgical sponge causing small intestine and large intestine obstruction,” ANZ Journal of Surgery, vol. 80, no. 10, pp. 756–757, 2010.

[17] F. H. Taylor, R. W. Zollinger, T. A. Edgerton, H. D. Harr, and V. B. Shenoy, “Intrapulmonary foreign body: sponge retained for 43 years,” Journal of Thoracic Imaging, vol. 9, no. 1, pp. 56–59, 1994.

[18] S. Jouini, R. Gourdie, K. Ayadi, M. Suleiman, and E. Wachuku, “Giant abdominal cystic textiloma mimicking hydatid cyst,” Annals of Saudi Medicine, vol. 21, no. 1-2, pp. 62–64, 2001.

[19] A. E. Duman, O. Ersoy, O. Abbasoglu et al., “Misdiagnosis of gossypiboma as hydatid cyst,” Indian Journal of Gastroenterology, vol. 30, no. 6, p. 285, 2011.

[20] W. Hosch, M. Stojkovic, T. Jänisch, G. W. Kauffmann, and T. Junghanss, “The role of calcification for staging cystic echinococcosis (CE),” European Radiology, vol. 17, no. 10, pp. 2538–2545, 2007.

[21] M. Stojkovic, K. Rosenberger, H.-U. Kauczor, T. Junghanss, and W. Hosch, “Diagnosing and staging of cystic echinococcosis: how do CT and MRI perform in comparison to ultrasound?” PLoS Neglected Tropical Diseases, vol. 6, no. 10, Article ID e1880.