



Pseudocyst of Spleen; **An Uncommon Clinical Entity, although Common Theoretically**

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Abstract

Splenic pseudocyst is not common entity in clinical practice. First case is 28 years old woman, admitted to our clinic with complaint of epigastric pain. In radiologic examinations, a cystic lesion was detected in the spleen with the size of 12 cm and performed splenectomy. Second case is 26 years old man, hospitalized after detected a mass in the spleen during performed ultrasonography due to complaint of dysuria. In further radiologic examinations revealed a cystic mass in the spleen with the size of 10 cm and splenectomy was performed. In histopathological examination, the splenic pseudocyst was detected in both patients. In this paper, we aimed to discuss differential diagnosis and treatment options of splenic cysts under the view of literature.

Key Words: Splenic cysts, pseudocyst, splenectomy

Dalağın Psödokisti

Özet

Dalak psödokistleri klinik pratikte nadir karşılaşılan lezyonlardır. Birinci olgu 28 yaşında kadın hasta, epigastrik ağrı şikayeti ile polikliniğe başvurdu. Radyolojik incelemeler sonucunda dalakta 12 cm çapında kistik lezyon saptandı ve splenektomi yapıldı. İkinci olgu 26 yaşında erkek hasta, disüri şikayeti nedeni ile yapılan üreter sistem ultrasonografisi sırasında dalakta kitle saptanması üzerine servise yatırıldı. Radyolojik incelemeler sonrası dalakta 10 cm çaplı kistik lezyon saptandı ve splenektomi yapıldı. Her iki olgunun da histopatolojik incelemesi dalak psödokisti olarak sonuçlandı. Bu yazıda, dalak kistlerinin ayırıcı tanımlarını ve tedavi seçeneklerini irdelemeyi amaçladık.

Anahtar kelimeler: Dalak kisti, psödokist, splenektomi

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INTRODUCTION

Pseudocysts of spleen are reported to be common in the literature, but in clinical practice, they are not so commonly faced. Classification is made as parasitic and non-parasitic in general, but some authors prefer to classify according to origin of the cells or the content of the cyst (Table 1) (1,2). It is important to distinguish pseudocysts from other splenic benign or malign cyts, especially from hydatid cyts in order to follow the right treatment and management options (3-6). The aim of this paper is to discuss the criteria to differentiate splenic pseudocysts from other cysts of spleen, radiologic findings of splenic cyts and treatment choices of cystic lesions of spleen according to data present in literature.

CASE 1

A 28 year-old woman presented with epigastric pain and discomfort located at her left upper abdomen. A well demarcated, immobile mass with an impression of splenomegaly and it was palpated on left upper quadrant by physical examination. The patient had a cesarean section history a year ago which has ended in intensive care unit because of renal failure due to acute tubular necrosis that has dissolved in about a week. Complete blood count, biochemical parameters and tumor marker levels were normal. A cystic mass with a diameter of 12 cm which was covering the left upper quadrant, so appearing as originating from the spleen was detected on abdominal computerized tomography (CT). A magnetic resonance imaging (MRI) was performed and a cystic mass with a diameter of 11-12 cm originating from the spleen was detected. The lesion was reported hypointense by T1-weighted images, and hyperintense by T2-weighted images (Figure 1). It was reported homogenous inside, with a little heterogenous part at a peripheral side. Operation was planned with a diagnosis of splenic pseudocyst. Laparoscopic exploration was performed at first and a cystic mass extending from the centre of the spleen towards posterior of the stomach, which was adherent to the adjacent tissues was observed. The cyst was filled with serous fluid surrounded with a thick capsule. Because of splenic hilus was not visualized as laparoscopically, operation was switched to open procedure and splenectomy was performed following a left subcostal incision. The patient was discharged postoperatively on 7th day without complication. Histopathological examination of the splenectomy

Table 1. Classification of splenic cysts according to etiology and content of the cyst.

| Class | Etiology and content |
|---------------|------------------------------|
| Congenital | Real cyst |
| Inflammatory | Pyogenic, echinoccal, fungal |
| Vascular | Peliosis |
| Posttraumatic | Hematoma, pseudocysts |
| Neoplastic | |
| Benign | Hemangioma, lymphangioma |
| Malign | Lymphoma, metastasis |

material was reported as pseudocyst originating from posterior of the spleen.

CASE 2

A 26 year-old man was admitted after detection of a cystic mass at spleen while he was having an abdominal ultrasonography for dysuria (Figure 2a). This patient had a history of hospitalization for 4 days after a traffic accident inside the car 2 years ago. Routine blood tests and tumor marker levels were normal. A well demarcated cystic mass with 8-9 cm diameter covering the center of the spleen was detected on MRI. The lesion was homogenous, not contrasting and hypointense by T1-weighted images (Figure 2b), hyperintense by T2 weighted images (Figure 2c). Operation was planned with a diagnosis of posttraumatic splenic pseudocyst. A cystic mass of 10 cm in diameter filled with serohemorrhagic fluid covering the central part of the spleen was observed during the operation and splenectomy was performed. No complication occurred and the patient was discharged on post-operatively 6th day. Histopathology of the splenectomy material was consistent with a pseudocyst originated from the centre of the spleen.

DISCUSSION

Cystic lesions of the spleen, parasitic or not, rarely cause clinical symptoms, they are usually diagnosed incidentally by abdominal imaging for another purpose. Localization, size and type of the cyst are the factors to make them symptomatic. It is accepted generally that a cyst with a diameter more than 4 cm will be symptomatic, although there is no consensus on this (7). The diameter of the cysts were over 4 cm in both of our

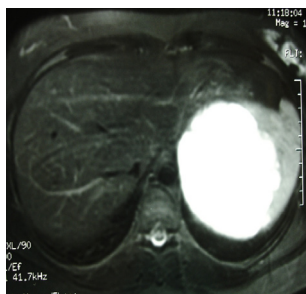


Figure 1. MRI image in transverse section : Obvious hyperintense signaling with fat suppressed T2-weighted images, a cystic mass covering the center of the spleen with a peripheral heterogeneity probably due to some debris.

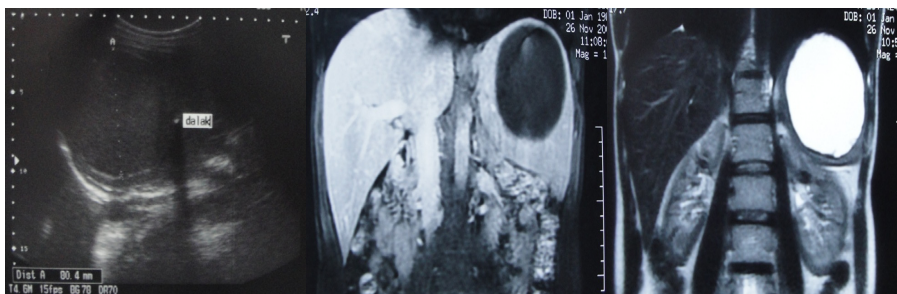


Figure 2. Ultrasonographic findings (a), cystic mass in spleen with 8 cm diameter, homogeneous internal structure and peripheral brightly echogenic foci with distal shadowing due to calcifications within the wall. MRI image in coronal section: (b) a cystic mass covering the splenic center, well defined border, homogeneous inside and obvious hypointense signaling with T1-weighted images, (c) obvious hyperintense signaling and homogenous inside with fat suppressed T2 analysis

cases. One case was totally asymptomatic and diagnosed incidentally on ultrasonography, while the other one had some non specific abdominal symptoms.

Pseudocysts of spleen have thick, fibrous walls without epithelium, which may be calcified. The most accepted etiologic theory is long term effect of posttraumatic hematomas (1, 5). But history of trauma is not always present for all cases. According to other suggested etiologic theories pseudocysts may be formed after infarction, inflammation or thrombosis of real cysts or they may be primary real cysts that have lost their epithelium (8). Debris inside the pseudocysts may cause a heterogenous image on ultrasonography and if there is calcification on the wall a peripheral shiny echogenicity with distal shading may also be an ultrasonographic finding. They appear as well demarcated, usually homogenous, not contrasting cystic masses on CT (1). MRI findings are usually similar with CT findings. The difference may be signal changes due to proteinous material and blood amount inside the cyst with T1 analysis and hyperintense signaling with T2 analysis. Cystic masses having homogenous structure on ultrasonography, not contrasting on CT which are hypointense with T1 and hyperintense with T2 on MRI were present in both of our patients.

Epithelial cysts of spleen are called real cysts that appear during the first decades of life. They may be congenital or sporadic, as well as familial. Surrounding wall

contains columnar, cuboidal, or squamous epithelium (5). CEA or CA 19-9 may be detected in the epithelium immunohistochemically as these markers may be elevated in the serum (7). They are usually asymptomatic and do rarely complicate. They appear as anechoic, well demarcated cystic masses on ultrasonography. Unilocular, not contrasting if there is no trabeculation, low density cystic mass with an ill defined wall is the image seen on CT. MRI shows hyperintense signaling with T2 which is a typical finding for all cystic lesions (1). Ages of our cases were consistent with the age of real cysts, but tumor marker levels were normal and no family history of cystic disease were present, while both of them had history of trauma on the other hand that referred the diagnosis of pseudocyst.

Abscess of spleen is diagnosed more commonly nowadays, because of increasing numbers of immunocompromised patients due to either AIDS or therapeutic agents. Abscesses located on spleen may be solitary, multiple or multilocular. Ultrasonography presents hypoechoic or anechoic mass image with an ill defined border depending on the density of purulent material inside. If air is involved, echogenicity will be increased with some posterior shading. CT image is usually a lesion with a better border that is not contrasting. Hypointense with T1, hyperintense with T2 is the image obtained on MRI. Peripheral contrasting may accompany if a capsule is existing (9). Stomach ache, fever and elevated sedi-

mentation rate are related clinical and laboratory features that help to differentiate abscesses from cysts, but abscesses still must be considered for differential diagnosis of cysts on spleen regarding their radiologic appearance.

Hydatid cyst disease should never be overlooked for differential diagnosis of cystic disease on spleen especially in endemic countries. Lesions developed by *Echinococcus granulosus* may appear solitary or multiple in number. Diagnosis is usually made preoperatively by patient history, laboratory findings or radiologic images, but sometimes these findings fail to make a diagnosis. It could be favorable if there exists calcification on cyst wall or female cysts radiologically. But the radiologic images may also be non-specific, especially Gharbi-type 1 hydatid cysts may easily be confused with other splenic cysts (6). Hydatid cyst must be considered especially when a patient demonstrates other cysts on different organs including liver simultaneously concomitant with cysts on spleen in an endemic area. The lesions of our patients were solitary and no co-existing cysts on other organs were present. Patients showed no risk factor for hydatid cyst as well as no positive serological test result. Structures like germinative membrane or female vesicles were not detected as well. Cystic lesions on spleen may also occur after lymphomas with central necrosis, metastatic tumors, huge hemangiomas, and infarctions covering large areas besides those major cystic diseases.

Surgical intervention is the traditionally suggested approach if the diameter is more than 4 cm for a cyst which makes it easier to complicate (3). Splenectomy was the operation for all types of lesions in the past, but more conservative solutions are suggested currently because the immunological functions of the spleen is better understood and it is obvious that the risk of sepsis increases after splenectomy. Total splenectomy is suggested only for huge cysts, as well as cysts covered with splenic parenchyma or cysts located on splenic hilus (7). Partial splenectomy may be the procedure for cysts which are not so large or the ones located on superior or inferior pole, and for the cases that bleeding is not a problem. This procedure allows to preserve the spleen at least partially. Since it is realized that post operative benefits are obvious, laparoscopic intervention is the suggested surgical operation for splenic cysts currently (10). Furthermore, partial splenectomy choice is recommended to be preserved for relapsing cases or suspi-

cious primary cysts; just decapsulation is reported to be enough for the rest of the cases (11). Total splenectomy was performed after the vaccinations completed via left subcostal incision since splenic hilus was not visualized as laparoscopic exploration, the cysts were large in size and located centrally in both of our cases.

Splenic cysts are rather rarely encountered in clinical practice. These cysts may be totally benign diseases or may present infectious diseases, primary malignancies and metastatic disseminations as well. The most crucial point clinically is to make the differentiation preoperatively. Therapeutic approach should allow to preserve the spleen as much as possible, but when this is not accessible, splenectomy is a treatment option especially the endemic areas of hidatyidosis and if the cyst is in very large or in hilus.

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