EPIDERMOID CYST OF THE SPLEEN: CASE REPORT

Felix Garfunkel

Greene Memorial Hospital and Wright State University School of Medicine, Xenia, Ohio

The classification, incidence, and clinical, radiologic, and scintigraphic findings of benign nonparasitic splenic cysts are presented with a case report of epidermoid cyst of the spleen.

Cysts of the spleen are rare causes of splenomegaly, and epidermoid cysts of the spleen are rare among the splenic cysts. Fifty-six cases have been published (1), 22 of which involved patients under 15 years of age.

In 1829, Andral (2) reported the first case of a splenic cyst, which he found at autopsy. Fowler's classification (3) for splenic cysts first appeared in 1940 and was modified by McClure and Altemeier (4) in 1942. Over the past 30 years case reports outlining either etiologic factors and pathogenesis or improved methods for preoperative diagnosis have appeared (5). The following is a report of an epidermoid cyst of the spleen. The classification, incidence, and clinical, radiographic, and scintigraphic findings are presented. Epidermoid masses in the left upper quadrant of the abdomen are rare and spleen scintigraphy is considered a valuable tool in the diagnosis of obscure abdominal masses (6).

LITERATURE DATA

Classification. Nonparasitic splenic cysts were classified by Fowler (7) as follows:

- I. Primary (with cellular lining)
 - A. Congenital
 - B. Traumatic
 - C. Inflammatory
 - D. Neoplastic
 - 1. Epidermoid
 - 2. Dermoid
 - 3. Lymphangioma
 - 4. Hemangioma
- II. Secondary (with no cellular lining)
 - A. Traumatic
 - B. Degenerative
 - C. Inflammatory

Splenic cysts are classed as primary or secondary depending on the presence or absence of a cellular wall lining. Secondary cysts, or pseudocysts, are considered false cysts that arise after trauma, inflammation, or vascular insult to the spleen (8).

In 1958 Martin (9) offered a simplified clinical classification system in which nonparasitic primary cysts are divided into congenital and neoplastic:

- I. Primary (true cysts)
 - A. Parasitic
 - B. Nonparasitic
 - 1. Congenital
 - 2. Neoplastic
- II. Secondary (false cysts)

This classification has been endorsed in several recent reports (10,11).

Incidence. Over two-thirds of the splenic cysts throughout the world are parasitic hydatid cysts caused by Taenia echinococcus (3), which is very rare in this country (8). Pseudocysts are four times more frequent than nonparasitic true cysts (9,12). Hemangioma is the most common primary cyst (13,14) and dermoid is the rarest (3).

Epidermoid cysts account for 10% of nonparasitic cysts (3,9,15). Such cysts have been found at ages from 6 months to 50 years; two-thirds of the reported cases occurred in the second and third decades (16). The rarity of this clinical entity is substantiated by Custer (17), who found only five epidermoid cysts of the spleen in 5,000 autopsies.

Clinical findings. Nonparasitic splenic cysts may be completely asymptomatic or may present with acute abdominal symptoms (5): either pain in the left upper quadrant of the abdomen or increasing abdominal girth (18,19). The average duration of symptoms was found by Lee and Arnspiger to be 1.49 years (20). All clinical symptoms can be explained by the displacement of surrounding struc-

Received April 30, 1975; revision accepted Sept. 26, 1975. For reprints contact: Felix Garfunkel, Greene Memorial Hospital, Xenia, Ohio 45385.



FIG. 1. Upper gastrointestinal study shows rightward displacement of stomach and duodenal loop.



FIG. 2. Anterior scintiscan shows splenomegaly and splenic defect.

tures by the enlarging splenic mass (21). Physical examination usually reveals a palpable mass in the left upper quadrant of the abdomen, generally non-tender. The sizes of the other abdominal organs must be carefully evaluated because of their significance in the differential diagnosis.

Radiographic and scintigraphic findings. If the splenic cyst is large, the main finding will be a large

space-occupying lesion in the left upper quadrant. First there will be evidence of splenomegaly. If a spleen is very large, it may not be readily recognizable on abdominal films. The left diaphragm may be elevated. The stomach, the duodenojejunal junction, the splenic flexure of the colon, and the left kidney may be displaced downward and occasionally to the right, as shown by contrast studies (19, 22, 23). McNamara et al (24) considered that displacement of the stomach to the right, rather than superiorly, anteriorly, or to the left was more consistent with a splenic mass than with pancreatic or retroperitoneal masses. Contrast studies, such as upper gastrointestinal series, barium enema, or intravenous pyelogram, confirm the presence of a mass in the left upper abdomen and may locate the mass at the spleen.

With these diagnostic procedures alone, however, a correct preoperative diagnosis of splenic cyst has been quite rare (13). Needle aspiration of the mass in the left upper abdomen and splenography have been described, but an abdominal catastrophe due to damage of a hydatid cyst still may occur with these techniques. Splenic scintigrams using ¹⁹⁸Aucolloid, ⁵¹Cr-tagged red blood cells, or ^{99m}Tc-sulfur colloid have been recommended as very valuable tools in the diagnosis of obscure upper abdominal masses (24–26). Selective abdominal arteriography is also emphasized for definitive diagnosis (13,27).

CASE REPORT

A 19-year-old girl was admitted with the finding of a large supraumbilical mass in the upper abdomen. She had first noted the mass 4 months before admission, and since then it had gradually increased in size. It first became painful 11 days before admission. The mass was movable and seemed to be on the left side. She had no other symptoms. Menarche was at age 12. There had been no previous surgery. Family history was noncontributory.

On examination she looked as if she were 7 months pregnant. She appeared slightly pale. In the markedly distended upper abdomen a bloated mass was felt, slightly to the left side. Liver size was normal. The spleen was thought to be not palpable.

The white blood count was 6,000 per mm³, hemoglobin 12.2 gm per 100 ml, hematocrit 36%, and platelet count adequate. A screening test for renal function gave normal results. Chest radiograph was normal. An intravenous pyelogram disclosed a large uncalcified mass in the left upper abdomen, extending across the midline into the right upper quadrant and measuring about 20×25 cm in diameter. The left kidney was displaced inferiorly. An upper gastrointestinal series revealed marked displacement of the



FIG. 3. Left lateral scintiscan of spleen shows defect anteriorly.

stomach and duodenojejunal junction to the right (Fig. 1). A barium enema showed downward displacement of the splenic flexure of the colon. A splenic scan with ^{99m}Tc-sulfur colloid (3 mCi) was performed; routine anterior, posterior, and left lateral views showed a large oval area, 19 cm in diameter, devoid of activity in the medial posterior aspect of the spleen. It occupied the upper two-thirds of the organ, but left a rim of normal splenic activity in the uppermost aspect (Figs. 2 and 3). This intrasplenic defect was believed to represent the above-described mass, which had very markedly enlarged and had displaced normal splenic tissue. The liver was normal in every respect. Neither arteriography nor ultrasonography was performed.

At laparotomy a very large cyst of the spleen, occupying the entire left upper abdomen, was found. Prior to removal of the spleen, 3,300 ml of serous fluid were aspirated by trocar and cannula.

The excised organ measured $26 \times 16 \times 8$ cm and weighed 1,400 gm. The capsule was smooth. The decompressed cyst measured 15×8 cm and still contained some cloudy reddish-yellow fluid. The inner surface was yellowish white with a focal trabecular pattern. The microscopic diagnosis was epidermoid cyst of spleen. The postoperative course was uneventful and the patient was discharged 8 days after the operation.

CONCLUSIONS

This case is reported because of the relative rarity of this type of cyst and because it presents such characteristic findings as young age group, female, and the absence of acute symptoms except for the spaceoccupying lesion in upper abdomen, its recent rapid growth, and vague abdominal pain. The spleen scintigram is felt to be the most contributory diagnostic tool in that it pinpoints the location of the mass, which in conjunction with the displacement of the stomach to the right, is most suggestive of a splenic mass. Although not performed in this case, an ultrasonic scan of the left upper abdomen would have been most useful. Such findings (i.e., a splenic mass with ultrasonic cystic pattern) would definitely point toward inclusion of epidermoid cyst of the spleen in the differential diagnosis.

ACKNOWLEDGMENTS

I thank Allan Ashare, Chairman of Radiological Sciences at Wright State University School of Medicine, for reviewing this paper and Harold Tharp for permission to use this case.

REFERENCES

1. BLANK E, CAMPBELL J: Epidermoid cysts of the spleen. Pediatrics 51: 75-84, 1973

2. ANDRAL G: Precis d'anatomie pathologique, Paris, Gabon, 1829, p 432

3. FOWLER RH: Hydatid cysts of spleen. Int Abstr Surg 96: 105-116, 1953

4. MCCLURE RD, ALTEMEIER WA: Cysts of the spleen. Ann Surg 116: 98-102, 1942

5. SIRINEK KR, EVANS WE: Nonparasitic splenic cysts. Am J Surg 126: 8-13, 1973

6. ANSINGH HR, STAPLETON JE: Epidermoid cyst of the spleen. Diagnosis by photoscanning. Northwest Med 66: 461-463, 1967

7. FOWLER RH: Cystic tumors of spleen. Int Abstr Surg 70: 213-223, 1940

8. QURESHI MA, HAFNER CD, DORCHAK JR: Nonparasitic cysts of the spleen. Arch Surg 89: 570-574, 1964

9. MARTIN JW: Congenital splenic cysts. Am J Surg 96: 302-308, 1958

10. DAS GUPTA T, BRASFIELD R: Splenic cysts. NY State J Med 66: 632-635, 1958

11. CLARKE JM, TOLBERT JL: Neoplastic noncancerous cystic tumors of spleen. Am J Surg 35: 488, 1969

12. BRON KM, HOFFMAN WJ: Preoperative diagnosis of splenic cysts. Arch Surg 102: 459-461, 1971

13. PINES B, RABINOVITCH J: Hemangioma of spleen. Arch Pathol 33: 487-503, 1942

14. WHITLEY RD, WINSHIP T: Splenic hemangioma with subsequent fatal hemangiosarcoma. Surgery 35: 787-792, 1954

15. BOSTICK WL, LUCIA SP: Nonparasitic noncancerous cystic tumors of spleen. Arch Pathol 47: 215-222, 1949

16. COLEMAN WO: Epidermoid cyst of spleen. Report of two cases. Am J Surg 100: 475-479, 1960

17. CUSTER RP: Practice of Pediatrics, vol 3, chap 20, Brennemann J, ed, Hagerstown, Md, Prior, 1944

18. QURESHI MA, HAFNER CD: Clinical manifestations of splenic cysts, study of 75 cases. Am J Surg 31: 605, 1965

19. HOFFMAN E: Nonparasitic splenic cysts. Am J Surg 93: 765-770, 1957

20. LEE RE, ARNSPIGER LA: Epidermoid cyst of spleen. Report of a case. Arch Surg 77: 10-12, 1958

21. MONTGOMERY AH, MCENERY ET, FRANK AA: Epidermoid cysts of spleen. Ann Surg 108: 877-884, 1938

22. ALLEN RP, CONDON VR: Epidermoid cyst of the spleen in children. Am J Roentgenol Radium Ther Nucl Med 86: 534-539, 1961

23. FORDE WJ, FINBY N: Splenic cysts. Clin Radiol 12: 49-54, 1961

24. MCNAMERA JJ, MURPHY LJ, GRISCOM NT, et al: Splenic cysts in children. Surgery 64: 487-491, 1968

25. PEARSON HA, TOULOUKIAN RJ, SPENCER RP: The binary spleen: A radioisotope scan sign of splenic pseudocyst. J Pediatr 77: 216–220, 1970

26. JOHNSON PM, HERION JC, MOORING SL: Scintillation scanning of the normal spleen, utilizing sensitized radioactive erythrocytes. *Radiology* 74: 99–101, 1960

27. POLLER S, WHOLLEY MH: Splenic cysts: Confirmation by selective visceral angiography. Am J Roentgenol Radium Ther Nucl Med 96: 418-420, 1966

THE SOCIETY OF NUCLEAR MEDICINE 23rd ANNUAL MEETING

June 8-11, 1976

Dallas Convention Center

Dallas, Texas

FIFTH CALL FOR ABSTRACTS FOR SCIENTIFIC EXHIBITS

The Scientific Exhibits Committee welcomes the submission of abstracts for the display of scientific exhibits for the 23rd Annual Meeting of the Society of Nuclear Medicine. This year there will be three categories of exhibits: 1) regular and small viewbox; 2) residents and fellows; and 3) jiffy exhibits containing materials presented in scientific papers.

Regular exhibits may be large or small, free standing or illuminated by viewbox, and offer a means whereby attendees can take their time to view the material, assimilating and digesting the information at their own pace. The special residents and fellows category is designed to encourage physicians who are training in the field of nuclear medicine to participate in the national meeting. The jiffy exhibits, whose introduction last year was so successful, offer an opportunity for those presenting scientific papers to present the salient features of their paper in exhibit format so that participants in the meeting can review the data at their leisure.

Scientific exhibits award: The Society is pleased to announce the presentation of awards in the following categories: 1) clinical nuclear medicine; 2) instruction; 3) biophysics and instrumentation; 4) residents and fellows exhibits. In each category there are gold, silver, and bronze medal awards for outstanding exhibits. Judging is based on scientific merit, originality, display format, and appearance. Judging will occur on the first full meeting day.

Abstract format: Abstracts must be submitted on a special abstract form for scientific exhibits which is available from the Society of Nuclear Medicine, 475 Park Avenue South, New York, New York 10016.

Abstract deadline—April 1, 1976. Abstract deadline for Jiffy Zibits—May 1, 1976. Send all abstract forms to:

> H. William Strauss, M.D. Division of Nuclear Medicine and Radiation Health The Johns Hopkins Medical Institutions 615 North Wolfe Street Baltimore, Maryland 21205