www.jmscr.igmpublication.org

Impact Factor 3.79 ISSN (e)-2347-176x



Giant Mesenteric Cyst, Presentation as Acute Appendicitis in Adult: A Rare Case Report

Authors

Dhukiya Ramniwas¹, Kajla Ramakishna², Dhukiya Saroj³, Sinwar Prabhu Dayal⁴ Chouhan S.P⁵

¹Senior Resident, Department of General Surgery, Sardar Patel Medical College Bikaner
²AssociateProfessor, Department of General Surgery, Sardar Patel Medical College Bikaner
³Senior Demonstrator, Department of Physiology, Sardar Patel Medical College Bikaner
⁴ Senior Resident, Department of General Surgery, Sardar Patel Medical College Bikaner
⁵Professor, Department of General Surgery, Sardar Patel Medical College Bikaner
⁶Drofessor, Department of General Surgery, Sardar Patel Medical College Bikaner
⁶Drofessor, Department of General Surgery, Sardar Patel Medical College Bikaner
⁶Drofessor, Department of General Surgery, Sardar Patel Medical College Bikaner

S/o Ladu Ram Dhukiya V.P.O.-Binasar, Tehsil and District-Churu Rajasthan 331031 India Email: dr.ramniwas1985@gmail.com

ABSTRACT

Introduction - Mesenteric cysts are uncommon benign abdominal lesions with no classical clinical features. The preoperative diagnosis requires the common imaging modalities but the final diagnosis established only during surgery or histological analysis. The treatment of choice is complete surgical excision. Case report-We report a 25-year-old male with a non-specific abdominal pain and discomfort since 15 days. USG finding suggestive of large collection in right ileac fossae and pelvis. Expletory laperotomy was done and during intraoperative a large mesenteric cyst was found. The cyst was completely removed and histology report confirmed mesenteric cyst without evidence of malignancy. Conclusion- Mesenteric cyst is rare tumor that more common in children. They may be presented various clinical symptom. Mesenteric cyst best treated as resection anastomosis

Keyword: Mesenteric cyst, Large collection, Malignancy, laparotomy, Resection anastomosis

INTRODUCTION

A mesenteric cyst is a cystic mass developing in that is observed in one in 100,000- 250,000 hospital admissions $^{[1,2]}$. The cause of a

Dhukiya Ramniwas et al JMSCR Volume 2 Issue 10 October 2014

mesenteric cyst is still not clear; nonetheless, obstruction of lymph ducts, injury of lymph ducts, degeneration of lymph nodes, proliferation of ectopic lymphoid tissues, failure of mesenteric leaves to fuse have been considered. As its etiology, congenital diseases, diverticula, surgery in the pelvic area, and pelvic inflammatory diseases have been cited $^{[3,4]}$. It is asymptomatic in most cases and is detected incidentally by using radiological diagnostic tests. Although uncommon, complications, such as infection, hemorrhage, volvulus, perforation, and ileus, have been reported [3,5,6]. The choice of treatment is a complete surgical resection with or without bowel resection. We experienced 1 patient who developed an acute abdomen induced by infected mesenteric cysts that had developed in the mesentery of the small bowel.

CASE REPORT

An 25-year-old male presented to the surgical clinic of PBM Hospital Bikaner with the history of dragging pain in the lower part of abdomen in right ileac fossa 4 month back. The Patient investigated and USG finding suggestive of appendicular abscess with lump formation. Patient treated conservatively. Patient discharge from hospital with advised interval appendectomy. He lost in follow up. Patient readmitted with pain in right ileac fossa since 15 day. The patient also have fever last 5 day, with sign of intestinal obstruction.He denied any history of trauma or previous abdominal surgery. The patient had tachycardia Pulse 110/minute, BP 116/76 mm of Hg. Her abdominal examination demonstrated slight distension with acystic and mass measuring 10x12 cm in the in right ileac fossa region. Bowel sounds were audible with exagraded frequncy. In investigations, TLC counts raised, 18000 cu/mm and RFT, LFT in normal limits and chest x-ray was normal. Ultrasonography of abdomen and pelvis suggestive of lump in right ileac fossa and large collection measuring 15x12 cm in right ileac fossa and 10x12 cm in pelvis with intervening septation no visualisation of appendix. Abdominal drain inserted in right ileac fossa in local anaesthesia.but patient condition not improved. The patient plan for exploratory laparotomy. Intraoperative a large mesenteric cyst originating from small bowel mesentery, occupying in right ileac fossa and pelvis(figure 1,2) with proximal small bowel loop dilated. Resection and anastomosis done along with removed of mesenteric cyst which was not found to be infiltrating the surrounding structures (Figure-3) Layered closure of the abdomen was done after the placement of a suction drain. The patient made uneventful recovery and was discharged home on 9th postoperative day. The histological report showed cystic wall composed of fibrocollagenous and adipose tissue with chronic inflammatory cell neutrophils and eosinophils.Intestinal surface is devoied of lining epithelium. It is made up of acute inflammatory cell consisted with mesenteric cyst lined by columnar epithelial cells (figure 4,5,6).



[Figure-1] Mesenteric cyst



[Figure-2] Mesenteric cyst



[Figure-4] Mesenteric cyst(normal intestine10x H&E)



[Figure-5] Mesenteric cyst with infiltration of chronic inflamatery cell 40x H&E



[Figure-3] Mesenteric cyst



[Figure-6] Mesenteric cyst with infiltration of chronic inflamatery cell 10x H&E

DISCUSSION

Majority of mesenteric cysts are congenital, but may be related to previous abdominal surgery, pelvic diseases, and trauma [7,8]. The most common presentation is no specific abdominal pain (55-82%), followed by the complaint of abdominal mass (54-61%), and abdominal distension [9,10,11] (17-61%). Physical examination is often unremarkable but reveals a mass in 66.8% of cases12 while the average duration of symptoms are 2-6 months (range; 12 h to 12 months)[12]. Viola et al. hypothesized that mesenteric cystic lymphangioma is an acquired anomaly due to chronic intermittent volvulus [13]. The most widely accepted classification was coined by Beahrs et al. in 1955[14].16 According to this classification, there are four types of mesenteric traumatic, cysts; developmental, infective, and neoplastic. The malignant cystic mesothelioma is the only mesenteric cystic tumor that carries malignant potential and has tendency to recur after surgical excision[15]. Diagnosis is established via one of three routes: Firstly, patients may be asymptomatic and the disease is discovered coincidentally on imaging or intraoperatively. Secondly, patients may present with non-specific symptoms prompting investigation or thirdly, approximately one third of patients develops complications and present acutely as in our case. Complications include bowel obstruction volvulas, hemorrhage, infection and rupture. Rupture of a spontaneously infected mesenteric cyst is extremely rare. Mesenteric cysts have been classified based on their histopathological findings [16], the first type consisting of cysts of lymphatic

origin also known as lymphangioma, It has been suggested that ectopic lymphatic tissue or blocked lymphatic channels may be the causative developmental abnormality. The mechanism that such can infect a cyst is unclear. The histopathology in this case did however reveal patent lymphatic channels extending to the bowel wall and therefore lymphangitis with infection from bowel organisms would seem a plausible explanation.

CONCLUSIONS

The high mortality in the treatment of this condition is due to the late recognition of the disease, usually during an emergency operation, at which time the serious complications of intestinal obstruction and peritonitis have already developed. Under such circumstances, resection of the bowel naturally carries with it a very high primary mortality. These tumors are sufficiently common to justify consideration of the condition whenever an abdominal tumor is observed. The symptoms and signs are not characteristic and the diagnosis must be made by the exclusion of the more common tumors met with in the abdomen.

REFERENCES

- Liew SC, Glenn DC, Storey DW. Mesenteric cyst. Aust N Z J Surg 1994;64:741-4.
- Sato M, Ishida H, Konno K, Komatsuda T, Konno S, Watanabe S, et al. Mesenteric cyst: sonographic findings. Abdom Imaging 2000; 25:306-10.

JMSCR Volume||2||Issue||10||Page 2571-2575||October-2014

2014

- Sardi A, Parikh KJ, Singer JA, Minken SL. Mesenteric cysts. Am Surg 1987;53:58-60.
- de Perrot M, Brundler M, Totsch M, Mentha G, Morel P. Mesenteric cysts. Toward less confusion? Dig Surg 2000;17:323-8.
- Vanek VW, Phillips AK. Retroperitoneal, mesenteric, and omental cysts. Arch Surg 1984;119:838-42.
- Tan JJ, Tan KK, Chew SP. Mesenteric cysts: an institution experience over 14 years and review of literature. World J Surg 2009;33: 1961-5.
- De Perrot M, Brundler MA, Totsch M, et al. Mesenteric cysts. Towards less confusion? Dig Surg 2000;17:323-8.
- Wykypiel HF, Margreiter R. Chylous cyst formation following laparoscopic fundoplication. W ien K1 inh Wochenschr 2007;119:729-32.
- Liew SC, Glenn DC, Storey DW. Mesenteric cyst. ANZ J Surg 1994;64:741-4.
- Shimmura H, Ueda J, Ogawa Y, et al. Total excision of mesent eri c c yst s by

laparoscipoc surgery; report of 2 cases Surg Laparosc Endosc 1997;7:173-6.

- 11. Tan JY, Tan KK, Chew SP. Mesenteric cysts: An institution experience over 14 years and review of literature. World HJ Surg 2009;33:1961-5.
- Kurtz RJ, Heimann TM, Beck AR, Holt J. Mesenteric and retroperitoneal cysts. Ann Surg 1986;203:109-12.
- 13. W eeda VB, Booij KA, Arons on DC. Mesenteric cystic lymphangioma: A congenital and an acquired anomaly? Two cases and a review of the literature. J Pediatr Surg 2008;43:1206-8.
- Beahrs OM, Judd ES Jr, Dockerty MB. Chylous cysts of the abdomen. Surg Clin North Am 1950;30:1081-96.
- 15. Mennemmeyer MS. Multicystic peritoneal m esotheliom a: a report with electronmicroscopy of a case mimicking intraabdominal cystic hygroma (lymp hangiona).Cancer 2006;44:692-8.
- 16. de Perrot M, Brundler M, Totsch M, Mentha G, Morel P. Mesenteric cysts. Toward less confusion? Dig Surg 2000;17:323-328