Case Report and Review of the Literature

Small Bowel Mesenteric Pseudocysts: Report of a Case and Review of the Literature

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ABSTRACT

Background: Mesenteric pseudocyst (MP) are rare clinical entity presenting a thick fibrotic wall with internal septae without epithelial lining cells.

Case Report: We described one case of MP of a 50-year-old woman with persistent, non-specific, and non-responding to medical therapy abdominal pain. Laboratory tests, clinical examination and the abdominal US were un-conclusive while the CT scan of the abdomen showed a cystic lesion of the jejunal mesentery. We performed a laparoscopic surgical excision of the lesion with the resolution of clinical symptoms. The lesion resulted in inflammatory mesenteric cyst at the histological examination. At 5 years CT scan follow up we did not record any recurrences.

Conclusion: MP present themselves without specific symptoms, diagnostic imaging techniques such as US or CT scan may define its features, location or size but fail in determining their benign or malign nature. In fact, even if, the majority of these lesions are benign, a rate of malign transformation of around 3% has been reported, which is why the complete surgical excision is the gold standard treatment.

Introduction

The term “mesenteric cyst” refers to a heterogenous group of cystic lesions with different etiopathogenetic roots, which appears in the abdominal cavity or in the retroperitoneum without knowledge of its abdominal organ of origin. The mesenteric cyst is a rare entity, and its reported incidence ranges from 1 per 100,000 to 1 per 250,000 hospital admission [1]. Various classifications for this kind of lesions have been proposed. The term “mesenteric pseudocyst” (MP) was used for the first time by Ros et al. in their classification of mesenteric cysts; in 2000 de Perrot et al. introduced a classification based on the histological features of the internal epithelium and consisted of 6 groups: lymphatic cyst, mesothelial cyst, enteric cyst, urogenital cyst, mature cyst teratoma and pseudocyst [2, 3]. MP might have an infectious, traumatic, or even degenerative aetiology and is similar to pancreatic pseudocyst, from a histological point of view, presenting a thick fibrotic wall with internal septae without epithelial lining cells [3]. They are sited more frequently in the mesentery of the small bowel (50-80% of cases), colon (15-30%) or in the retroperitoneum (7-20%) [4].

Iida et al. reported in 2003 that the total number of MP’s records in Japanese literature was only 15 cases [5]. Yamabuki et al. reported in 2017 that the total number of MP records in the literature was about 20 cases [6]. In our report, we present the case of a woman with a jejunal MP and abdominal pain, who underwent a laparoscopic surgical procedure to excise the lesion and a concomitant review of the relevant literature.

Case Report

A 50-year-old female patient presented to our observation for the appearance of an abdominal cystic neoformation. The patients described diffuse abdominal pain for about 2 months without the association with any other gastrointestinal or general symptoms. The US of the abdomen...
was unfortunately vitiated by the subcutaneous fat and intestinal meteorism. An injected CT scan of the abdomen was carried out showing a cystic lesion (Figure 1) with a thick wall located in the mesentery of the small bowel in the right upper abdominal quadrant, which did not take contrast and appeared dissociable from the surrounding structures. His past medical history was remarkable for essential hypertension, while his surgical one was unremarkable. He had a BMI of 30 kg/m² so the physical examination showed only a non-localized abdominal tenderness without signs of peritonitis; the laboratory tests were negative.

We performed a complete laparoscopic surgical excision of the lesion, which was located in the jejunal mesentery 30 cm distant from the Treitz ligament. The histological examination pleads in favour of the MP (Figure 2). The post-operative course was uneventful, the patient was discharged on postoperative day 2 and the 5 years CT scan follow up did not show any recurrence. Moreover, at 6 months, clinical follow up the symptoms disappeared.

**Discussion**

MPs are a rare entity, which usually occurs in the 4th decade in both sexes with equal frequency [7, 8]. Their size may vary from few to 30 centimetres; their pathogenesis remains unclear [9, 10]. Most authors support the hypothesis of a traumatic or infectious intestinal origin, in our case, both patients did not report any history of abdominal trauma or inflammatory disease. MPs are often asymptomatic and could be incidentally discovered during surgical procedure or ultrasound (US) and CT scan carried out for other reasons [6, 11].

At the physical examination, according to the patient’s habitus, we might not find any alteration, while sometimes, the lesion could disappear and reappear afterwards in a different location within the abdomen [12]. MP present non-specific symptoms such as abdominal pain (55-82%), abdominal lumps (44-61%) or might cause complications such as bleedings, volvulus or infections requiring an emergency surgical approach [4, 8, 9, 13-17]. US may be useful in the initial evaluation of the lesions providing information on its size and with high sensitivity on the presence of septation or debris inside it; nevertheless, it might fail the diagnosis if the MP is covered by intestinal loops. As a matter of fact, the T scan is more sensitive in terms of size, site and relation with neighbouring organ definition. Ayyappan et al. showed the utility of chemical shift MRI in detecting the lipid content of MP [18]. All MPs are benign lesions, except for the malignant cystic mesothelioma, an extremely rare entity [3]. However, an incidence rate of malignant mesenteric cyst of approximately 3% has been reported; therefore, given the impossibility of excluding the malignant nature with the imaging technique or laboratory tests, it is advisable to surgically excise even the smaller MP [3, 19, 20]. The complete surgical excision of MPs, both with a laparoscopic and open surgical approach, is associated with partial resection of the bowel in the event they are non-dissociable; it is the treatment of choice. The laparoscopic approach should be preferred and the conversation to an open one is rare. Chandradevon et al. recently reported a case of MP successfully treated with the robotic approach [6, 21]. From a review of the cases reported in the literature, after complete excision of the MPs recurrences have not been observed.

**Conclusion**

MPs are an extremely rare entity; given the lack of pathognomonic clinical, laboratory and imaging features, surgical excision is advisable due to the risk of complications and malignant transformation and a laparoscopic approach should be preferred.

**Author Contributions**

All authors contributed to the study; Geraci G. and Di Vita G. conceived, devised and designed the manuscript; D’Orazio B., Bonventre S. and Cudia B. collected the case and iconography; D’Orazio B. and Geraci G. wrote the manuscript and participated in the sequence alignment; Di Vita G., Sciumé C. and Martorana G. revised and approved the final manuscript and its conclusions.

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Competing Interests

None.

Conflicts of Interests

None.

Consent

An informed written consent was obtained from all individual participants included in this study.

Ethical Approval

All procedures performed in studies involving human participants were in accordance with the ethical standards of the 1964 Helsinki Declaration and its later amendments or comparable ethical standards.

Data Availability

The datasets generated during and/or analysed during the current study are available from the corresponding author on reasonable request.

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