Introduction

Appendicular tumours are found in less than 1% of appendix pathology. Mucocele of the appendix is a rare entity, with 0.2 to 0.3% incidence, which are obstructive appendicular dilatation of mucous material. Simple mucocele represents 29%, mucinous cystoadenoma contributes to 31 to 34% and cystoadenocarcinoma is even less frequent, representing 22 to 23% of the malignant appendix pathology. Although cystoadenomas are a benign lesions, any proliferation or rupture of cystoadenomas have a risk of peritoneal dissemination, reason why the complete excision without rupture is important to prevent complications [1].

Case Presentation

A 49-year-old man with abdominal pain of 6 days of evolution. He has medical history of diabetes mellitus type 2 and hypertension in treatment. He presents to the emergency department complaining of pain in right lower quadrant and reports radiation to the ipsilateral renal fossa. Ultrasonography demonstrated an oblique cystic fusiform lesion suspicious of abscess. Simple computed tomography concluded probable mesenteric cyst. An exploratory laparotomy was performed finding a 12 X 8 cm tumour lesion that protruded from the appendix. Tumour was dissected and right hemicolectomy with side-end ileocolic anastomosis was performed. The histopathology study reveals an appendiceal mucinous neoplasm of low grade. After 5 days the patient was discharged without complications. After 8 months follow up the patients is asymptomatic and without evidence of tumour activity.

Case Report

Giant Appendicular Mucinous Cystoadenoma: Case Report and Review of the Literature

Abstract

Background: Appendiceal tumours present an incidence of 0.2 to 0.3% among all appendectomies. It is a rare condition, commonly found as an incidental diagnosis by imaging studies due to his lack of symptoms; nevertheless it is associated with many complications when undiagnosed like pseudomixoma peritonei; a condition with high morbidity and mortality. This is the importance of recognize it to establish an opportune diagnose and treatment.

Case: A 49-year old man with abdominal pain of 6 days of evolution. He has medical history of diabetes mellitus 2 and hypertension in treatment. He presents to the emergency department complaining of pain in right lower quadrant and reports radiation to the ipsilateral renal fossa. Ultrasonography demonstrated an oblique cystic fusiform lesion suspicious of abscesses. Simple computed tomography concluded probable mesenteric cyst. An exploratory laparotomy was performed finding a 12 X 8 cm tumour lesion that protruded from the appendix. Tumour was dissected and a right hemicolectomy with side-end ileocolic anastomosis was performed. The histopathology study reveals an appendiceal mucinous neoplasm of low grade. After 5 days the patient was discharged without complications. After 8 months follow up the patients is asymptomatic and without evidence of tumour activity.

Conclusion: Appendiceal tumours are a rare pathology with an insidious presentation representing a difficult diagnosis; requiring high index of suspect and knowledge about the correct surgical management to obtain better outcomes.

Discussion

Appendicular tumours have been divided in three main categories:
1) adenoma, 2) mucinous neoplasms of uncertain malignant potential or low-grade mucinous neoplasm and 3) adenocarcinoma [1-3].

Mucocele of the appendix can be divided into simple mucocele, which is seen in approximately 29%, epithelial hyperplasia and mucinous cystoadenoma, 31 to 34% respectively, cystoadenocarcinoma 5% and retention cysts [3]. Appendix mucoceles are characterized by a thin walled round cystic mass, confined to the mucosa, encapsulated and calcifications areas in 55% of cases. Mucoceles smaller than 2 cm are rarely malignant, but in larger ones, >6 cm in size, cystoadenoma or cystoadenocarcinoma must be suspected, with a perforation rate of 20%. Enhancing nodules in the mucocele wall suggests diagnosis of adenocarcinoma [4].

The mean age of presentation is 49.1 years and showed a female preponderance. In 50% of cases, mucocele of the appendix is asymptomatic and picked up as an incidental diagnosis. When symptomatic, 80% is presented as abdominal pain, palpable mass, nausea, vomiting, weight loss, generalized peritonism and intestinal obstruction. The right lower abdominal pain may simulate an acute appendicitis, a mass or pseudomixoma peritonei [1,4].

When rupture occurs there is extravasation of mucin and/or neoplastic epithelium, resulting in an inflammatory reaction and fibrosis with formation of a mass localized to the right lower quadrant [3].

Ultrasound may be helpful and may show an encapsulated cystic lesion, firmly attached to the caecum and sometimes an “onion skin” like appearance. CT scan shows round low-density, thin walled encapsulated mass communicating with the cecum. Colonoscopy may show a pathognomonic “sign of the volcano” image, an erythematous soft mass with a central crater from which mucus is discharged [4,5].

Carcinoembryonic antigen (CEA), Ca 19-9 and Ca-125 are potential tumour markers in epithelial appendicular neoplasm. CEA is raised on 56.1% and Ca 19-9 in 67.1% of cases. One or any combination of them can be elevated in 60% of patients. A Ca-125 elevation can be an important survival indicator. A 3-year survival rate in patients with elevated versus non elevated Ca-125 level were 83%, versus 52% respectively [1,5]. Synchronous colon cancers have been recognized to occur with appendicular mucoceles in 19.5 to 21.4%, and these patients require full evaluation for the presence of other tumours [4,6].

It is estimated that based on imaging studies only 15 to 29% of appendicular mucinous cystoadenomas are correctly diagnosed prior to surgical intervention. Cystoadenoma of the appendix remains a diagnosis that is rarely considered prior to elective surgery [7].

Perforation occurs in approximately 20% leading to mucin distribution either locally or throughout the peritoneal cavity, knowing as pseudomixoma peritonei [1].

Complete excision of the mucocele without rupture is of paramount importance because complications with rupture increase the risk of malignancy. Simple appendectomy and resection of the mesentery is also thought to be adequate even in patients with cystadenocarcinoma without mesenteric lymph nodes or adjacent organ involvement. Whenever there is doubt, caecal resection or

![Figure 1: CT scan axial view with a tumour image of 109.7 mm of length.]

![Figure 2: CT scan coronal view showing the extension from right to left abdomen, apparently right colon origin and septum inside.]

![Figure 3: Smooth and pearly colour surface, 12 x 8 cm tumor protruding from the appendix.]

![Figure 4: 15 x 7.5 x 6 cm cecal appendix with abundant mucus inside and fibrous septum that divide his light.]

right hemicolectomy is advised. Right hemicolectomy is indicated for: invasive adenocarcinoma, tumors near the cecum with lesions larger than 2 cm, mucin production, invasion of the lymphatics, if a non-mucinous histological type is identified by hystopathological examination and serosa, or mesoappendix or cellular pleomorphism with a high mitotic rate. However, if invasive cystoadenocarcinoma has already seeded into peritoneum at diagnosis time, there are no better outcomes or benefit in patient mortality with hemicolectomy compared to simple appendicectomy [1,4,8,9].

Low-grade adenomas need a completely atraumatic removal to minimize the risk of peritoneal seeding. If exploration reveals a ruptured appendicular mucocele, the primary resection should be accompanied by removal of all gross implants, increasing 5-year survival rates to 91-100%. This 5-year survival rate is markedly diminished due to complications of pseudomyxoma peritonei, decreasing to 32 to 58%. A survival advantage was shown for patients treated by appendectomy alone (median survival 18 years) compared with those who underwent right hemicolectomy (median survival 10 years) if this patients did not have the previous mentioned hemicolectomy indications [1,8].

Aggressive treatment strategy involving cyto-reductive surgery and perioperative intra-peritoneal chemotherapy showed no benefit in the group of patients who had right hemicolectomy compared with appendectomy alone. Patients who had right hemicolectomy in the absence of intra-peritoneal chemotherapy had a survival disadvantage [9].

Conclusions

Appendix tumors are a rare pathology with an insidious presentation representing a diagnostic challenge inclusive with appropriate image studies like CT scan, making difficult surgery planning and requiring high index of suspect and knowledge about the best surgical management to obtain good outcomes without complications or morbidities associated with the different treatment options. Although many cystoadenomas are benign, any proliferation or rupture of cystoadenomas have a risk of peritoneal dissemination, reason why the complete excision without rupture is important to prevent complications and in cases of laparoscopic approach the risk of rupture must be valued.

Informed Consent

Informed consent for publication of this case was signed.

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References