CASE REPORT

A weeping umbilical hernia: bilateral ovarian mucinous cystadenoma with disseminated peritoneal adenomucinosis

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SUMMARY
A young woman was referred to us for the management of an umbilical hernia with macerated overlying skin through which massive ascites was leaking. On examination we found a jelly-like substance seeping out of the opening. A contrast-enhanced CT scan of the abdomen revealed bilateral ovarian cystadenocarcinoma with pseudomyxoma peritonei. The patient underwent cytoreductive surgery along with hysterectomy and bilateral oophorectomy followed by hyperthermic intraperitoneal chemotherapy with mitomycin C. Histopathological examination showed bilateral borderline cystadenoma with peritoneal adenomucinosis. Follow-up at 1 year showed no signs of recurrence.

BACKGROUND
Pseudomyxoma peritonei is rarely encountered in day-to-day practice and is poorly understood. Most cases usually present as either an appendicular or ovarian pathology. Presentation as umbilical and bilateral inguinal hernias is rare.

CASE PRESENTATION
A young female patient aged 26 years from a very poor socio-economic background was referred to us for investigation and management of massive ascites which had eroded through a pre-existing umbilical hernia. A thorough history revealed a gradually distending abdomen initially, followed by a rapid increase in the size of the abdomen over the previous 6 months. The patient only visited the hospital when the umbilical hernia ruptured and started leaking. She also complained of a constant dull ache in her abdomen which had never upset her enough to visit hospital. She had no other abdominal complaints other than early satiety for the previous 6 months. She had achieved menarche at the age of 15 years and her menstrual cycles were normal in terms of duration and flow. She was married for almost a year and had no children. There were no other gynaecological complaints. Examination of the abdomen was remarkable as it was hugely distended and fluid thrill. Bowel sounds were normal. There was an umbilical hernia which was perforated and weeping constantly. On close examination we found a jelly-like substance protruding from the perforation site. The patient also had bilateral inguinal herniae (figure 1). A working diagnosis of pseudomyxoma peritonei was made.

INVESTIGATIONS
The patient’s haemoglobin was 6 g/dL and total leukocyte count was 5600/mm3 of blood with a lymphocyte count of 58%. Serum creatinine and blood urea were 1.7 and 78 mg/dL, respectively. The urine examination was unremarkable and a chest X-ray did not reveal any abnormality except a markedly elevated diaphragm, which was consistent with the clinical examination.

Ultrasonography (figure 2) and a contrast-enhanced CT scan (figure 3A, B) of the abdomen revealed bilateral adnexal multiloculated cystic space-occupying lesions, a thickened omentum with multiple small cystic lesions and multiloculated intraperitoneal collections causing compression of the liver, pancreas and spleen resulting in a scalloped margin along with the umbilical hernia. Features were suggestive of bilateral ovarian cystadenoma or cystadenocarcinoma with pseudomyxoma peritonei.

TREATMENT
The patient was supported with packed cell transfusions and parenteral nutrition, before cytoreductive surgery was carried out. Intraoperative findings included cysts of varying sizes filled with mucoid and gelatinous material filling the entire peritoneal

Figure 1 Pseudomyxoma peritonei with umbilical hernia with leaking gelatin-like fluid. Note the inguinal herniae.
cavity (figure 4A, B). Both ovaries were enlarged and cystic, and filled with similar material (figure 5). However, the most interesting finding was that the appendix could not be located. Surgery consisted of total omentectomy, right and left parietal peritonectomy, pelvic peritonectomy, hysterectomy and bilateral salpingo-oophorectomy, lesser omentectomy and stripping of the visceral peritoneum. As much diseased tissue as possible was removed. This surgery was followed immediately by hyperthermic (42.5°C) infusion of mitomycin C, which was followed by 5-fluorouracil infusion for 5 days. The patient was discharged after 10 days of uneventful post-operative recovery.

OUTCOME AND FOLLOW-UP
Histopathological examination of both ovaries showed multiple cysts lined with mucin-secreting columnar cells and at places stratification (two to three layers) and mild nuclear atypia suggesting borderline cystadenomas involving both ovaries (figure 6).

Random sections from the omental mass showed a predominance of acellular mucinous material and strips of epithelium composed of mucin-secreting columnar cells with stratification, mild nuclear atypia and intracellular mucin in some places. Sections from the omphalocele showed normal-looking stratified squamous epithelium. The subepithelial zone showed fibrocollagenous tissue admixed with mucinous material and strips of epithelium lined by mucinous cells.

DISCUSSION
Pseudomyxoma peritonei, a term first used by Werth1 that actually means false tumour of the peritoneum, usually has its origins in an appendiceal adenocarcinoma or ovarian...
cystadenoma, although other sites have also been described. It is a progressive disease and mostly affects women in their fifth to sixth decades. The earlier suggestion of a common ovarian origin is now debated and it has been proposed that women may have synchronous involvement of the appendix and ovary, with ovaries being secondarily involved following appendix rupture due to an enlarging mucinous adenocarcinoma. Clinical presentation varies and usually starts with non-specific symptoms which may include unexplained weight loss quickly followed by abdominal distension and pain. The most common presentation which prompts patient to present to hospital is features of appendicitis (27%) followed by increased abdominal girth (23%). Presentation as an ovarian mass (20%) is the third most common followed by hernia (14%). Various other rare presentation have also been reported including recurrent incisional hernia.

A classification was proposed on the basis of histopathology, and grouped patients into disseminated peritoneal adenomucinosis (DPAM), peritoneal mucinous carcinomatosis (PMCA) and PMCA with intermediate or discordant features (PMCA-I/D). DPAM has a far better prognosis than the other two conditions. Various prognostic factors were proposed including the presence of epithelial cells, but it was later found that virtually all cases of pseudomyxoma peritonei demonstrate some epithelial cells. Prognosis depends on the nature of the primary tumour and histological appearance. Completeness of cytoreduction therapy is a strong determinant of prognosis which is more likely achievable in DPAM.

The treatment strategy has moved from multiple surgeries aimed at debulking and palliation to a more aggressive approach involving peritonectomy and perioperative chemotherapy intended for cure. The technique of peritonectomy involves greater omentectomy and right parietal peritonectomy with or without right colon resection, pelvic peritonectomy with or without sigmoid colon resection as well as hysterectomy and bilateral salpingo-oophorectomy, lesser omentectomy and dissection of the hepato-duodenal ligament with or without antrectomy and cholecystectomy, right upper quadrant peritonectomy and Glissonian capsule resection, left upper quadrant peritonectomy and left parietal peritonectomy with or without splenectomy and other intestinal resection and/or abdominal wall resection. Intrapерitoneal chemotherapy is indicated to eradicate microscopic disease. Intrapерitoneal chemotherapy is combined with hyperthermia (hyperthermic intraperitoneal chemotherapy, HIPEC). The intraperitoneal temperature is maintained at 42.5°C. Commonly used drugs are cisplatin, mitomycin C, doxorubicin and 5-fluorouracil. In earlier series, the recurrence rate was reported to be high but this decreased considerably after the introduction of heated chemotherapy, and prognosis has improved over the years to reach a 5-year survival rate of more than 80%.

**Learning points**

▸ Pseudomyxoma peritonei is a relatively uncommon presentation of appendiceal adenocarcinoma or ovarian cystadenomia.

▸ Presentation as hernia is even more rare and can include virtually any type of ventral hernia.

▸ Unexplained weight loss followed by massive abdominal distension must be thoroughly investigated.

▸ Early detection followed by combined modality treatment with cytoreductive surgery and hyperthermic intraperitoneal chemotherapy can help achieve 5-year survival.

**Contributors** KN and VKS were clinically associated with the patient and performed the surgery. BKS carried out the radiological investigation and RL performed the histopathological examination. All the authors actively participated in drafting the manuscript.

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REFERENCES