Idiopathic sclerosing encapsulating peritonitis; rare cause of intestinal obstruction
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INTRODUCTION
Sclerosing encapsulating peritonitis (SEP) is an uncommon peritoneal condition that is characterized by the encasement of the small bowel by a fibro-collagenic membrane leading to clustering of the bowel, and thus may rarely cause intestinal obstruction. Only few cases of this disease have been reported in world literature. The etiology of this condition is unknown, and it is usually diagnosed incidentally on exploration. Surgery (membrane dissection and extensive adhesiolysis) is the treatment of choice. We report a case of SEP in a 38-year-old male patient.

CASE PRESENTATION
A 38-year-old male presented in the emergency room with two-day history of abdominal pain, vomiting and abdominal distension. The abdominal radiograph showed multiple air-fluid levels in the central abdomen. Ultrasonography of the abdomen revealed clumping of small bowel loops covered by an encapsulating membrane (Figure 1-A). The patient gave a history of a similar episode a year back which subsided without treatment in two days. He was explored for acute intestinal obstruction. Intra-operatively, complete encasement of the small bowel loops and omentum by a membrane with loose inter-bowel adhesions was found (Figure 1-B). Using blunt and sharp dissection, the adherent membrane overlying the small bowel loops was stripped off and adhesiolysis of bowel was performed. Histology of the membrane showed fibro-collagenous tissue with chronic inflammatory reaction. The patient has been followed for ten months post-operatively and is doing well.

DISCUSSION
Sclerosing encapsulating peritonitis (SEP), also known in the literature as abdominal cocoon, is an uncommon peritoneal condition that is characterized by the encasement of the small bowel by a fibro-collagenic membrane leading to clustering of the bowel, and thus may rarely cause intestinal obstruction. Only few cases of this disease have been reported in world literature. The disease primarily involves small bowel, but can extend to involve other organs like the large intestine, liver and stomach. This condition...
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can be idiopathic or secondary (most importantly and most frequently due to chronic ambulatory peritoneal dialysis). The idiopathic form is commoner and was first described by Foo et al. Patients with abdominal cocoon are generally adolescent girls. Only few idiopathic cases in males have been reported. The condition has been found to be more common in tropical and subtropical countries of the world, especially China, Malaysia, Singapore, Pakistan, India, Nigeria, Kenya, Saudi Arabia, Israel, and South Africa. The typical age range is between 12 and 18 years; however, patient ages range from 6 to 65 years in the literature.

Although the etiology of abdominal cocoon is unknown, subclinical primary viral peritonitis, as an immunological reaction to gynecological infections, or due to retrograde menstruation, and in collagen tissue diseases like scleroderma, have been suggested. Multiple etiologies for abdominal cocoon have been established, although the underlying pathogenesis is not fully understood. It may occur as a serious complication of chronic ambulatory peritoneal dialysis. It has also been described in association with prolonged practolol therapy, sarcoidosis, systemic lupus erythematosus, indwelling abdominal catheters (specifically ventriculoperitoneal and peritoneovenous shunts), orthotopic liver transplantation, liver cirrhosis, intraperitoneal instillation of drugs, leiomyomata of the uterus, endometriotic cyst or tumours of the ovary, recurrent peritonitis and tuberculous pelvic inflammatory disease.

It can rarely occur in the absence of medical or surgical risk factors, as in this case. This condition presents with recurrent episodes of acute or sub-acute small intestinal obstruction that is seldom complete, attacks of colicky abdominal pain, weight loss, nausea and anorexia, and at times with a palpable abdominal mass. Early clinical features of sclerosing peritonitis are generally nonspecific and are frequently not recognized until the patient develops partial or complete small bowel obstruction.

The condition is usually diagnosed incidentally on exploration, as in our case, although a preoperative diagnosis is feasible by a combination of barium follow-through which shows a concertina pattern or cauliflower sign along with delayed transit of contrast medium, and computed tomography of the abdomen. However, preoperative diagnosis is difficult and requires a high index of clinical suspicion. Pre-operative CT findings may include clumping of small bowel loops in the centre of the abdomen encased by a soft-tissue density mantle, peritoneal thickening, calcification, peritoneal enhancement, small bowel tethering, and loculated fluid collections.

Surgery (membrane dissection and extensive adhesiolysis) is the treatment of choice, and there is usually no need for bowel loop resection, especially when a preoperative diagnosis is feasible. Resection of the bowel is unnecessary and it increases morbidity and mortality. Resection is indicated only if the bowel is non-viable. Careful dissection and excision of the thick membrane to release the small intestine leads to recovery. An excellent long-term postoperative prognosis is most of the times guaranteed.

CONCLUSION

Idiopathic sclerosing encapsulating peritonitis, also called abdominal cocoon, is a rare cause of intestinal obstruction in adults.

References

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