INTRODUCTION

Abdominal cocoon is a rare disease which causes small-bowel obstruction that is characterized by a thick, fibrotic and cocoon-like membrane encapsulating total or partially the small bowel.\(^1,2,3\) Its etiology remains unknown.\(^4,5\) Surgical management is the treatment of choice\(^1\). In this report, it is described a young patient with abdominal cocoon who presented clinical signs and paraclinical findings of intestinal obstruction.

CASE REPORT

In July 2010 a 25 year-old man was admitted with 2-month history of intermittent colicky abdominal pain and bilious vomiting and abdominal distension. He also complained from chronic constipation, anorexia and 14 kg weight loss during the previous two months. He had no surgical or other medical history except using of a body building supplement drugs which contain creatin.

On physical examination he was anxious. Vital signs were normal (blood pressure 110/75, pulse rate 85/min, temperature 37.3° C). Abdominal distention with hyperactive bowel sounds and mild tenderness in the whole of the abdomen especially in left upper quadrant were detected at physical examination. There were no abdominal scars, palpable abdominal mass or organomegaly. Digital rectal examination was normal. Laboratory blood analysis ascertained a total leukocyte count 8500 cell/ml, hemoglobin 16 mg/dl, normal serum chemistry and normal urine analysis. Up right plain abdominal X-ray showed multiple air-fluid levels without free gas under the diaphragm. The barium X-ray showed jejunal and ileal loops dilatation with the contrast medium passing slowly (Figure 1). Contrast-enhanced abdominal computed tomography showed small bowel loops and dilatation (Figure 2).

Due to signs of mechanical small bowel obstruction an emergency laparotomy was performed through a midline incision. During surgery the small bowel dilatation was observed; its mesentery was edematous and the whole small bowel covered by a dense whitish
membrane which gave the appearance of a cocoon (Figure 3). Careful sharp and blunt dissection with excision of the membrane enveloping the small bowel was performed and the whole small intestine was freed and followed from duodenojejunal angle to ileocecal junction. The histopathology of the peeled off membrane revealed a chronic non-specific inflammatory reaction with proliferation of fibroconnective tissue.

![Intraoperative photograph showing small intestine encapsulated in white, thickened cocoon-like membrane](image)

The patient was discharged from the hospital on the 8th postoperative day. After 24 months regular follow-up he was in satisfactory condition and the abdominal plain showed normal appearance.

**DISCUSSION**

Sclerosing encapsulating peritonitis can be classified as idiopathic and secondary. The idiopathic class or abdominal cocoon was first described by Foo et al in 1978. The idiopathic type primarily affects young females from tropical and subtropical countries. The cause of this rare condition is unknown although a number of hypotheses have been mentioned. Abdominal cocoon is characterized by a thick shiny fibrotic membrane, with total or partial encasement of the small intestine.

In clinical presentation, abdominal cocoon is presented by recurrent attacks of acute or subacute small bowel obstruction, nausea, weight loss and anorexia and sometimes with palpable abdominal mass. Most patients are diagnosed incidentally at laparotomy and preoperative diagnosis requires a high index of clinical suspicion.

This patient had used some supplements for body buildings such as creatin for four years. No reference was found in the literature any association between medication given and abdominal cocoon.

WBC count, C-reactive protein level, hypoalbuminemia and anemia are common findings in SPE cases but no significant sign was observed in this case. Imaging has an important role in the diagnosis of abdominal cocoon. Barium X-ray and contrast CT are useful for the diagnosis of abdominal cocoon preoperatively. Dilated bowel loops with multiple air-fluid level may be shown in conventional abdominal X-ray but they are nonspecific. The classic findings of abdominal CT consist of small bowel loops congregation in the center of abdomen with a non-enhancement fibrous membrane surrounding the bowel loops that is best visualized on computed axial tomography scan. Surgical dissection and excision of the membrane and adhesiolysis remain the cornerstone in the treatment of abdominal cocoon. In reported case with an insignificant history, a dilated small bowel with an edematous mesentery and a dense whitish membrane coverage on bowel observed in operation which advised idiopathic sclerosing peritonitis. Because of increasing morbidity and mortality following bowel resection, if it is nonviable, resection of the bowel is indicated. In this case an emergency laparotomy was performed and an edematous mesentery covered by a dense whitish membrane was observed and incised carefully, separated from the intestinal serosa by sharp and blunt dissection.

In general, prognosis of abdominal cocoon after surgery is satisfactory and excision of the thick membrane and release of the small intestine leads to complete recovery.

In conclusion sclerosing encapsulating peritonitis is not common and definite preoperative diagnosis is difficult. The recurrent episodes of small bowel obstruction combined with relevant preoperative findings at imaging studies and lack of other causes can facilitate preoperative diagnosis of abdominal cocoon.

**REFERENCES**