ASCARIS INFECTION, A RARE CAUSE OF NECROTIZING PANCREATITIS IN A CHILD

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CASE REPORT

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INTRODUCTION

Acute pancreatitis is an inflammatory illness of variable severity, characterized by acute inflammation of the pancreatic parenchyma and peripancreatic tissues and it is characterized by epigastric pain, nausea and vomiting. Recent studies indicate that prevalence is more in children of acute pancreatitis. In the majority of cases, acute pancreatitis resolves with bowel rest and does not result in severe complications in children. It can rarely be complicated by the development of necrosis.

Acute necrotizing pancreatitis is a serious condition associated with high morbidity and mortality rates in adults. It is well described in adult but the literature is very limited with regards to pediatrics population. It has been estimated that necrotizing pancreatitis occurs in less than 1% of children with acute pancreatitis. From five large pediatric series in the United States, pancreatic necrosis was observed in only 3 out of 1014 children. There are a few case reports of necrotizing pancreatitis in children, however the data are limited regarding etiology, course, and outcome.

CASE REPORT

A 5-year-old boy presented to the emergency department of National Medical College and Teaching Hospital with history of upper abdominal pain and multiple episodes of vomiting for 1 day. The pain was sudden in onset, severe, with no known aggravating or relieving factors. The vomitus was non bilious. There was history of non-passage of stool for the same duration. On examination, the patient was in a state of shock. Abdominal examination revealed fullness of the epigastrium, generalized tenderness, absent bowel sounds and an empty rectum. Following resuscitation in the emergency department, the child was shifted to ICU for further care. His CBC showed a total count of 18,500 cells/mm³ with neutrophil of 85%. Liver Function Test showed total bilirubin is 2.30 mg/dl with direct bilirubin of 1.08 mg/dl, alkaline phosphate 490 IU/L, SGOT (146 IU/L), SGPT (87 IU/L). Serum amylase was 1700 IU/L. Renal Function Test were normal. X-ray of abdomen revealed few dilated loops of intestine. USG of abdomen revealed gallbladder and CBD ascariasis with acute pancreatitis and complex ascites. A CT Scan was planned after the stabilization of the child.

On 3rd day of admission, the patient developed features of generalized peritonitis and exploratory laparotomy was planned.
Upon entering the peritoneal cavity, about 500ml of reddish fluid was encountered and was drained. There was multiple calcification in the omentum and mesentry (Figure 1).

Figure 1: Calcification of mesenteric and omentum

Gall bladder and CBD were free of worms but few worms were noticed in the proximal jejunum that were milked into the large intestine. Upon entering the lesser sac, a necrotic mass of 4x4cm was seen along the stomach bed ensuring a mass effect on the stomach & transverse colon (Figure 2). The mass was arising from the tail of pancreas. The Necrosectomy was done and drains were placed in the lesser sac, retroperitoneal space and in the pelvis after thorough peritoneal lavage.

Figure 2: Necrotizing mass between transverse colon and stomach

Post-operatively, the child was managed in surgical ICU. Inj. octreotide along with broad spectrum antibiotic were started on 3rd postoperative day. Serum amylase was repeated on 5th POD and it was within normal limits (80IU/L). The drain from lesser sac initially put out 100ml/day of hemorrhagic fluid, but the output tapered down to 5ml/day of serous for two days prior to its removal on 10th days. On 10th post-operative day sips were allowed and on 12th day liquid diet was started. On 14th post-operative day soft to normal diet was allowed and the patient was discharged on 15th POD. His biopsy report of necrotic tissue shows feature of necrotizing pancreatitis. The patient was doing well on his follow up visit.

DISCUSSION

Necrotizing Pancreatitis, a severe form of acute pancreatitis is very rare in pediatric population. It has a variable etiology in children. Various literature have shown that necrotizing pancreatitis in children can be caused by use of valproate, M. pneumoniae and secondary to Crohn’s disease. Although Ascaris infection may be a cause of acute pancreatitis, progression to necrotizing Pancreatitis as result of it is rarely seen, as was noticed in our case.

Ascaris infection occurs in about 25% of world population and most infection are asymptomatic. A wide range of symptoms including fever, nausea, vomiting, diarrhea, pain abdomen may be present when patient is symptomatic. About 500 cases of hepatobiliary and pancreatic disease due to ascariasis in adult have been reported by Khurro et al. Pancreatitis due to ascariasis may be caused by the obstruction of papilla of vater, invasion of CBD, invasion of pancreatic duct USG, CT or MRCP, ERCP along with relevant lab investigation may help in the diagnosis of pancreatitis. Most case can be managed conservatively with antispasmodics, analgesics, IV fluids, bowel rest and anthelminthic therapy. Rarely endoscopic intervention may be required. In our case since the child has developed feature of peritonitis, we choose for surgical treatment after which the child was successfully managed.

In summary, we reported a case of necrotizing pancreatitis with peritonitis due to infestations with ascaris, which is rare cause in young child. The child was surgically managed and recovered completely.

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