A Case Report: Neonatal Sigmoid Volvulus

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Abstract: Sigmoid volvulus (SV) is an extremely rare cause of bowel obstruction in the new born period. We report a neonatal case of SV in an one month old girl presented with evidence of small bowel obstruction. At laparotomy, the classical findings of a SV was observed without gangrene of the Gut. The operative procedure consisted of simple derotation followed by sigmoidopexy.

Keywords: Bowel obstruction, new borne, sigmoid volvulus (sv).

INTRODUCTION
Sigmoid Volvulus is the most common large bowel volvulus in adults. However colon volvulus is uncommon in children[1]. SV is a disease of the elderly, often in those who are institutionalized and debilitated with neurologic and psychiatric condition[2]. The first report of SV in a 14 days old boy was published in 1961 where as the youngest patient reported is a 1 day old boy with anal stenosis[3,4]. SV is rarely considered in the differential diagnosis of abdominal pain in neonates and children and this could be responsible for the devastating results. We report a neonate (01 month old) in which acute SV occurred and was successfully managed surgically by derotation and fixing.

CASE REPORT
A one month old female infant presented with pain abdomen (crying and denying food) absolute constipation of one week duration. Vomiting was present for 03 days. Abdominal distention was present for 02 days duration. The laboratory data was unremarkable. Plain abdominal X ray showed distended bowel loops and air fluid level in intestine.[Fig-1] the baby was resuscitated by nasogastric tube, intravenous fluids and broad spectrum antibiotics. She was taken up for emergency surgery for bowel obstruction. Abdominal exploration revealed clock wise SV. Pathological finding consisted of a redundant sigmoid loop rotated around its narrow, elongated mesentery. There was no gangrene. So derotation was done with loop fixation. Recovery was uneventful. The patient was discharged from the hospital after complete oral feeding was resumed. She is doing well for last 06 months of post op follow up.

DISCUSSION
SV is a rare cause of bowel obstruction in new borne and children. There is strong male preponderance (Male : Female = 35 : 1) with wide geographic variation[5]. Our patient was a female. The presentation can range from acute to recurrent abdominal pain that is often
relieved by passage of stool or flatus. The diagnosis is usually missed or delayed with devastating consequences [6]. Our patient presented with abdominal pain and diagnosis was quick with good result after surgery. The possibility of SV should be suspected in presence of abdominal pain, constipation, nausea, vomiting with abdominal distention, tenderness and mass. Signs of peritonitis and fever suggests worst condition and perforation. Bowel peristalsis is compromised and rectum is empty [7]. The etiology of SV is different in children and adults with complication being the most common in elderly [8]. Although the etiology in children is not completely understood, the hypothesis is the presence of congenital elongation of the sigmoid colon with pathological long colonic mesentery in association with narrow base or lack of fixation of the part of colon [9]. Constipation can be considered as a cause with progressive colonic redundancy or a result. However in our case either long elongation of sigmoid colon or history of constipation was cause of volvulus.

The diagnosis is often difficult considering history, physical examination and plain abdominal radiograph. The gas pattern on X-ray is nonspecific due to absence of single U-shaped sigmoid colon loop as in adult and the inconsistent presence of the “coffee – bean” sign. The diagnosis is possible by CT scan by whirl pattern caused by dilated sigmoid colon and its meso colon and vessels and a bird peak appearance of the afferent and efferent colonic segment. However, the classic imaging are not uniformly seen. In our case, CT scan was not possible, the reaction time was less and the poor general condition of the patient forced us for emergency laparotomy.

The management of SV remains controversial, partly due to its rare occurrence. Barium enema has both diagnostic and therapeutic importance but can be complicated by perforation and shouldn’t be attempted in any patient with possible peritonitis [2]. We did not do this investigation procedure as we took up the case for surgery immediately. Reduction by endoscopy and decompression by rectal tube are other means of non operative procedure, but also carry the risk of perforation. Furthermore, non operative procedures have a high recurrence (35%) and not definitive [10].

The neonate reported here was in very poor general condition with clinical evidence of mechanical obstruction of gut. Hence, taken up for early surgery. Non resection approach with de rotation gave us a good prognosis. A volvulus preventive procedure, like sigmoidopexy was added to its good prognostic factor [11].

The overall mortality rate for SV is 6% while operative and neonatal mortality has been reported as 8.1% and 14% respectively [5]. The most common cause of death in SV is sepsis. Other causes include pneumonia, intra cranial haemorrhage, malnutrition, renal failure or hepatic failure. However, our patient is doing well for last one year of follow up.

CONCLUSION

In our experience and in accordance to literature review, children with SV do well in experienced hands of surgeon’s if operated early (prior to ischemia) in absence of co morbidities.

REFERENCES