Short and Hypertrophic Ligament of Treitz: A Rare Cause of Superior Mesenteric Artery Syndrome

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ABSTRACT
Superior mesenteric artery syndrome (SMAS) is a rare form of upper intestinal obstruction in which the third part of the duodenum is compressed between the superior mesenteric artery and the aorta, secondary to any condition decreasing the angle between these two arteries. We recently cared for a young male who came with features of proximal small bowel obstruction. On investigation, there was extrinsic duodenal obstruction. Exploratory laparotomy was done which revealed a short and hypertrophic ligament of treitz leading to compression of 3rd part of duodenum. Release of the ligament with duodenojejunostomy was done. Postoperatively, patient recovered well. This case report highlights the occurrence and importance of hypertrophic and contracted ligament of treitz as a rare cause of SMAS.

CASE REPORT
A 20-year-old thin built male was admitted with six months history of epigastric fullness, recurrent abdominal pain associated with vomiting which were improved with lying prone. He had a history of multiple admissions for suspected intestinal obstruction, wherein he was managed conservatively. On general examination, there was pallor, tachycardia and pedal oedema and BMI of 10.8 kg/m².

On abdominal examination, there were visible peristalsis in upper abdomen and hyper peristaltic bowel sounds. However, there was no sign of peritonitis. The patient was further investigated and put on parenteral nutrition and expectant management meanwhile. Haematological investigations revealed anaemia, hypoalbuminemia and hypokalemia. Ultrasonographic evaluation of the abdomen depicted duodenum to be grossly dilated with collapsed small bowel loops. Upper GI endoscopy showed dilated stomach and duodenum with no obvious intraluminal pathology. On barium meal study, stomach and duodenum up to its 3rd part were grossly dilated with abrupt cutoff in 3rd part [Table/Fig-1a]. There was no passage of contrast on films taken after one and half hour [Table/Fig-1b]. A provisional diagnosis of superior mesenteric artery syndrome was made and patient was taken up for exploratory laparotomy. Intraoperative findings confirmed dilated stomach and duodenum with a narrow aorto-mesenteric angle due to short and hypertrophic ligament of Treitz [Table/Fig-2]. Short and hypertrophic ligament of Treitz was divided and duodenojejunostomy was done. In postoperative period, parenteral nutrition was continued for a week and the patient had an uneventful recovery.

DISCUSSION
SMA syndrome was first described by Von Rokitanski in 1861 [1]. The incidence of this condition varies from 0.013-0.3% in barium series of upper GI tract. In human beings, due to the erect posture, the superior mesenteric artery leaves the aorta at an acute downward angle. It is through this vascular angle, between the aorta and the SMA that the third part of the duodenum passes and is thus vulnerable to becoming pinched in between the SMA anteriorly and the aorta and vertebral column posteriorly [2]. The aorto-mesenteric angle and aorto-mesenteric distance are reduced in SMA syndrome with values of 6 – 15 degree (normal 28-65 degree) and 2-8 mm (normal 10-34mm) respectively [3]. The common causes include exaggerated lumbar lordosis, external compression by a body cast used for treatment of vertebral fracture or lumbar lordosis [3]. However, a rare aetiology is a short and hypertrophic ligament of treitz, which brings the duodenum into the vascular angle between the SMA and the aorta [4].

Patients usually present acutely with features of duodenal obstruction, with symptoms getting relieved on lying prone. Plain abdominal X-ray demonstrates gastric dilatation. Endoscopic examination usually does not indicate the diagnosis. On barium meal, a positive diagnosis of SMAS can be made in the presence...
CONCLUSION

SMA syndrome is an uncommon cause of duodenal obstruction that should be considered once luminal pathologies have been ruled out by endoscopy. Short and hypertrophic ligament of Treitz is one of the rare causes of SMAS. Surgery offers a safe and complete cure for the same.

REFERENCES