A Case Report on Intussusception Due to Vanek’s Tumour

Dr. Ashutosh Talwar1, Dr. Nitin Nagpal2, Dr. Sarita Nibhoria3, Dr. Rahul Jain4, Dr. Anshul5

1Assistant Professor Surgery, 2Professor & Head Surgery, 3Professor & Head Pathology, 4Postgraduate Resident Surgery, 5Postgraduate Resident Surgery, GGS Medical College, Faridkot 151202 Punjab, India

DOI: 10.36347/sasjs.2020.v06i01.008 | Received: 20.01.2020 | Accepted: 27.01.2020 | Published: 30.01.2020

*Corresponding author: Dr. Ashutosh Talwar

Abstract

Inflammatory fibroid polyp (IFP), or Vanek’s tumor, is a rare benign lesion of the gastrointestinal tract. Clinical manifestations of IFP vary based on size and location within the GI tract. Review of the literature indicates that early surgical intervention is the treatment of choice for intussusception caused by IFP. Lesions are typically reported as solitary, and resection is curative.

Keywords: Intussusceptions, inflammatory fibroid polyp, adenocarcinoma, gastrointestinal tract.

INTRODUCTION

Intussusception is uncommon among adult patients, which accounts for only 1 to 5% of the bowel obstruction cases [1, 2]. Majority of adult intussusceptions occur due to malignancies i.e. Colon adenocarcinoma. Small bowel intussusceptions are caused by benign neoplasms, and lipoma is the most common one. The inflammatory fibroid polyp (IFP), which is a reaction condition, is a rare cause of ileal intussusception [1]. Vanek’s tumor, or inflammatory fibroid polyp (IFP), is a rare lesion found throughout the gastrointestinal (GI) tract but is most commonly found in the gastric antrum [3]. Peak incidence occurs in the sixth and seventh decades of life predominantly in males [4]. Clinical manifestations of IFP ranges from abdominal pain, vomittings, intestinal obstruction, intussusception, and rarely GI bleeding [5, 6]. The authors report a case of terminal ileo ileal intussusception caused by IFP in a young patient.

CASE REPORT

We present a 32-yr male presented in emergency in GGS Medical College, Faridkot with history of pain whole abdomen for 4-5 days, which was sudden in onset and progressive in nature. Per abdominal examination -mild distension of whole abdomen and hyperactive bowel sounds were present.

X ray abdomen showed dilated small bowel loops with air fluid levels, centrally placed. Right colonic gas shadow couldn’t be visualised. Ultrasound abdomen showed telescoping of bowel loops in the right hypochondrium/ lumbar region- bowel within bowel appearance with? intussusception. Computed tomography of the abdomen and pelvis with oral contrast revealed ileoileal intussusception. Multiple dilated proximal small bowel loops were seen with air fluid levels - small bowel obstruction.

The patient was taken for laparotomy. OT Findings were as follows- there was ileoileal intussusception with adhesions present between the intussuscepted segment of the ileum and intussucipient segment (Fig 1). On limited manipulation of this segment intussusception was reversed and the lead point was identified. Intussusception part of the ileum was covered with hot sponges and active peristalsis was confirmed without any discoloration of the gut. (Fig 2) Intraluminal mass around10cm proximal to ileoceal junction within the healthy small intestine bowel loop, mass was adherent to intestine luminal wall, not movable with firm consistency. Incision was given at antimesentric border. On exploration pedunculated mass of 3-4cm was found extending upto the muscularis mucosa (Fig 3). Wedge resection was done of the involved ileum segment and primary Anastomosis was done. The specimen sent for histopathological examination which confirmed the diagnosis of inflammatory fibroid polyp. Postoperative period was uneventful.

© 2020 SAS Journal of Surgery | Published by SAS Publishers, India
Histopathological findings

Gross Examination showed polypoidal tissue with attached part of small intestine measuring 3x2.5x2 cm. The subjacent mucosa was noted to be edematous and hyperemic. Cut section – homogenous, grey white in colour. Upon microscopic examination- there were features of inflammatory fibroid polyp consisting of eosinophils. (Fig 4)

DISCUSSION

IFP was first described by Vanek, in 1949, as a "gastric submucosal granuloma with eosinophilic infiltration [1]". Confusion in the literature stems from the number of different names this rare benign lesion is known by, including Vanek’s tumour, eosinophilic granuloma, fibroma with eosinophilic infiltration, haemangiopericytoma and polypoid myoendothelioma[4]. Helwig and Ranier devised the generally accepted term, inflammatory fibroid polyp in 1953[5].

The fifth to seventh decade of life is the most common age at which patients present, both sexes being equally affected [4]. The most common site is the gastric antrum (60–70%), followed by small bowel (18–20%), colorectum (4–7%), and far less commonly (1%) in esophagus, duodenum, gallbladder, and appendix [2]. The polyps are typically solitary, but rare metachronous lesions have been reported in familial cases [7, 8]. Most IFPs grow intraluminally and are smaller than 4 cm, but case reports have discussed polyps up to 20 cm [9]. In this case, the polyp measured 5 cm in greatest dimension, with the majority of the mass extending in the lumen.

Clinical manifestations depend largely on tumor location and size. Often IFPs are asymptomatic and are identified incidentally during endoscopic or surgical procedures. Epigastric pain, vomiting and bleeding are commonly observed when the lesions are located in the stomach. In contrast, patients with IFP in the small bowel are more likely to present with chronic colicky abdominal pain, small bowel obstruction, intussusception, and weight loss [1]. Such situation occurs due to the presence of intraluminal mass, which can cause a motility disorder between both intestinal segments, thus causing bowel intussusception[1]. GI bleeding is a rare presenting symptom, and if present, it may indicate significant ulceration or ischemia [7].
X-ray abdomen and CT Abdomen are the imaging techniques for adults with intussusception, with sensitivity between 50 and 100%; however, visualization of a mass within the intussusception is rare [10]. In the adult population, once intussusception is diagnosed, prompt surgical intervention is warranted to avoid complications of ischemia, necrosis, and perforation. Debate in the literature on appropriate surgical treatment for adult intussusception focuses largely on initial resection of the intussuscepted segment versus reduction followed by a more limited resection. Current recommendations favor reduction and limited resection in small bowel intussusception, but only if the bowel is easily reduced and the lead point appears grossly benign [11]. In our patient we could reduce the intussusception and excise the polyp (alongwith wedge resection due to muscularis involvement).

A discussion highlighting the management, histology and features of inflammatory fibroid polyps makes surgeons aware of this uncommon disease.

REFERENCES


