
Inflammatory Fibroid Polyp Of The Jejunum Presenting As Intussusception: A Case Report And Review Of The Literature.

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Citation

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Abstract

Adult intussusceptions are rare entities, almost always associated with a demonstrable lead point. There has been a notable association between adult intussusceptions and inflammatory fibroid polyps (IFPs). These are benign mesenchymal tumor masses, composed of a spindle cell proliferation, accompanied by an inflammatory reaction and variably prominent blood vessels. The histogenesis of these tumors remain controversial, but they are believed to occur in response to local noxious stimuli.

INTRODUCTION

Adult intussusception is a rare entity and causes only 1% of all bowel obstruction¹. In the small bowel, benign lesions account for 70%, with malignant lesions responsible for the remainder². Inflammatory fibroid polyp (IFP) is a type of non-neoplastic inflammatory pseudotumor that occurs mainly in the gastric antrum or in the ileum, however, they are considered to be very rare in the jejunum and duodenum³. IFP has been reported to cause gastrointestinal bleeding and simple mechanical obstruction, but rarely cause intussusception.

Here, we present a rare case of Inflammatory fibroid polyp (IFP) confined to the jejunum presenting as intestinal intussusception in an adult female patient.

CASE PRESENTATION

A 63-year-old female patient presented to the hospital after experiencing colicky abdominal pain, nausea and vomiting that had begun 36 hours earlier. Similar attacks of pain she had experienced time after time over the past four months which had resolved spontaneously. There was no history of prior abdominal operations. Between the episodes the patient reported normal bowel habits. Gastroscopy which had been

undertaken previously was found to be normal.

Significant abdominal findings were confined to the abdomen that was distended with mild peri-umbilical tenderness and hyperactive bowel sounds. Digital rectal examination did not reveal the presence of feces, mucus or blood. An upright plain abdominal film revealed multiple loops of dilated small bowel in the left upper quadrant of the abdomen (Fig.1).

Figure 1

Plain abdominal radiograph showing distended loops of small bowel.



Abdominal ultrasonography failed to reveal the intussusception. Computed tomography (CT) was performed and showed a typical “sausage”-shaped soft tissue mass suggesting small bowel intussusception.

Vitals were stable: pulse 78/min, blood 140/80mm Hg, respiratory rate 16/min and temperature normal.

Laboratory investigations showed : Urea 22mg/dl(normal: 15-42 mg/dl, creatinine 0.7mg/dl(normal:0.6-1.3mg/dl), SGOT(ALT) 23u/l(normal:5-37u/l), SGPT(AST)52(normal 12-70u/l), Sodium 137mmol/l, potassium 4.4 mmol/l, haemoglobin 10.8g/dl, total leucocyte count 10.8x10³/mm³(normal: 4-11.000/l).

The clinical symptoms and CT images were consistent with small bowel obstruction. Thus urgent laparotomy with midline incision was performed. On exploration, an intussusception at the jejunum was found 40 cm distal to the duodeno- jejunal flexure(Fig.2), with an intraluminal obstructive mass as a lead point (Fig.3). The small bowel was dilated proximally. The intussuscepted segment was resected and end-to-end

anastomosis was carried out. The resected specimen was sent for histopathology.

Figure 2

Jejuno-jejunal intussusception.



Figure 3

IFP after section of the S-bowel.

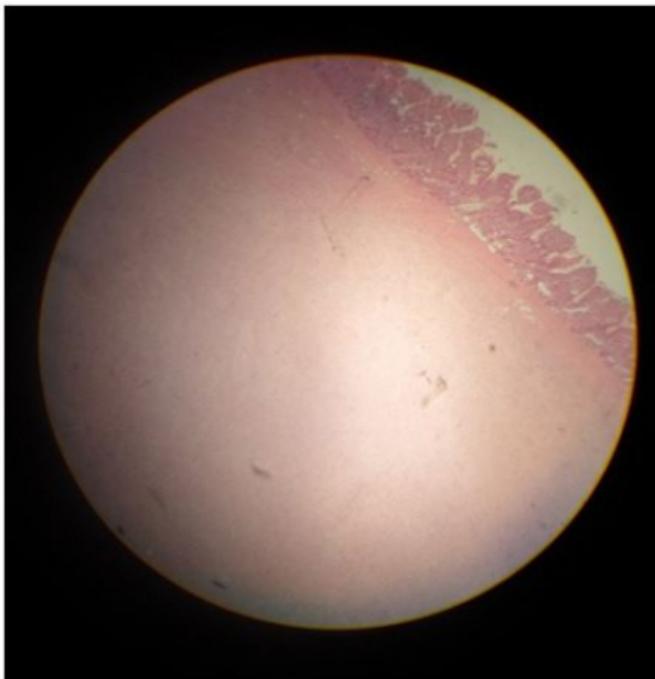


Gross pathologic examination of the resected segment of the jejunum revealed a 25x20x15mm sessile polypoid mass, located perpendicular to the plane of mucosa. Sectioning of the lesion revealed a homogeneous, hemorrhagic, tan with glistening surface. Sixteen lymph nodes were present in the attached mesentery.

Light microscopic examination showed portion of small bowel containing a submucosal mass(Fig. 4) with extension into the muscularis propria. The overlying mucosa was hemorrhagic with patchy area of ulceration. The mass was composed of a proliferation of spindle-shaped and stellate fibroblasts embedded within loose, oedematous stroma, which contained numerous blood vessels and an abundant inflammatory infiltrate comprising eosinophils, plasma cells and lymphocytes. All resected mesenteric lymph nodes showed reactive changes only.

Figure 4

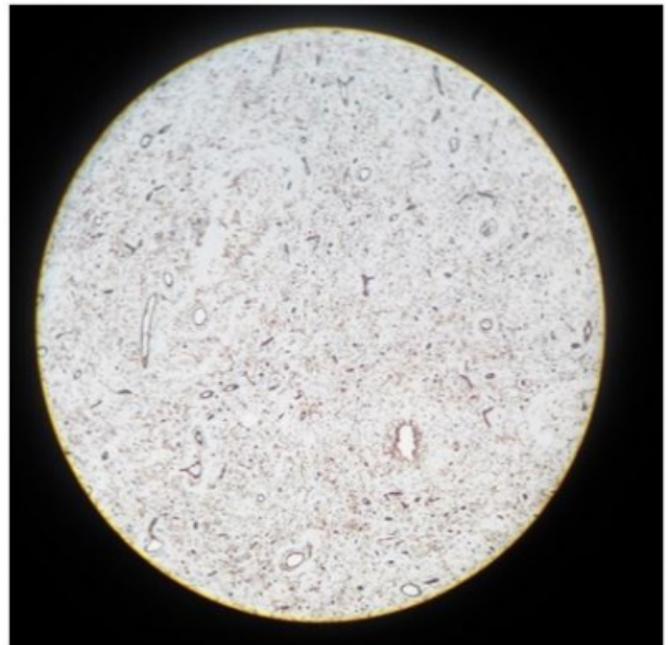
Histological appearance of IFP(H-E stain, original magnification x100).



The primary diagnoses were doubted between gastrointestinal stromal tumor(GIST) and inflammatory fibroid polyp. Immunohistochemical study was done on the specimen. The spindle cells were negative for CD117(C-kit), smooth muscle actin, desmin and S-100. Vimentin, fascin and cyclin D1 were strongly and diffusely positive. CD34 highlighted the vascular supply, although some of the spindle cells located between vessels were positive (Fig. 5).

Figure 5

Positive Immunostaining of spindle cells and vascular supply for CD 34(Immuno- histochemical stain, original magnificationx100).



The differential diagnosis also included neural and smooth muscle tumors. Immunohistochemistry performed on this lesion was negative for S100 and desmin, excluding a neurogenic and smooth muscle tumors. A final diagnosis of an inflammatory fibroid polyp of the jejunum was made.

The patient had an uneventful post-operative recovery, and was discharged 7 days after surgery . Six months after the surgery she remained well and symptom free.

DISCUSSION

Intussusception is a rare condition in adults, but can be the first manifestation of a tumor. It represents only 1% of all cases of intestinal obstructions in adults.1,2,3 The jejunum is the least likely location for intussusceptions to occur because the leading pathology is rare at this location of the gastrointestinal tract4,5,6,7. A jejuno-jejunal intussusception resulting from an IFP was first described by Winkler et al. in 1986.4 Since then, to the best of our knowledge, about 17 cases of jejunal IFPs, producing intussusception , have been described 2,4,5,6,7,8,9,10,11,12,13. IFPs can affect any portion of the GI tract, but they are most common in the gastric antrum (70%) and the terminal ileum(20%), but rarely in the jejunum. Akbulut et al.13

reported that among 85 patients with IFPs of the small bowel ileal intussusception was found in 63 patients and 17 had jejunal, three had colonic and two had ileojejunal intussusception. IFPs affect both sexes and all ages, with peak incidence in the fifth and sixth decades. This is likely because the lesions produce local symptoms that are size dependent.

The exact aetiology and pathogenesis of IFPs has not yet been clarified. Most regard IFPs as a reactive or inflammatory process in response to local noxious stimuli, with incomplete differentiation of myofibroblasts and primitive submucosal stromal cells^{14,15}. An allergic aetiology has been proposed due to the association with eosinophils¹⁰.

Immunohistochemical study with positive expression of spindle shaped cells with CD34 favor the hypothesis that IFPs represent a reactive proliferation of primitive perivascular/vascular cells¹⁴.

Macroscopically, IFPs tend to appear as solitary polyps, that are well-circumscribed. Larger lesions may be pedunculated, and the mucosal surface is often ulcerated on its apex. Size up to 18 cm in the largest dimension has been reported⁹. Histologically, the epicenter of the tumor is usually in the submucosa, but it can infiltrate into the overlying mucosa and the muscularis propria or serosa^{6,10,11}. The microscopic picture entails a loosely arranged fibroconnective tissue with a prominent thin-walled vascular component and an inflammatory response dominated by eosinophils. Cellular atypia and significant numbers of mitoses are not seen.

The GIST is an important consideration in the histological differential diagnosis of IFPs. In morphological ambiguous cases, immunohistochemistry is helpful to make a distinction between the two entities. Both tumors are positive for CD34 and Vimentin, but GISTs are positive for CD117(c-kit), while IFPs are not^{5,10,15}.

Negative staining to S-100 protein distinguishes an IFP from a neurogenic tumors which are positive for this marker. Some cases of IFPs fail to show positive staining for CD34, and this may be related to the evolution of the tumor^{14,16}.

Many IFPs are asymptomatic, and are typically identified during endoscopy or laparotomy. When

symptomatic, the clinical picture depends on the size of the tumor and on the location. Most of the jejunal IFPs have been found to cause small bowel obstruction as a result of intussusception^{3,4,,5,6,7,9}. If the overlying mucosa ulcerates then GI bleeding, anaemia and alteration of bowel habits were occasionally. Small bowel lesions are not usually diagnosed pre-operatively because they present with vague symptoms of bowel obstruction due to intussusception. Laboratory investigations and plain radiographs are not helpful in making the diagnosis as they will demonstrate nonspecific findings that are more in keeping with bowel obstruction. Abdominal CT scanning, is currently the diagnostic instrument of choice¹⁷, allowing in some series a correct preoperative diagnosis in up to 80% of the cases¹. The CT appearance is complex. According to the cut axis, the intussusception appears as a "sausage" or a "target" mass. There are no unique identifying features on imaging that can distinguish the nature of the lead point. The absence of pathognomonic clinical or radiological features makes operative resection and microscopic examination of the resected lesion mandatory.

Operative resection of the involved intestine with free margins is sufficient treatment. In conclusion, IFP is a rare benign cause of enteric intussusception in adults. This diagnosis should be considered by Surgeons when intussusceptions are encountered in adults.

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