

Inflammatory Myofibroblastic Tumor of the Colon with an Unusual Presentation of Intestinal Intussusception

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Abstract

Keywords

- ▶ inflammatory myofibroblastic tumor
- ▶ intussusception
- ▶ childhood

Inflammatory myofibroblastic tumor (IMT), also known as inflammatory pseudotumor is unusual, benign solid tumor. This tumor is commonly reported in the lungs but can be present in extrapulmonary sites as well. We present the case of a 7-year-old girl with IMT in an unusual location. The patient was admitted with abdominal pain, and ultrasound showed a solid mass in the abdomen. She was operated and colocolic intussusception secondary to a mass was found. Histologic evaluation of mass revealed IMT.

Introduction

Inflammatory myofibroblastic tumor (IMT) is rare and may be grouped within the family of fibroinflammatory disorders. This tumor affects predominantly children and young adults, but patients of any age and sex can be affected.¹ Its clinical symptoms are also diverse according to the location of the lesion. A mass, fever, weight loss, malaise, pain, and site-specific symptoms were the most common presenting complaints or manifestations.¹ However, IMT seldom presents in the colon and intussusception is a rare complication of this tumor. We document this unusual presentation of IMT in this case report. There are few pediatric IMT cases in the literature such as our patient. Because of this presentation, IMT should be considered in patients with colocolic intussusception.

Case Report

A 7-year-old girl was admitted with abdominal pain to our center. On physical examination, the patient had a palpable mass in the midline of the abdomen. Abdominal ultrasound showed a solid mass which was 6 × 5 × 4 cm, hypervascular

and partly calcified areas in the center. The white blood cell count was 7,300/mm³, hemoglobin 6.8 g/dL, platelet count 554,000/mm³, C-reactive protein 154 mg/L, erythrocyte sedimentation rate 27 mm/h, biochemical values were normal. Immunoglobulin (Ig) A was 222 mg/dL (normal, 70–312 mg/dL), Ig M was 114 mg/dL (normal, 56–352 mg/dL), IgG was 1,030 mg/dL (normal, 639–1,349 mg/dL). Patient was operated and colocolic intussusception was found. Involved segment was beginning from 4 cm distal of the ileocecal valve and including a polypoid mass. Resection of the involved segment of the ascending colon was done with end-to-end anastomosis.

The polypoid mass was 5.5 × 4.5 × 3.5 cm wide, sessile, sticking to the mucosa with an area of 3.5 × 3 cm macroscopically. The tumor covered the bowel wall at full thickness (▶Fig. 1). The tumor composed of fibroblast/myofibroblast-like cells which were spindle, vesicular nuclei, some places with prominent nucleoli, and unclear boundaries of the cytoplasm. The tumor cells contained prominent mixed inflammatory cells including numerous plasma cells (▶Fig. 2). Immunohistochemistry was positive for actin and negative for CD117, desmin, S-100, pancytokeratin, bcl-2, and

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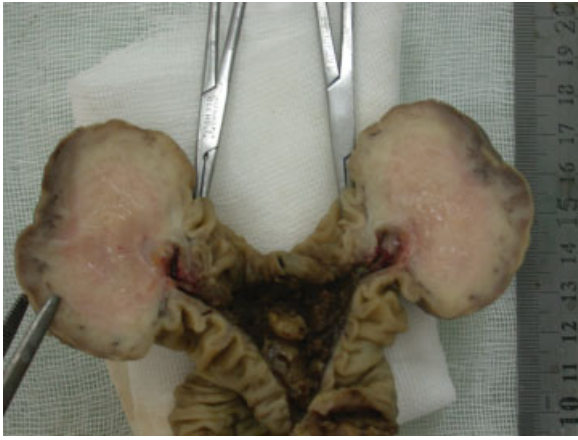


Fig. 1 Macroscopic view of polypoid mass emanating from bowel wall. The surface seems ulcerated and cut surface exhibits solid, white, fibrous texture.

anaplastic lymphoma kinase (ALK) (–Fig. 3). Tumor cells were negative for CD34. Evaluation of mass revealed IMT of colon.

After surgery, laboratory values of the patient improved dramatically. One year later, the patient was evaluated through the colonoscopy. There were two polypoid structures on anastomosis line and their pathological examinations were nonspecific. We are following up the patient with biochemical examinations and abdominal scanning closely for tumor recurrence.

Discussion

IMT was first described in 1937 as a primary lung tumor. The tumor consists of the proliferation of fibroblastic–myofibroblastic cells admixed with inflammatory infiltrate.¹ Independent of the location, IMT is more common in children and young adults, but patients of any age and sex can be affected.¹ IMT is rare and may be grouped within the family of fibroinflammatory disorders. This benign tumor typically resides in

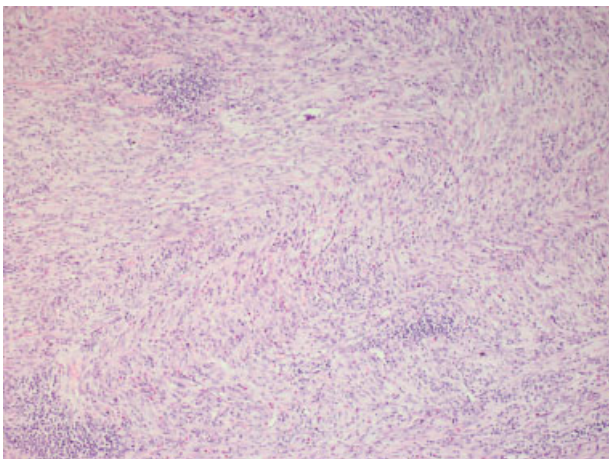


Fig. 2 A fascicular pattern in which the lesional spindle cells were arranged in moderately cellular with intersecting fascicles with a heavy infiltrate of lymphoplasmacytic cells in the background. Their nuclei were elongated with occasional small nucleoli (hematoxylin and eosin stain; $\times 10$).

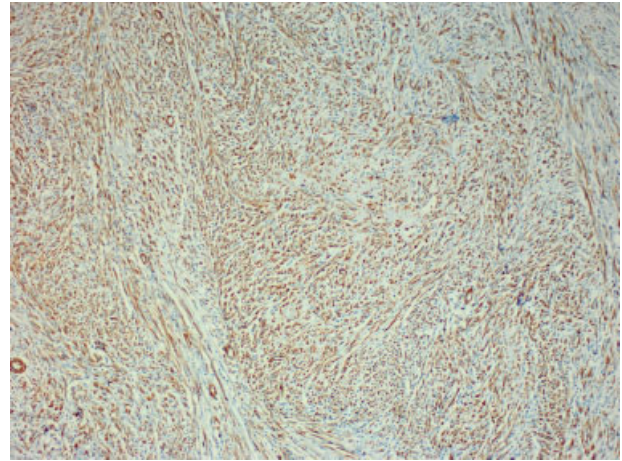


Fig. 3 The expression of smooth muscle actin by immunohistochemistry in the spindle cells (antismooth muscle actin; $\times 10$).

the lung; however, multiple extrapulmonary manifestations have been reported.^{2,3} Sites of extrapulmonary IMT may include the upper respiratory tract, mesentery/omentum, genitourinary tract, gastrointestinal tract, mediastinum, retroperitoneum, pelvis, trunk, extremities, head, neck, spleen, brain, pancreas, and liver.⁴

The etiologic factors responsible for development of IMT are not clearly established. Some investigators believe it is a true neoplasm, and the others believe that it represents an immunologic response to an infectious or noninfectious agent.^{1,5} But the etiology of the mechanism still remains unclear.

IMT is typically considered as benign lesion, but a spectrum including locally destructive variants is expectable. Biselli et al demonstrated that chromosomal aberrations may be present in IMT.⁶ Cessna et al describes abnormalities of chromosome 2p23 with expression of p80 and ALK1 in up to 40% of IMTs.⁷ These studies have added molecular credence to the clinical observation that there is a spectrum of aggressiveness among IMTs, and some may exhibit aggressive local behavior and rarely metastasis. Up to 71% are positive for ALK1, and approximately 50% have clone rearrangement involving the ALK locus on chromosome 2p23 that provides support for the neoplastic nature of IMT. The ALK gene encodes a receptor tyrosine kinase that has strong oncogenic activity when constitutively activated.^{7–9} However, as a predictor, ALK expression has not demonstrated a clear relationship with prognosis. On the one hand, ALK1-negative tumors occur in older patients and display greater nuclear pleomorphism and atypical mitoses; in two recent series, metastatic IMTs were confined to ALK1-negative lesions.^{8,10} Whereas on the other hand, positive ALK1 status may be more frequent in younger males and associated with a higher recurrence rate.¹¹ In our young case, we were unable to demonstrate ALK cytogenetic analysis, ALK immunohistochemistry analysis was found negative and there was no atypia or recurrence.

Microscopically, IMT also shows a similarity to a gastrointestinal stromal tumor. An IMT consists of the proliferation of spindled to epithelioid myofibroblasts with admixed inflammatory cells, predominantly of mononuclear type. Tumor

cells without atypia or hyperchromatism can invade the muscularis propria and even the adventitia. For final diagnosis of an IMT, immunohistochemistry analysis is very important. Tumor cells are characteristically positive for vimentin and do not express CD117 and CD34 in IMT.^{1,12} The cells are positive for smooth muscle actin (SMA) with or without desmin expression and S100 positive.¹³ In our case, tumor cells were common positive for actin, negative for S100, ALK, and there was no desmin, CD117, CD34 expression.

Bonnet et al reported that IMT cases documented in the mesentery/omentum were in fact mostly located in the mesentery of the small bowel.¹⁴ Clinical presentation of IMT varies markedly, depending on the site at which the tumors originated. At the intestinal location, the onset may be insidious or rapid and may be accompanied by weight loss, malaise, a mass, and a variety of laboratory abnormalities.⁵ Childhood IMT cases with an intramural location of the tumor have interestingly presented with diarrhea and intestinal obstruction.¹⁵ Rarely, the presentation may be a complicated picture such as intestinal obstruction and intussusception.^{16,17} Colonic IMT is a rare condition in abdominal IMT and the most common site has been the right colon. Colonic IMT can be served as a lead point for intussusception.^{18,19} There are few pediatric cases of colonic IMT with intussusception in the literature.

The incidence of local recurrence has been reported to be 15 to 37% in large series of children presenting with IMT of mesentery and retroperitoneum.¹ We evaluated our case through colonoscopy for this recurrence risk and no recurrence was detected in the colon.

Patients may have hypochromic microcytic anemia, thrombocytosis, polyclonal hyperglobulinemia, and elevated erythrocyte sedimentation rate in IMT. However, following a complete resection abnormal laboratory tests recover rapidly.²⁰ Hyperglobulinemia was not observed in our case. She has hypochromic microcytic anemia, elevated erythrocyte sedimentation rate, and elevated platelet count. After surgery, all laboratory tests improved.

Surgical excision should be the primary therapy for cases with IMT.¹⁷ Complete resection is associated with less than 10% risk of local recurrence. We have performed surgical excision of the mass in our patient and no recurrence has occurred yet. Chemotherapy has been reserved for patients in whom resection is morbid, impossible or in patients who have incomplete resections. There is no evidence to prove that chemotherapy is effective when used singly, however, it may play a role following complete resection. There is little evidence within the literature regarding chemotherapy for IMT, and the majority of data are within the pediatric population.⁴ Radiation is typically reserved for palliation, to alleviate the mass effect of the IMT, or in conjunction with chemotherapy for cure in patients who are not suitable to resection. Radiation treatment has been shown to be of some benefit in pulmonary IMT.²¹ As with chemotherapy, there is currently no evidence to support routine use of radiotherapy in patients who have complete resection. Steroids may be added to reduce surrounding inflammation if deemed necessary, particularly in cases that involve the central nervous system. Nonsteroidal anti-inflammatory drugs (NSAIDs) have been

successful in the treatment of IMTs when resectability is limited due to the tumor invading vital structures.²² Chemotherapy combined with oral NSAIDs may be an effective therapeutic option for patients with unresectable IMT.²³

In conclusion, colonic IMT is a rare condition and intussusception is a rare complication of this tumor. Therefore, presentation of this case is very important for the literature. Because of recurrence risk the clinical and laboratory follow-up should be done in these cases.

Conflict of Interest

None.

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