Inflammatory myofibroblastic tumor (IMT) is a histologically idiosyncratic lesion having an intermediate biological behavior and potential for local recurrence. IMT belongs to a group of pseudosarcomatous conditions with many names including plasma cell granuloma, plasma cell pseudotumor, inflammatory myofibrohistiocytic proliferation, omental-mesenteric myxoid hamartoma and, most commonly, inflammatory pseudotumor. These lesions are mostly found in the viscera and soft tissue of young adolescents and usually pursue a benign clinical course. Age range is broad for the initially described lesion of the lung; however, extrapulmonary IMTs show a predilection for children with a mean age of 10 years.

We are reporting an unusual case of appendiceal IMT in a 14-year-old boy who presented with symptoms and signs of acute appendicitis. A review of the literature and comparison with other similar published cases is also included.

**CASE REPORT**

A 14-year-old boy was referred to the emergency room with acute abdomen with right-sided iliac fossa pain and tenderness and fluctuating nausea and vomiting for two days. On physical examination, the rebound tenderness sign was not significantly positive.

Ultrasonographic examination showed a mildly echogenic oval-shaped area measuring 31 × 26 mm in the right iliac fossa and a connection with an attached hypoechoic tubular structure. This was reported as mucocele of the appendix [Figure 1]. Contrast-enhanced computed tomography scan was suggested to enable further evaluation. However, the surgeon preferred to proceed with surgery based on a strong clinical preoperative diagnosis of acute appendicitis.

An emergency laparotomy was performed during which the surgeon found a firm to solid-looking mass encircling the tip and firmly adherent to the appendiceal wall. This was difficult to excise intact with healthy surgical resection margins. This intraoperative scenario gave the impression of sarcoma or an appendicular tumor.

Microscopic examination showed histological features of a mesenchymal neoplasm composed of loosely arranged plump to spindle-shaped myofibroblastic cells in an edematous myxoid background with proliferating blood vessels and an infiltrate of plasma cells, lymphocytes, and eosinophils [Figure 2]. The features were suggestive of a low malignant potential/borderline...
gastrointestinal stromal tumor (GIST) or a low-grade fibrohistiocytic neoplasm. The presence of neutrophils within the muscularis propria of the appendiceal wall was evidence of acute appendicitis. A panel of immunohistochemical stains were performed. The tumor cells were focally positive for desmin [Figure 3] and negative for CD117, CD34, DOG-1, and activin receptor-like kinase (ALK-1) protein. Based on the immunohistochemistry and morphological features a final diagnosis of IMT was confirmed with the exclusion of GIST and mucinous neoplasms of the appendix. The patient’s postoperative course was uneventful.

**DISCUSSION**

The appendix is a very rare site for IMT and this is one of the reasons for writing this case report. To the best of our knowledge, only 11 cases of appendiceal IMT have been reported excluding one recent report, which had a preceding event of gastric cancer treatment. The last case was reported in 2012 and was associated with a rare finding of mucosal dysplasia. Extrapulmonary IMTs show a tendency for occurring in children, with a mean age of approximately 10 years. Females are affected slightly more commonly than males. However, the gender in almost all reported cases of appendiceal IMT is male.

Current literature review seems to support a neoplastic origin for IMT with intermediate malignant potential in a small subset of cases described in the recent World Health Organization classifications. The occurrence of chromosomal aberrations (30–40% of cases) in these tumors suggests that IMT is a neoplastic proliferation.
of clonal origin and is associated with aggressive clinical behavior.8

Bonnet et al,8 reviewed 74 abdominal cases of IMTs in children. They found only a single case with appendiceal origin which was the first reported case by Narasimharao et al.9 The patient was an 8-year-old boy who presented with symptoms of appendicitis. The last and eleventh case was reported in 2012 but with a very rare finding of dysplasia in addition to appendiceal IMT, which had not been observed in any previously described cases.7 In our case, laboratory findings showed significantly increased neutrophils, which is consistent with a finding of leukocytosis. This was also highlighted in a case reported by Oeconomopoulou et al,10 in 2016. In our case, clinical and preoperative impression of acute appendicitis were somewhat confused and changed to an appendiceal tumor during intraoperative analysis.

It is well-known that the clinical findings, laboratory investigations and even radiological examinations in most instances give no diagnostic clue; hence a definitive diagnosis is based on the histopathological and immunohistochemical findings performed on the resected tumor. Microscopically, IMT is composed of myofibroblastic spindle cells arranged in haphazard fascicles accompanied by an inflammatory infiltrate of plasma cells, lymphocytes, and eosinophils.3 Three histological patterns have been described including fibromyxoid, which was identified in our case. Atypia is rare in these tumors with few or no mitotic figures.3,10

Immunohistochemistry plays an important role in better defining the lesion with the presence of positive vimentin, smooth muscle actin, muscle-specific actin, desmin, and focally for cytokeratins in the majority of cases. Overexpression of ALK-1 protein is seen in 50–60% of IMTs, which is considered significant if positive but in our case was negative.1 This immunophenotypic feature is associated with a local recurrence rate of 15–37% in a large series of children.14 Another panel of immunohistochemical markers including CD117, S-100, CD34, and DOG-1 was performed to exclude other spindle cell neoplasms with GIST as a primary suspect. Hence, the final pathological diagnosis was IMT.

CONCLUSION

Appendiceal IMT is a rare mesenchymal neoplasm of intermediate biological potential, which may occur alone or coexist with acute appendicitis. Although radiology is not characteristic of these lesions, it can be an adjunct to pre- or intraoperative differential diagnosis. The final diagnosis should be based on histopathological and immunohistochemical findings. Complete surgical excision of appendiceal IMT remains the mainstay of treatment. A careful one-year follow-up is recommended because of the risk of local recurrences.

Disclosure

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