Case Report

Gastric calcifying fibrous tumor of the stomach presenting as a subepithelial tumor

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Abstract We describe a case of a 48-year-old Korean woman who had a subepithelial mass incidentally discovered by endoscopic examination. Endoscopic mucosal resection revealed a well-circumscribed whitish solid mass within the submucosal space. Microscopically, the tumor was comprised of sparse spindle cells in the dense collagenous stroma with several calcifications and lymphoid aggregates. Immunohistochemical analysis showed that the tumor cells are negative for c-kit, smooth muscle actin, desmin, S-100 and CD34. Based on these findings, the tumor was diagnosed with calcifying fibrous tumor.

Key words: Calcifying fibrous tumor, Stomach, Subepithelial tumor

INTRODUCTION

Calcifying fibrous tumor (CFT), also called calcifying fibrous pseudotumor, is a benign mesenchymal tumor characterized by the presence of abundant hyalinized collagen, interspersed bland fibroblastic spindle cells, psammomatous or dystrophic calcifications and lymphoplasmacytic infiltration.¹⁾ It usually affects children and young adults and shows a predilection for the subcutaneous and deep soft tissues in the extremities. However, gastric CFT is very rare. Here, we report a case of gastric CFT that has been incidentally detected as a subepithelial mass by esophagogastroduodenoscopy examination.

CASE REPORT

A 48-year-old female patient was admitted for gastric

tumor that had been incidentally found at medical check-up. The patient underwent an endoscopic evaluation, and which revealed a round subepithelial tumor at greater curvature of high body, and which was an 1-cm sized and relatively well-circumscribed mass. Contrast-enhanced computed tomography scan confirmed a 1cm sized, non-enhanced, and homogenous nodular lesion in the same location. Endoscopic mucosal resection was performed.

Cross sections revealed an oval-shaped nodule in the submucosal area, and which showed a whitish hard mass. Grossly, the tumor was covered by intact mucosa. On section, it was a well-defined white mass located in the submucosa, measuring 1 × 0.6 cm size. Microscopically, the tumor was characterized by a sparse spindle cell lesion in dense collagenous stroma which was highlighted by Masson's trichrome special stain. Spindle cells had ovoid vesicular nuclei with fine chromatin, but did not show any significant nuclear atypia or pleomorphism. In addition, no mitotic activity was observed. A couple of lymphoid follicles harboring germinal centers or small lymphoid aggregates were scattered in the tumor and a

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Figure 1. (A) Endoscopic evaluation reveals a polypoid subepithelial mass with intact overlying mucosa at high body of stomach (B) A computed tomography image shows a non-enhanced and homogenous small nodule in the stomach (indicated by red arrow).

Figure 2. Histologic findings of calcifying fibrous tumor. (A) The tumor is a well-defined oval shaped, with dense collagenous tissue in the sub-mucosa. H&E (× 1.25) (B) Special staining highlights the dense fibrotic background. Masson’s trichrome (× 1.25) (C) A lymphoid follicle is observed within the tumor. H&E (× 100) (D) Sparse spindle cells are interspersed among thick collagen bundles with dystrophic calcification. H&E (× 400)

Figure 3. Immunohistochemical analysis for calcifying fibrous tumor. (A) Tumor is characterized by a paucicellular fibroblastic lesion with marked sclerotic stroma. (B-F) Spindle tumor cells showed no immunoreactivity for CD34, S-100, smooth muscle actin, desmin, and c-kit. ($\times 200$)

couple of dystrophic calcifications were observed. Various inflammatory cells including plasma cells, mast cells, and eosinophils were also found around small vessels. Immunohistochemically, the spindle cells were all negative for CD34, smooth muscle actin (SMA), desmin, S-100, and c-kit.

This study was approved by the Institutional Review Board of Jeju National University Hospital (IRB No. 2018-12-002) and informed consent was waived.

DISCUSSION

CFT was originally defined as benign, soft, and fibrous mass with psammomatous calcifications in two young girls.²⁾ It is characterized by a heavily collagenized paucicellular spindle cell lesion, psammomatous and/or dystrophic calcifications and varying degrees of inflammatory infiltrates. CFT was initially considered to represent a reactive process, it was also described as calcifying fibrous pseudotumor.¹⁾ However, other studies claimed that it is neoplasm with a tendency of non-destructive local recurrence.³⁾

The cause and pathogenesis of CFT remains unclear, but it has been hypothesized that CFT represents a sclerosing end stage of inflammatory fibroblastic tumor (IMT). Despite histologic similarities, including fibroblastic pro-

liferation and mixed inflammatory infiltration, CFT has distinctive histologic and immunohistochemical features that are different from IMT. For example, while CFT demonstrates varying degrees of calcifications, IMTs rarely contain calcifications.⁴⁾ Immunohistochemically, CFTs are diffuse positive for Factor XIIIa and negative for SMA, and CD34, while IMTs are diffusely positive for SMA, variable positivity for CD34 and focal positivity for Factor XIIIa.⁴⁾

Although gastric CFT is very rare, there have been several reports of CFTs that developed in the stomach.⁵⁻¹³⁾ Gastric subepithelial tumors are mostly asymptomatic and detected incidentally during endoscopic examinations. The most common subepithelial tumors of stomach are gastrointestinal stroma tumors (GISTs) that derived from interstitial cells of Cajal, and other differential diagnosis includes a variety of mesenchymal tumors such as leiomyoma, schwannoma, inflammatory fibroid polyp, inflammatory myofibroblastic tumor, and plexiform fibromyxoma. Histologically, sclerosing GIST, sclerosing leiomyoma and ancient schwannoma may be in part similar to CFT. However, psammomatous calcifications and lymphoplasmastic infiltrates are quite unique features of CFT. In addition, the majority of resected cases are diagnosed as follows based on the immunohistochemical positivity for c-kit (GIST), S-100 (schwannoma), and SMA (leiomyoma). The subepithelial tumor in our patient did

not express any of these markers and showed typical sclerosing stroma and calcifications, which are consistent with CFT.

Gastric CFT tends to be under-recognized in part due to its low incidence and small size. However, characteristic histology of gastric CFTs may help to recognize them. For example, the calcification detected by endoscopic ultrasonography can be considered a useful feature for suspecting CFTs.

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