Gastric plexiform fibromyxoma resected by endoscopic submucosal dissection after observation of chronological changes: A case report

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Abstract

A 66-year-old man was diagnosed with a gastric submucosal tumor. Endoscopic ultrasound (EUS) revealed an iso/hypoechoic mass in the third layer. No malignant cells were detected in a histological examination. Yearly follow-up endoscopy and EUS showed the slow growth of the tumor. Endoscopic submucosal dissection (ESD) was performed and a glistening tumor was resected. The lesion showed a multinodular plexiform growth pattern consisting of spindle cells with an abundant fibromyxoid stroma that was rich in small vessels. The tumor was diagnosed as plexiform fibromyxoma (PF) by immunohistochemistry. Although difficulties are associated with reaching a diagnosis preoperatively, chronological changes on EUS may contribute to the diagnosis of PF. ESD may also be useful in the diagnosis and treatment of PF.

Key words: Plexiform fibromyxoma; Plexiform angiomyxoid myofibroblastic tumor; Endoscopic ultrasound; Endoscopic submucosal dissection; Gastrointestinal stromal tumor
Core tip: Plexiform fibromyxoma (PF) is a very rare gastric submucosal tumor. Therefore, difficulties are associated with diagnosing PF preoperatively, particularly in a differential diagnosis of gastrointestinal stromal tumors with cystic changes. We suggest that the chronological changes observed by endoscopic ultrasound contribute to the preoperative diagnosis of PF. Furthermore, endoscopic submucosal dissection needs to be considered for the diagnostic treatment of PF without muscle invasion.

INTRODUCTION

Plexiform fibromyxoma (PF), also known as a plexiform angiomyxoid myofibroblastic tumor (PAMT), is a very rare gastric submucosal tumor (SMT) with a unique plexiform growth pattern of bland spindle cells[1-3]. Few studies have described the endoscopic ultrasound (EUS) characteristics of PF, and its chronological changes also remain unclear. We herein report a case of PF resected by endoscopic submucosal dissection after observation of chronological changes: A case report. World J Gastrointest Oncol 2017; 9(6): 263-267 Available from: URL: http://www.wjgnet.com/1948-5204/full/v9/i6/263.htm DOI: http://dx.doi.org/10.4251/wjgo.v9.i6.263

CASE REPORT

A 66-year-old man was referred to our institute for the management of a gastric tumor. An endoscopic examination revealed a SMT, approximately 20 mm in diameter, located in the antrum (Figure 1A). EUS showed an iso/hypoechoic mass in the third layer (Figure 1B). Computed tomography (CT) displayed a poorly enhanced lesion (Figure 2). Endoscopic biopsy and endoscopic ultrasound-guided fine-needle aspiration (EUS-FNA) were performed. Histological findings showed no malignant cells, and no further diagnosis was made.

Yearly follow-up endoscopy revealed the slow growth of the tumor, which became pedunculated and showed transpyloric prolapse (Figures 1C-F). EUS revealed gradual increases in the solid and multicystic components without muscle invasion. Based on these findings, our preoperative diagnosis was a hamartomatous inverted polyp[4-6]. In order to avoid outlet obstruction and reach a histological diagnosis, ESD was performed (Figure 3).

On dissection, a glistening, 40 mm × 30 mm tumor covered with normal gastric mucosa was identified. Microscopically, the lesion showed a multinodular plexiform growth pattern, and consisted of bland spindle cells separated by abundant intercellular myxoid or fibromyxoid matrix. The stroma was rich in small vessels (Figure 4).

Immunohistochemical tests revealed that tumor cells were focally positive for smooth muscle actin (SMA), muscle-specific actin (HHF35), and calponin, but were negative for c-kit, CD34, DOG-1, desmin, the S-100 protein, CD10, and h-caldesmon. The Ki-67 labeling index was approximately 2% (Figure 5). The pathological assessment led to a diagnosis of PF. Resected margins were histologically tumor-free. Although vascular invasion was positive, the patient did not undergo surgery due to the reportedly good prognosis of PF[1,7], and remained under careful observation.
by endoscopy and CT follow-up. There was no recurrence or metastasis in the 12-mo follow-up.

**DISCUSSION**

Gastric PF is a new benign mesenchymal tumor that has been adopted by the 2010 WHO classification of tumors of digestive system\(^8\). The term PAMT is also used for this type of tumor. The distinction between these terms has been controversial\(^7,9\). Previous studies reported that most cases of this tumor are found in the antrum, with approximately half extending into the extragastric...
soft tissues or proximal duodenum\textsuperscript{2-7}. The diagnosis of PF is based on its histological features, including immunohistochemical findings\textsuperscript{1}. Its histology indicates a plexiform growth pattern composed of spindle cells, fine small vessels, and a myxoid matrix. Tumor cells are typically immunoreactive for SMA and HHF35, whereas c-kit, CD34, DOG-1, and the S-100 protein are nearly completely negative. Focal immunoreactivity for CD10, caldesmon, or desmin has occasionally been detected\textsuperscript{1,3,7}.

In the present case, endoscopy and EUS showed that the tumor grew gradually, with increases in the solid and multicystic components. Spindle cells, with a rich vascular myxoid stroma, were considered to be detected as an isoechoic lesion and fluid leakage was observed as a hypoechoic lesion.

Previous studies reported the lack of recurrence or metastasis of PF after excision\textsuperscript{1,7}; however, Miettinen et al\textsuperscript{1} demonstrated that some plexiform elements showed intravascular involvement, suggesting that PF occasionally spreads through vessels. Since our case also exhibited vascular invasion, follow-up examinations were carefully performed. Since no patients have developed recurrence, annual endoscopy and CT are considered to be sufficient to monitor patients.

Although PF is considered to be benign, distal or partial gastrectomy is generally performed under the assumption of the presence of GIST\textsuperscript{7}. Although GIST typically appears as a solid mass, few studies have described myxoid GIST that also shows a plexiform growth pattern\textsuperscript{10}, and some cases of GIST have shown cystic changes as a result of degeneration or necrosis\textsuperscript{11-13}. Thus, it may be difficult to distinguish PF from these GIST by performing EUS only once. The chronological changes observed in the present case may contribute to a preoperative diagnosis of PF and the elucidation of its growth process. In this case, even though contrast-enhanced EUS was not performed, it may also be useful for reaching a differential diagnosis\textsuperscript{14,15}. The distinction of PF from a hamartomatous inverted polyp is also important. EUS-FNA is the first choice for a definite diagnosis of SMT\textsuperscript{16}. Nevertheless, ESD remains an important option for diagnostic treatment, including that for cases of gastric SMT of the submucosal layer\textsuperscript{6,17}. Since EUS-FNA revealed no abnormalities in the present case, ESD was selected as a second choice. We performed en bloc ESD, which allowed for the diagnosis of PF. To the best of our knowledge, this is the first case report to describe the successful resection of PF by ESD. Further studies are needed in order to establish the appropriateness of ESD for PF.

### COMMENTS

#### Case characteristics

A 66-year-old man presented with a gastric tumor located in the antrum.

#### Clinical diagnosis

Gastric submucosal tumor.

#### Differential diagnosis

A hamartomatous inverted polyp, myxoid gastrointestinal stromal tumor (GIST), and GIST with cystic degeneration.

#### Laboratory diagnosis

Laboratory test results were within normal limits.

#### Imaging diagnosis

Endoscopic ultrasound revealed an iso/hypoechoic mass of 20 mm in diameter in the third layer, and it showed gradual increases in the solid and multicystic components without muscle invasion.

#### Pathological diagnosis

Plexiform fibromyxoma.

#### Treatment

Endoscopic submucosal dissection was performed as a diagnostic treatment.

#### Related reports

Few studies have described plexiform fibromyxoma, also known as a plexiform angiomyxoid myofibroblastic tumor. Patients with plexiform fibromyxoma have generally undergone distal or partial gastrectomy.

#### Term explanation

Plexiform fibromyxoma is a new mesenchymal tumor entity that shows a unique...
plexiform growth pattern of bland spindle cells.

**Experiences and lessons**

Plexiform fibromyxoma needs to be considered in a differential diagnosis of gastric submucosal tumors, and follow-up endoscopic ultrasound (EUS) may be able to distinguish plexiform fibromyxoma from other gastric submucosal tumors.

**Peer review**

The rarity of the case could be enriched with a brief review of the literature, due to the scarce number of papers reporting similar tumors. Moreover it could be interesting to expand data about EUS, for example explaining the characteristics of elastometry and eventual contrast enhancement. The quality of the article is augmented by the images, which are impressive and clear. Overall it is a good paper.

**REFERENCES**

1. Miettinen M, Makhlouf HR, Sobin LH, Lasota J. Plexiform fibromyxoma: a distinctive benign gastric antral neoplasm not to be confused with a myxoid GIST. *Am J Surg Pathol* 2009; 33: 1624-1632 [PMID: 19675452 DOI: 10.1097/PAS.0b013e3181a0660a]
2. Sakamoto K, Hirakawa M, Atsumi K, Mimori K, Shibata K, Tobo T, Yamamoto H, Honda H. A case of gastric plexiform fibromyxoma: radiological and pathological findings. *Jpn J Radiol* 2014; 32: 431-436 [PMID: 24744134 DOI: 10.1007/s11604-014-0315-z]
3. Takahashi Y, Shinzui S, Ishida T, Aita K, Toida S, Fukusato T, Mori S. Plexiform angiomyxoid myofibroblastic tumor of the stomach. *Am J Surg Pathol* 2007; 31: 724-728 [PMID: 17460456 DOI: 10.1097/01.pas.0000213448.54643.22]
4. Yang TC, Hou MC, Chen PH, Liao WC, Li AF. Gastric hamartomatous inverted polyp mimicking ectopic pancreas on endoscopy and endosonography. *Endoscopy* 2014; 46 Suppl 1 UCTN: E119-E120 [PMID: 24676819 DOI: 10.1055/s-0034-134888]
5. Aoki M, Yoshida M, Saikawa Y, Otani Y, Kubota T, Kumai K, Wakabayashi G, Omori T, Mukai M, Kitajima M. Diagnosis and treatment of a gastric hamartomatous inverted polyp: report of a case. *Surg Today* 2007; 34: 532-536 [PMID: 15170552 DOI: 10.1007/s00595-004-2761-1]
6. Odashima M, Otaka M, Nanjo H, Jin M, Horikawa Y, Matsushima T, Ohba R, Koizumi S, Kinoshita N, Takahashi T, Shima H, Watanabe S. Hamartomatous inverted polyp successfully treated by endoscopic submucosal dissection. *Intern Med* 2008; 47: 259-262 [PMID: 18277026]
7. Takahashi Y, Suzuki M, Fukusato T. Plexiform angiomyxoid myofibroblastic tumor of the stomach. *World J Gastroenterol* 2010; 16: 2835-2840 [PMID: 20556828 DOI: 10.3748/wjg.v16.i23.2835]
8. Miettinen M, Fletcher CD, Kindblom LG, Tsai WM. Mesenchymal tumours of the stomach. In: Bosman FT, Carneiro F, Hruban R, Theise ND, editors. Mesenchymal tumours of the stomach. WHO classification of tumours of the digestive system. Lyon: IARC; 2010: 74-79
9. Li P, Yang S, Wang C, Li Y, Geng M. Presence of smooth muscle cell differentiation in plexiform angiomyxoid myofibroblastic tumor of the stomach: a case report. *Int J Clin Exp Pathol* 2014; 7: 823-827 [PMID: 24551311]
10. Li B, Zhang QF, Han YN, Ouyang L. Plexiform myxoid gastrointestinal stromal tumor: a potential diagnostic pitfall in pathological findings. *Int J Clin Exp Pathol* 2015; 8: 13613-13618 [PMID: 26722584]
11. Miettinen M, Sobin LH, Lasota J. Gastrointestinal stromal tumors of the stomach: a clinicopathologic, immunohistochemical, and molecular genetic study of 1765 cases with long-term follow-up. *Am J Surg Pathol* 2005; 29: 52-68 [PMID: 15613856]
12. Naitoh I, Okayama Y, Hirai M, Kitajima Y, Hayashi K, Okamoto T, Akita S, Gotot K, Mizusima M, Sano H, Ohara H, Nomura T, Joh T, Yokoyama Y, Itoh M. Exophytic pedunculated gastrointestinal stromal tumor with remarkable cyctic change. *J Gastroenterol* 2003; 38: 1181-1184 [PMID: 14714258 DOI: 10.1007/s00535-003-1228-2]
13. Okano H, Tochio T, Suga D, Kumazawa H, Isono Y, Tanaka H, Matsuasaki S, Sase T, Saito T, Mukai K, Nishimura A, Baba Y, Murata T. A case of a stomach gastrointestinal stromal tumor with extremely predominant cyctic formation. *Clin J Gastroenterol* 2015; 8: 197-201 [PMID: 26112771 DOI: 10.1007/s12328-015-0577-8]
14. Hirooka Y, Itoh A, Kawashima H, Ohno E, Itoh Y, Nakamura Y, Hiramatsu T, Sugimoto H, Sumi H, Hayashi D, Ohnini Y, Miyahara R, Nakamura M, Funasaka K, Ishigami M, Katano Y, Goto H. Contrast-enhanced endoscopic ultrasoundography in digestive diseases. *J Gastroenterol* 2012; 47: 1063-1072 [PMID: 23001249 DOI: 10.1007/s00535-012-0662-4]
15. Kitano M, Sakamoto H, Kudo M. Contrast-enhanced endoscopic ultrasound. *Dig Endosc* 2014; 26 Suppl 1: 79-85 [PMID: 24118242 DOI: 10.1111/den.12179]
16. Mekky MA, Yamao K, Sawaki A, Mizuno N, Hara K, Nafeh MA, Osman AM, Koshikawa T, Yatabe Y, Bhatia V. Diagnostic utility of EUS-guided FNA in patients with gastric submucosal tumors. *Gastrointest Endosc* 2010; 71: 913-919 [PMID: 20226456 DOI: 10.1016/j.gie.2009.11.044]
17. Hoteya S, Izuza T, Kiduchi D, Yahagi N. Endoscopic submucosal dissection for gastric submucosal tumor, endoscopic sub-tumoral dissection. *Dig Endosc* 2009; 21: 266-269 [PMID: 19961528 DOI: 10.1111/j.1443-1661.2009.00905.x]

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