Case Report

Multiple Gastrointestinal Stromal Tumors in Neurofibromatosis Type 1 Treated with Laparoscopic Surgery

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Gastrointestinal stromal tumors (GISTs) have been reported to occasionally occur in patients with neurofibromatosis type 1 (NF-1), and many cases have had multiple lesions predominantly involving the small intestine. We report herein a case of multiple GISTs associated with NF-1 from whom laparoscopic surgery was beneficial. In a 79-year-old female admitted with anemia and melena, the abdominal computed tomography revealed a tumor arising from the small intestine. Laparoscopic surgery was performed, and another small tumor was revealed during laparoscopic observation. Extracorporeal partial and wedge resection of the small intestine were undertaken. Both lesions were diagnosed as typical GISTs of low risk. Laparoscopic surgery would be useful for examination and a minimally invasive approach to tumors of the small intestine, especially on cases with the possibility of multiple tumors.

Key words: gastrointestinal stromal tumor, neurofibromatosis type 1, laparoscopic surgery

Gastrointestinal stromal tumors (GISTs) are mesenchymal tumors of the gastrointestinal tract arising from the Cajal’s cells expressing c-kit [1]. GISTs commonly occur sporadically but have been reported to occur with increased tendency in patients with neurofibromatosis type 1 (NF-1). NF-1, also known as von Recklinghausen disease, is one of the most common human single gene disorders, affecting at least 1 million persons throughout the world. It encompasses a spectrum of multifaceted disorders and may present with a wide range of clinical manifestations, including abnormalities of the skin, nervous tissue, bones, and soft tissues. The clinicopathological characteristics of GISTs in NF-1 are different from those of sporadic cases; most cases show multiple GISTs predominantly involving the small intestine, and mutations of c-kit or platelet-derived growth factor receptor A (PDGFRα) are rare [2–7]. Those multiple tumors are sometimes problematic for surgical treatments.

Laparoscopic surgery is a minimally invasive approach to common surgical problems in the abdomen that results in much less surgical trauma and postoperative pain. Another advantage of laparoscopic surgery is providing a magnified view of the patient’s internal organs on a television monitor, which helps with precise observation of the whole abdominal cavity. We present herein a case of multiple GISTs of the small intestine in which laparoscopic surgery was useful for examination and treat-

Received August 21, 2006; accepted October 24, 2006.
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Case Report

A 79-year-old female patient, with known NF-1, felt general fatigue and consulted a physician. Severe anemia and melena were found and the patient was referred to our institution for the exploration of gastrointestinal bleeding. The patient presented café-au-lait spots over her whole body, but no abdominal mass or tenderness was noted. Laboratory analysis showed severe anemia with a hemoglobin count of 4.8 g/dl (Table 1). Upper gastroendoscopic and colonoscopic examinations were negative. Computed tomography revealed a round, approximately 3.0 cm × 2.5 cm, lower abdominal mass that seemed to arise from the small intestine (Fig. 1). The tumor was suspected to be a GIST and the cause of gastrointestinal bleeding. After blood transfusion, laparoscopic surgery was undertaken.

A laparoscope was inserted through a 10-mm cannula on the left side of the navel. Two additional 5-mm side ports were inserted to manipulate the intestine. Laparoscopy clearly located the main tumor protruding from the ileum and showed another 5 mm × 4 mm tumor of the terminal ileum (Fig. 2). There was no sign of peritoneal dissemination. By lengthening the cannula incision, extracorporeal partial and wedge resection of small intestine were undertaken. The main tumor had ulcer formation on the mucosa (Fig. 3). Upon histological examination, both lesions were diagnosed as typical GISTs, with

<table>
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<th>Table 1 Preoperative data of blood test</th>
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<tr>
<td>WBC 6900 /mm³</td>
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<tr>
<td>RBC 216 × 10⁴ /mm³</td>
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<tr>
<td>Hb 4.8 g/dl</td>
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<tr>
<td>Ht 16.4 %</td>
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<tr>
<td>Plt 33.0 × 10⁴ /mm³</td>
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<tr>
<td>T.P 5.8 g/dl</td>
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<tr>
<td>Alb 3.6 g/dl</td>
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<td>AST 18 IU/l</td>
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<td>ALT 9 IU/l</td>
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<td>CRP 0.3 mg/dl</td>
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<td>ALP 240 IU/l</td>
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Fig. 1 The contrast-enhanced computed tomography revealed a lower abdominal tumor arising from the small intestine (arrow).

Fig. 2 A, Laparoscopic view of the main tumor, which protrudes from the ileum; B, Laparoscopic view of a smaller tumor of the terminal ileum.
a low risk of malignancy and CD117 (c-kit)-and CD34-positive. The patient had an uneventful post-operative course and was discharged on the 12th day following the surgery. No adjuvant treatment was performed, and there had been no evidence of recurrence at 1 year.

**Discussion**

Laparoscopic surgery is a minimally invasive approach to abdominal problems, including cancers, obesity, functional diseases of the gut, and parietal defects, carried out through small openings while watching the internal organs on a television monitor. Less surgical trauma results in less postoperative pain, a shorter hospital stay, a faster return to a solid food diet, and a quicker return of bowel function. Another advantage of laparoscopic surgery is providing a magnified view of the patient’s internal organs, which allows better visualization of the operative site for more precise work [8]. Nguyen et al. have attempted laparoscopic surgery for GISTs, and most tumors were successfully removed laparoscopically with intracorporeal or extracorporeal anastomoses [9]. In their study, the conversion to open surgery was associated with coincidental pathology in addition to the GIST. Recent consensus reports have stated that GISTs are of uncertain malignant potential, and that all tumors should be resected despite an uncertain diagnosis. Therefore, laparoscopic surgery is advantageous for observation and treatment for tumors in the small intestine. However, laparoscopic procedures also have several limitations. The surgeon should be careful not to destroy the capsule of the GIST because destruction of GIST may cause peritoneal dissemination. It is important to hold the adjacent normal intestine but to avoid directly holding the tumor with laparoscopic devices. The limited observation of stomach and colon, especially their posterior walls, may explain why multiple lesions are sometimes overlooked. We chose laparoscopic surgery in the present case because the patient had undergone upper gastroendoscopic and colonoscopy examinations, and the issue of limited observation also exists in open surgery.

18F-Fluorodeoxyglucose (FDG)-positron emission tomography (PET) is a useful technique for assessing GIST, and the FDG uptake is considered to predict the malignant potential of GIST because of the correlation between FDG uptake and both the Ki67 index and the mitotic index [10]. In addition, combined PET and CT offers several advantages over PET alone, and the use of this technique has spread widely over the past few years [11]. Some studies have described PET/CT imaging of GISTs [12]. The sensitivity has been found to be 93% for CT and 86% for PET for the diagnosis of GIST, and a false negative on PET scan appears to be related to small lesions. The minimum lesion detected has been reported to be 0.4 cm, which is almost the same size as our smaller tumor. PET/CT is often used to monitor the response to imatinib with malignant GIST, but our case also appeared to be a good candidate for follow-up with PET/CT scan, because the patients with NF-1 tend to have multiple GISTs.

The phenotypic and genotypic characteristics of GIST in NF-1 patients have been described recently. A great majority of tumors occur in the jejunum or ileum, and most cases are multi-focus. Though most tumors are stained for CD117 and CD34 and are morphologically indistinguishable from sporadic GISTs, few GISTs with NF-1 have a c-kit mutation or PDGFRα mutation, which is typically seen in sporadic GISTs [2-6, 13, 14]. These characteristics indicate that GISTs in NF-1 patients have a different pathogenesis than sporadic GISTs. The genetic disorder of NF-1 is a germ line mutations of NF1 which encodes neurofibromin and functions as a negative

![Fig. 3](image-url)  
**Fig. 3** Macroscopically appearances of large intestinal GIST involving the intestinal wall and small GIST protruding to the serosa.
regulator of Ras activity. Additional somatic inactivation of the wild-type NF1 allele has recently been shown in GISTs; the homozygous inactivation of NF1 may therefore favor GIST formation [15]. Because of these genetic disorders, patients with NF-1 are estimated to develop GISTs at a rate at least 45 times greater than sporadic cases, but most GIST patients with NF-1 have revealed a good prognosis with long-term follow up [2].

In conclusion, we have described a NF-1 patient with multiple GISTs for whom laparoscopic surgery was useful for finding another small lesion. Laparoscopic surgery appears to be useful for examination and a minimally invasive approach to tumors of the small intestine, especially in cases with the possibility of multiple tumors.

References