Supraclavicular lymph node metastases from malignant gastrointestinal stromal tumor of the jejunum: A case report with review of the literature

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Abstract

Gastrointestinal stromal tumors (GISTs) represent the most common mesenchymal tumors of the alimentary tract. These tumors may have different clinical and biological behaviors. Malignant forms usually spread via a hematogenous route, and lymph node metastases rarely occur. Herein, we report a patient with a jejunal GIST who developed supraclavicular lymph node metastasis. We conclude that lymphatic diffusion via the mediastinal lymphatic station to the supraclavicular lymph nodes can be a potential metastatic route for GISTs.

Key words: Gastrointestinal stromal tumor; Metastasis; Lymph nodes

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Core tip: Unlike gastrointestinal carcinomas, lymph node metastases rarely develop in patients with malignant gastrointestinal stromal tumors (GISTs). We
report a patient with a jejunal GIST who developed supraclavicular lymph nodes metastasis and review the related literature. We conclude that lymphatic diffusion via mediastinal lymphatic station to the supraclavicular lymph nodes can be a potential metastatic route of GISTs.

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INTRODUCTION
Gastrointestinal stromal tumors (GISTs) represent the most common mesenchymal tumors of the alimentary tract. These tumors may have different clinical and biological behavior. Malignant forms usually spread via hematogenous route, and lymph nodes metastases rarely occur. Herein, we report a patient with jejunal GIST who developed supraclavicular lymph nodes metastasis.

CASE REPORT
A 56-year-old man with a 24-h history of melena was admitted to the gastrointestinal department of the Yuhuangding Hospital affiliated to Qingdao University, China. Laboratory assessment revealed a hemoglobin level of 8 g/dL. Urgent gastroduodenoscopy, colonoscopy, and enhanced computed tomography (CT) did not reveal any source of bleeding. During the next 24 h, the patient had further episodes of melena and became hemodynamically unstable after receiving a transfusion of 5 units of packed red blood cells and hemostatic agents. Therefore, emergency laparotomy was performed. Upon surgical exploration, a bleeding solid mass was found in the jejunum. Resection of a small bowel loop measuring 20 cm in length was performed. Gross examination revealed a nodular well-encapsulated tumor measuring 2 cm. Histologic sections showed a GIST infiltrating through all bowel layers, and it had features of mixed spindle and epithelioid types of cells. The mitotic index was > 5/50 high-power fields (HPFs). There was no infiltration of the surgical margins, and two harvested lymph nodes were free of metastases (Figure 1). CD117, CD34, and Dog-1 were positive in immunohistochemical studies; the Ki67 index was 20%. A diagnosis of GIST of the small intestine with high-grade malignancy was established based on the modified NIH GIST criteria[1]. Postoperative total body CT scan and positron emission tomography-computed tomography (PET-CT) were negative for metastatic disease. Adjuvant imatinib therapy was prescribed, but he declined it due to family and economic reasons.

One year later, the patient was admitted to the thyroid department at the same hospital complaining of a left cervical mass that had been gradually enlarging over one month. Ultrasound revealed a hypoechoic mass above the left clavicle, measuring 3.1 cm x 4.6 cm; this mass was unenhanced in a contrast-enhanced CT scan (Figures 2 and 3). Interestingly,
there were no alterations of the thyroid in imaging studies. Core needle biopsy was performed, and histopathological examination with hematoxylin-eosin staining revealed lymph node metastasis from a GIST. In immunohistochemical studies, CD117, CD34, and vim were positive, whereas calponin, estrogen, progesterone, and thyroglobulin were negative. A PET-CT scan showed uneven 18F-fluorodeoxyglucose (FDG) uptake in the cervical mass and multiple lymph nodes in the mediastinum (Figure 4). The patient underwent surgical removal of the cervical mass. Gross examination of the excised lymph node demonstrated that it measured 5 cm x 6 cm x 8 cm and appeared irregular, smooth, and well encapsulated. Histopathologic examination again confirmed a metastasis from a GIST (Figure 5). Immunohistochemical studies of the resected lymph node showed positivity for CD117 and CD34 and negativity for Dog-1 and S-100. The Ki67 index was 30%. Insertion of GCC TAT in exon 9 of the c-KIT gene was identified with mutation analysis. Exons 11, 13, and 17 and the PDGFRα gene were wild type. After the second operation, the patient was regularly given imatinib at a dose of 400 mg per day. At 1 year following the surgery, the patient was asymptomatic, and there were no signs of tumor recurrence or progression (Table 1).

**DISCUSSION**

GISTs represent the most common neoplasms of mesenchymal origin of the gastrointestinal tract. GISTs may have different clinical and biological behavior, ranging from small benign tumors to aggressive forms that have a dismal prognosis. Approximately 20%-25% of GISTs are located in the stomach, and 40%-50% of those located in the small intestine were malignant neoplasms with features such as local recurrence after surgical removal, intraperitoneal dissemination and distant metastases. However, unlike gastrointestinal carcinomas, lymph node metastases (LNMs) rarely develop in patients with malignant GISTs. The mainstay of treatment for GISTs is complete surgical resection without a regional lymphadenectomy.

The rates of LNMs from GISTs range from 0% to 5%. A few studies reporting on this subject are summarized in Table 1. Most of the reported cases are peritumoral lymph nodes metastases, which have occasionally been discovered with histopathological examination of surgical specimens. We found only 3 cases that could be defined as distant LNMs, including 2 inguinal lymph nodes and 1 axillary lymph node. In our case, LNMs developed in the left supraclavicular and mediastinal basins. This behavior is similar to that observed in malignant gastrointestinal tumors of an epithelial origin. To the best of our knowledge, this is the first case report of the lymphatic spread of a gastric GIST to supraclavicular and mediastinum lymph nodes. This might indicate that a particular subgroup of GISTs has biological characteristics similar to carcinomas.

According to the modified NIH GIST criteria, our case was a high-grade malignancy. Ki67 expression changed from 20% in the primary tumor to 30% in the supraclavicular metastasis. Interestingly, Dog-1
was negative in the LNM, whereas it was positive in primary tumor. It could be speculated that the Ki67 and Dog-1 levels may be markers of a primary tumor de-differentiation tendency.

Activating mutations of the c-kit gene (especially exons 11 and 9) are present in most GISTs and probably play a fundamental role in the development of these tumors. Among the reported cases of LNsMs from GISTs, few gene detection results have been described, most of which are exon 11 mutations\cite{5,7,13,14}. In the study by Kong et al\cite{13}, the exon 11 mutation was linked to the likelihood of LNsMs. However, in our case, we found an exon 9 mutation. This genetic mutation in the LNsMs from GISTs has not been reported to date. The relationship between gene mutations and LNsMs is not clear, but many authors have stated that KIT exon 9-mutant tumors developed imatinib resistance more frequently than exon 11-mutant tumors\cite{20,21}. Cases with exon 9-mutant tumors should be treated with increased imatinib doses. Because the patient declined imatinib treatment after the first surgery, he was treated with 400 mg of imatinib per day after the second operation.

In conclusion, complete surgical resection remains the mainstay of treatment for resectable GISTs. Imatinib is currently indicated for the first-line treatment of patients with metastatic or unresectable KIT-positive GISTs. Adjuvant therapy with imatinib was deemed necessary for this patient following complete resection of a primary jejunal tumor because it was an aggressive, high-risk form of GIST. Unfortunately, he did not take imatinib after his first operation, and distant lymph node metastases occurred after 12 months. Following the second operation, the patient received imatinib treatment and had survived without disease progression at the 1-year follow up.

This case confirms that LNsMs in the mediastinum and supraclavicular lymph nodes is a potential metastatic route for malignant GISTs. Further studies are needed to clarify the mechanism of lymph node metastases in patients with GISTs.

### COMMENTS

**Case characteristics**

The patient was admitted to hospital, complaining of a left cervical mass, which had been diagnosed as a jejunal gastrointestinal stromal tumor (GIST) and cured by surgery 1 year before.

**Clinical diagnosis**

For the differential diagnoses of thyroid tumor, lymphoma, or metastatic carcinoma, the patient underwent computed tomography (CT), ultrasonound (US), positron emission tomography-computed tomography (PET-CT) and biopsy. US, CT and PET-CT revealed a hypoechoic, unenhanced and uneven FDG uptake mass above the left clavicle measuring 3.1 cm × 4.6 cm.

**Laboratory diagnosis**

After a biopsy of the cervical mass, this patient was diagnosed as having supraclavicular lymph node metastases from GISTs.
Pathological diagnosis
Core needle biopsy was carried out, and the histopathological examination using hematoxylin-eosin stain showed lymph node metastasis from GIST.

Treatment
The patient underwent surgical removal of the cervical mass and was regularly given imatinib 400 mg per day after the second operation.

Experiences and lessons
This case confirms that LNM in the mediastinum and supraclavicular lymph nodes are a potential metastatic route of malignant GISTs. Physicians should be aware of this during operation and chemotherapy. In this case report, we tried to give some but not sufficient evidence of the possible mechanisms of the supraclavicular lymph node metastasis.

Peer-review
This case report is well organized and had much information including genetic analysis data on primary GIST and metastatic lesion.

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