

Case Reports

Primary hepatic gastrointestinal stromal tumor

Introduction

Gastrointestinal stromal tumors (GIST) are uncommon, representing 0.1-3.0% of all gastrointestinal cancers. A primary GIST in the liver is even less frequent. We report a primary, malignant, hepatic gastrointestinal stromal tumor that was resected from the liver of a 41-year-old woman.

Case report

A 41-year-old female patient without any comorbidity, presented with a seven month history of pain in her right hypochondrium and right lower chest which was gradual in onset, severe, non-colicky and radiated to her back. It was not related to food intake and usually relieved with analgesics. She had lost approximately 6 kg bodyweight over the last 6 months and had anorexia since 3 months. Her medical history included an open cholecystectomy 16 years ago and abdominal tuberculosis 26 years back for which she completed a course of anti-tubercular therapy.

An ultrasound examination of her abdomen revealed a 6 × 5 cm mass in the right lobe of liver and fine needle aspiration cytology suggested a possible malignancy. Her serum alpha-fetoprotein was 23.3 ng/ml, and her viral markers were non-reactive. A triphasic CT angiography of the abdomen (**Figure 1**), revealed a 15 × 12 cm, heterogeneous mass lesion in the right lobe of liver, involving segments VI and VII, with cystic degeneration. The mass was compressing the right hepatic vein. There was a small satellite nodule in segment V.

Intra-operative findings revealed a large lesion involving segments VI and VII of the liver with central necrosis and a cystic component. There was a 4 × 3 cm lesion as well in segment V without any free fluid, peritoneal nodules or pelvic deposits. The patient underwent a right hepatectomy. Her post-operative course was uneventful and she was discharged four days after surgery. Histopathology (**Figures 2**) showed oval to spindle shaped tumor cells with vesicular, elongated nuclei and eosinophilic cytoplasm. These were arranged in sheets,

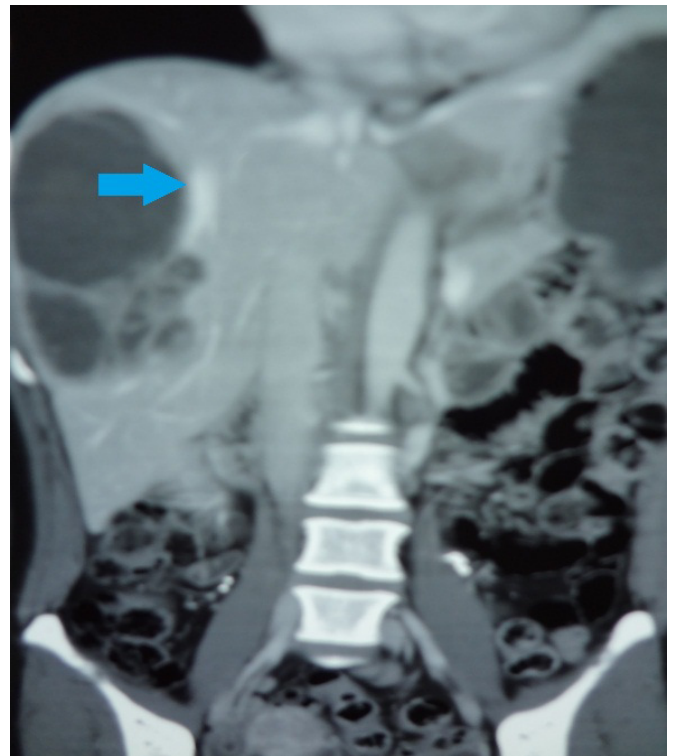


Figure 1: Triphasic CT angiography abdomen showing the tumor mass compressing the right hepatic vein (blue arrow).

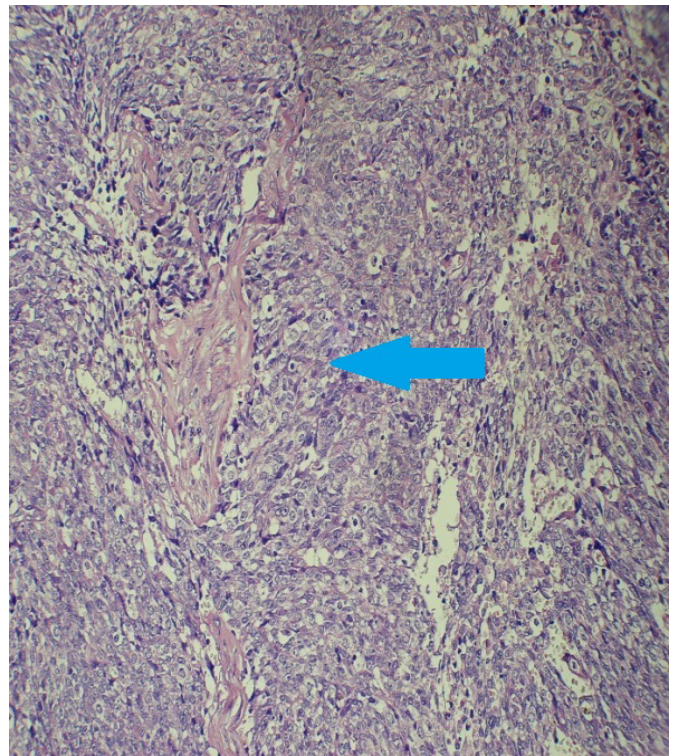


Figure 2: Histopathology depicting oval to spindle shaped tumor cells (blue arrow) with vesicular, elongated nuclei and eosinophilic cytoplasm. The cells are arranged in sheets, bundles and fascicles separated by interspersed fibrous septa.

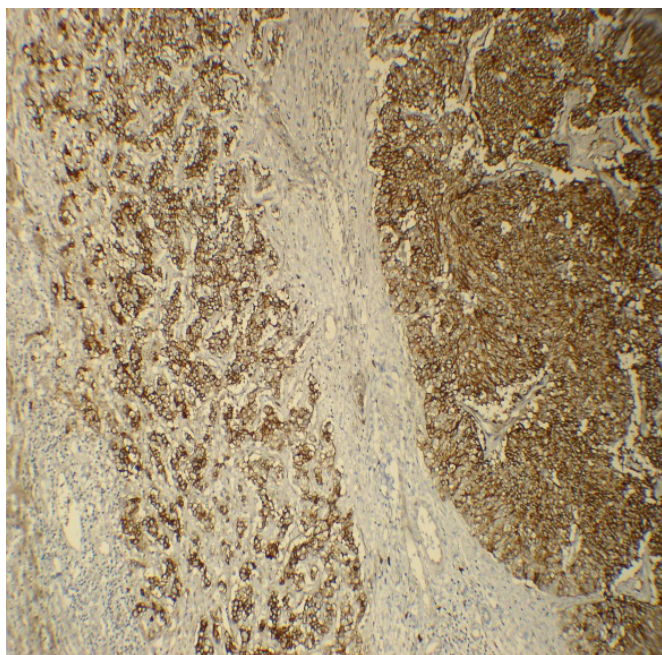


Figure 3: Immunohistochemistry showing tumor cells strongly positive for CD117 and focally for SMA.

bundles and fascicles separated by fibrous septa at places. Prominent cystic degeneration, areas of myxoid change and necrosis were also found. On immunohistochemistry (**Figure 3**) the tumor cells were strongly positive for CD117 and focally for SMA, but were negative for Hep-par 1 and vimentin. She was started on imatinib, but had to be stopped due to its side effects. At 5-month follow-up, the patient is doing well without any recurrence.

Discussion

Gastrointestinal stromal tumors are the most common mesenchymal neoplasms of the GI tract. They are defined as spindle cell, epithelioid or occasionally pleomorphic, mesenchymal tumors that arise from the gastrointestinal tract and usually express the KIT protein. Stomach (40–60%) and small intestine (30–40%) are the most common sites of affliction. The colon, rectum, oesophagus, appendix,¹ and rarely the mesentery and retroperitoneum are other sites of origin. Very few cases of hepatic GIST have been reported. Regardless of where they occur, almost all GIST have the same biological behavior marked by expression of c-kit (CD117) protein. C-kit is a transmembrane growth factor receptor with tyrosine kinase activity, which helps differentiate these tumors from other mesenchymal tumors. Surgery is the best and only curative option. GIST rarely metastasize to the lymph nodes, so formal lymphadenectomy is not necessary. There are only three case

reports of hepatic GIST in the literature, of which two underwent hepatectomy and one radiofrequency ablation therapy.²⁻⁴

TEJAL BHOY
SHAIENDRA LALWANI
JITENDER MISTRY
VIBHA VARMA
VINAY KUMARAN
SAMIRAN NUNDY
NAIMISH MEHTA

Correspondence: Dr Samiran Nundy
Department of Surgical Gastroenterology and Liver
Transplantation
Sir Ganga Ram Hospital, New Delhi - 110060
Email: snundy@hotmail.com

References

1. Miettinen M, Majidi M, Lasota J. Pathology and diagnostic criteria of gastrointestinal stromal tumors (GISTs): a review. *Eur J Cancer*. 2002;**38**:S39–51.
2. Hu X, Forster J, Damjanov I. Primary malignant gastrointestinal stromal tumor of the liver. *Arch Pathol Lab Med*. 2003;**127**:1606–8.
3. Ochiai T, Sonoyama T, Kikuchi S, Ikoma H, Kubota T, Nakanishi M, et al. Primary large gastrointestinal stromal tumor of the liver: report of a case. *Surg Today*. 2009;**39**:633–6.
4. Luo XL, Liu D, Yang JJ, Zheng MW, Zhang J, Zhou XD. Primary gastrointestinal stromal tumor of the liver: a case report. *World J Gastroenterol*. 2009;**15**:3704–7.

Hepatobiliary actinomycosis masquerading as malignancy

Introduction

Actinomyces are anaerobic gram positive filamentous bacteria known to cause chronic suppurative infections. The sites of infection are classified as cervicofacial, thoracic and abdominopelvic and rarely the central nervous system. Actinomycosis of the hepatobiliary system is rare and the clinical and