**Isolated Splenic Metastasis from Colon Cancer – Case Report and Literature Review**

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**ABSTRACT**

Solitary splenic metastases are very rare and sporadic. There are several explanations for this low incidence of splenic metastasis including anatomical, histological and immunological features of the spleen. In this paper we present a case of 70-year-old man with no history of previous diseases who was first operated under the diagnosis of acute abdomen revealing perforated colon tumor of splenic flexure with no metastases at that time. Left hemicolectomy was performed followed by postoperative complications demanding a subtotal colectomy and ileostomy. Primary tumor was classified as Dukes (Astler-Coller)-C2, T4N1M0. Patient was referred to oncologist and received chemotherapy (5FU, Leucovorin). 5 months later continuity of the gut was performed by ileosygmoanastomosis. 2 years after first surgical procedure, a CT scan and abdominal ultrasound, followed by needle biopsy, showed isolated metastasis in spleen, so splenectomy was performed. Pathological findings revealed sharply bordered, partially necrotic tumor inside of spleen tissue, spreading to, but not reaching splenic hilum. Histology showed low to medium differentiated adenocarcinoma tissue with desmoplastic stromal reaction. There were no protrusions of tumor cells through spleen surface. In splenic hilum 4 tumor free lymph nodes were harvested. No additional chemotherapy was conducted. The latest follow up, a year after diagnosis of metastasis showed no signs of cancer disease. Review of the literature showed that long term survival and prognosis of isolated splenic colorectal metastasis after splenectomy are rather optimistic, although these are the cases of distant metastasis. Due to small number of cases reported in literature, definitive conclusions and/or guidelines for the treatment of isolated splenic metastasis cannot be given, but splenectomy and chemotherapy are preferable in the treatment, promising long term survival at least for metachronous metastasis.

**Key words:** colon cancer, metastasis, spleen

**Introduction**

The incidence of splenic metastasis, due to better medical imaging and long term follow up has been increasing1. In most of the cases, they are part of multivisceral metastatic cancer2. There have been several autopsy studies considering prevalence of splenic metastasis ranging from 2.3 to 7.1%3. Also, there have been several studies considering prevalence in living patients showing 1.3% of 1280 splenic tumors as metastatic4, 9.8% of 122 diagnostic splenectomies were positive for metastasis4 and 1% of 1743 splenectomies were positive for metastasis5.

Majority of the cases are part of multivisceral metastatic diseases and usually originate from breast, lung, ovarian, colorectal and gastric cancer and skin melanoma3,5. Most of them are asymptomatic and diagnosed during regular follow up by ultrasound and CT scan, while PET scan is introduced as tool for revealing more asymptomatic cases1. Solitary metastases are very rare and sporadic. Incidence of colon and rectum metastasis in autopsy study of Berge3 was 4.4 and 1.6% respectively, however no solitary metastases were reported. Pisanu et al.6 reported in their case report and review of the litera-
ture, published in November 2007, only 42 cases of well
documented isolated splenic metastasis of colon and rec-
tum origin. There are several explanations for this low
incidence of splenic metastasis including anatomical, his-
tological and immunological features of the spleen7.

Case Report

In 2005, a 70-year-old man with no history of previous
diseases was referred to emergency department of our
Clinic suffering from abdominal pain, fatigue and vomit-
ing. During clinical examination distension and tender-
ness of abdomen were found and patient was immedi-
ately operated under the diagnosis of acute abdomen.
Laparoscopic exploration was done at first and it re-
vealed approximately 1 L of free intraperitoneal puss so
conversion to open laparotomy was done. After perito-
eal washing, a perforated tumor (approximately 10×5
cm) of splenic flexure was found, infiltrating surround-
ing fat tissue close to pancreas tail. No macroscopic
metastases were found. Left hemicolectomy was per-
formed with end to end hand sewed anastomosis. On
10th postoperative day patient developed signs of acute
abdomen, so relaparotomy was done revealing anasto-
ometic leakage. Since the rest of the colon was distended,
a subtotal colectomy and ileostomy was performed. Two
weeks after the second operation patient was discharged
from hospital.

Pathological finding revealed ulcero-infiltrative cir-
cumferential tumor, whose dimension was 11.5×4.5 cm.
Histology analyses showed a moderately differentiated
adenocarcinoma with high mitotic and apoptotic pattern
together with areas of necrosis. Tumor invaded visceral
peritoneum and surrounding fat tissue. Five lymph no-
des were harvested and two of them were positive for me-
tastasis. Resection margins of colon and mesocolon were
free of tumor. Overall, primary tumor was classified as
Dukes-C, Dukes (Astler-Coller)-C2, T4 N1 M0, stage
IIIB, according to TNM system and histological grade –
G2.

Following discharge from surgery department patient
was referred to oncologist and received chemotherapy
(5FU, Leucovorin). Four months after the initial exami-
nation, patient was admitted to our Clinic and continuity
of the gut was performed by ileosygmoanastomosis. The
free edge of sigmoid colon was resected during surgery
and histopathology did not reveal signs of colon cancer.
The patient was regularly followed up, having no symp-
toms of the disease. A scheduled CT scan in May 2007
showed hypovascular lesion of the spleen, 7.9 cm in di-
ameter (Figure 1). Other findings were normal except for
gallbladder stone. CEA level was 5.05 µg/L (ref. <3.4
µg/L). Abdominal ultrasound examination also showed
hypovascular lesion (Figure 2). In July 2007 an ultra-
sound guided needle biopsy was performed and cytology
analyses revealed adenocarcinoma cells. Patient was
again admitted to our Clinic and scheduled for surgery.
Laparotomy was done and exploration revealed spleen
with tumor mass inside with no protrusion through
spleen tissue. There were no signs of other metastatic le-
sions. Splenectomy was performed. Pathology findings
after splenectomy revealed an enlarged spleen (13×10×5
cm) (Figure 3a and 3b) with sharply bordered, partially
necrotic tumor (8×7×3 cm) inside of spleen tissue, spread-
ging to but not reaching splenic hilum (Figure 4). Histol-
ogy examination showed low to medium differentiated
adenocarcinoma tissue with desmoplastic stromal reac-
tion. There were no protrusions of tumor cells through
spleen surface. In splenic hilum four lymph nodes were
harvested, all without carcinoma cells.

Early postoperative myocardial infarction developed
and stenting of left anterior descending coronary artery
was performed. A week later a cerebrovascular infarction
developed with dysphasia and mild right hemiparesis.

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Fig. 1. Abdominal CT scan showing hypovascular splenic lesion.

Fig. 2. Abdominal ultrasound showing hypovascular splenic
lesion.
There were no surgical complications. In August 2007 patient was discharged from hospital and referred to specialized institution for further rehabilitation. No additional chemotherapy was conducted. The latest follow up in May 2008 showed no signs of cancer disease.

Discussion

As mentioned before, splenic metastasis of various cancers are very rare and usually part of multivisceral dissemination and isolated metastases are sporadic. There are several theories to explain this rarity and two main are: first including mechanical/anatomical and histological factors preventing implantation of blood borne cancer cells. These factors include constant flow of blood through spleen and rhythmic contractions of splenic capsule and splenic sinusoidal architecture, sharp angle of splenic and celiac artery preventing clamps of tumor cells from passing through and lack of afferent lymphatic vessels limiting lymphogenic metastases⁷,⁸. According to Indudhara⁹ neoplastic cells can reach splenic vein and parenchyma by retrograde diffusion through the inferior mesenteric vein. The spleen parenchyma has no lymphatic vessels but they are present in capsular, subcapsular and trabecular regions⁸. Tumor cells might also reach the spleen through lymphatic system which explains the subcapsular localization of isolated splenic metastasis⁸. The second theory includes immunology factors and that is inhibitory effect of splenic microenvironment on the growth of cancer cells⁷,⁸ as the spleen is the second largest organ of lymphoreticular endothelial system, so immune surveillance appears to potentially inhibit tumor cell proliferation¹⁰. However, recent studies based on sensitive immunologic and molecular methods that can detect single cells showed that micrometastases can be detected at the time of tumor diagnosis¹¹,¹². Considering this, it could be assumed that implantation of cancer cells can occur but further growth is inhibited by microenvironment, explaining high prevalence of spleen micrometastasis found at autopsy and low prevalence of clinically detectable metastases².

The prevalence of splenic metastasis, although very low, is increasing with the improvement of imaging techniques². About 20% of colorectal carcinomas are metastatic at their clinical presentation¹³. The usual sites of metastasis are liver, lung and axial skeleton¹⁴,¹⁵. Microscopic splenic metastases were found in 7–34% of cancer patients⁴. In the same study, incidence of splenic colorectal micrometastasis was reported as 2% of 1019 colorectal tumors but all of these cases involved other organs as well³. In his article, Pisanu reported that up to his case review only 41 cases of isolated colorectal splenic metastasis have been reported⁶ and most of the cases were metachronous.

Majority of cases reported had a disease free survival period of 3–144 months after the diagnosis of primary tumor¹⁶–¹⁸. Long term survival after splenectomy in patients with metachronous splenic metastasis was 0.5–7 years¹⁷–¹⁹. So, prognosis of isolated splenic colorectal metastasis is rather optimistic, although these are the cases of distant metastasis¹⁰. For synchronous metastasis much
worse prognosis and shorter disease free survival has been reported\textsuperscript{18}.

Although due to small number of cases reported in literature, definitive conclusions and/or guidelines for the treatment of isolated splenic metastasis can not be given, splenectomy and chemotherapy are preferable in treatment, promising long term survival, at least for metachronous metastasis.

REFERENCES


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