Right side fixation of sigmoid colon with a hepato-sigmoidocolic fistula in patient with hepatocellular carcinoma and midgut malrotation

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The location of the sigmoid colon varies within the abdominal cavity, but its mesocolon is fixed to the left side. Right side fixation of the sigmoid colon is a very rare congenital positional anomaly. In addition, it has been reported that hepatocolic fistula is also a very rare disease that may present lower gastrointestinal bleeding. Here, the authors describe a case of a 71-year-old man who underwent surgery for hepato-sigmoidocolic fistula complicated by hepatocellular carcinoma and the right side fixation of the sigmoid colon.

Key Words: Right side fixation of sigmoid colon, Hepato-sigmoidocolic fistula, Hepatocellular carcinoma, Midgut malrotation

INTRODUCTION

From the embryonic period, most of the sigmoid colon is located in the left lower abdomen and some in the right. Regardless of its location, mesocolon of the sigmoid is always fixed to the peritoneum in the left side. Right side fixation of the sigmoid colon is a rare congenital malposition probably associated with midgut malrotation or extra-long sigmoid colon [1,2].

Hepatocolic fistula with lower gastrointestinal (GI) bleeding is a rare condition that is reported to occur by hydatid cyst or hepatocellular carcinoma [3-5]. Anatomically, if liver disease invades the colon to develop hepatocolic fistula, it may affect the ascending colon or hepatic flexure of the colon. However, in this case, right side fixation of the sigmoid colon developed a hepato-sigmoidocolic fistula. To our knowledge, this is the first successfully treated case of hepato-sigmoidocolic fistula complicated by hepatocellular carcinoma and concomitant cirrhosis.

CASE REPORT

The case was a 71-year-old male with a history of alcoholic liver cirrhosis. After being diagnosed with hep-
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atocellular carcinoma 6 years ago, he was treated with transarterial chemoembolization (TACE) 7 times at other hospital. He visited our emergency room with lower GI bleeding. At the time of admission, the level of hemoglobin (Hb) was 7.5 g/dL, aspartate aminotransferase/alanine aminotransferase 27.8/13.8 U/L, alkaline phosphatase 41 U/L, albumin 2.8 g/dL, a-fetoprotein 26 ng/mL, and bilirubin 1.6 mg/dL. After he stabilized with transfusion of packed red blood cell (PRC), emergency colonoscopy was performed. A solitary diverticular-like lesion or ulceration with a brownish base in the sigmoid colon, 25 cm from the anal verge, was detected on colonoscopy, but the evidence of active bleeding was not shown (Fig. 1). Endoscopic clips were applied at the lesion to prevent recurrent bleeding. After bleeding was controlled, an abdominopelvic computerized scan was checked for evaluation of liver mass and other abnormalities (Fig. 2). The symptom improved and he was discharged 3 days later with plans to follow up using colonoscopy 2 months after. However, lower GI bleeding recurred after 7 weeks and the patient was admitted again. At the time of admission, Hb was 7.8 g/dL and PRC transfusion was given to stabilize the patient. The patient was diagnosed with recurrent sigmoid diverticular bleeding and was scheduled for laparoscopic segmental resection of the involved lesion of sigmoid colon. His pre-operative Child-Pugh classification was grade A with serum albumin 3.3 g/dL, serum bilirubin 0.94 mg/dL, prothrombin time 12.3 seconds, no ascites, and no encephalo-
finding was consistent with midgut malrotation (Fig. 3).

Under laparoscopic operative condition, the operator had to move the position from the right side to the left side of the patient, and the right side fixation of the sigmoid colon was dissected. Laparoscopic surgery was limited due to serious adhesion of proximal sigmoid colon to the liver and thus converted to laparotomy with upper midline incision. The upper fixation area of the sigmoid colon could be mobilized with concomitant en bloc nonanatomical excision of the liver (segment 6) containing hepatocellular carcinoma. After complete mobilization of the sigmoid colon with a part of the liver, segmental resection of the sigmoid colon was performed with about 10 cm resection margins proximally and distally, respectively. After surgery, the patient was treated for ascites and hydrothorax for about 20 days, which were free of malignant cells, and recovered without additional complications. Gross findings of the specimen showed a perforation hole between the sigmoid colon and liver, and hepatocellular
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carcinoma tissue. The microscopic findings revealed a 4.5 cm × 5 cm sized metastatic hepatocellular carcinoma invading the sigmoid colon. The central part of the tumor presented with necrosis with the peripheral part constituting intact cancer cells, suggesting marginal recurrence at the previous tumor site after TACE (Fig. 4).

DISCUSSION

In this case, besides the fixation anomaly of the sigmoid colon, midgut malrotation was considered a congenital anomaly. So far, only a couple of cases of right side fixation of the sigmoid colon have been reported [6-9]. Four or 5 cases were detected during autopsy, and another case was detected during X-ray examination. To our knowledge, our case is the first detected during the actual surgery. In most cases, it was considered to occur congenitally, since the sigmoid colon was excessively long. In one case that was detected during X-ray examination, it was reported to have occurred due to midgut malrotation [8]. It may be inferred from such evidence when midgut malrotation is developed, the cecum and the ascending colon float freely within the abdominal cavity, similar to the small bowel. Consequently, the empty space in the right side is occupied by the redundant sigmoid colon, and the fixation between the mesocolon of the sigmoid colon and peritoneum of the right side may occur during the later period of the fetus; and thus, such congenital malposition is developed. Therefore, summarizing the cases reported until now demonstrates that the cause of right side fixation of the sigmoid colon may be as a result of extra-long redundancy of sigmoid colon and/or midgut malrotation.

In the era of laparoscopic colorectal surgery, if the right side fixation of sigmoid colon is encountered during surgery, it may prove problematic, such that the operator position would need to be changed to the left side of the patient or an additional trochar may be inserted in the left side of abdomen.

The colon is located quite close to the liver within the abdominal cavity, particularly the ascending colon or the hepatic flexure of the colon. Despite this anatomical proximity, hepatocolic fistula is extremely rare. Until now, only 3 cases of hepatocolic fistula have been reported. Two of which were caused by hydatid disease [4,5], the other by an invasion of hepatocellular carcinoma [3], which corresponds to this case. In that case, the patient with a history of alcohol abuse showed lower GI bleeding and was confirmed with hepato-ascending-colic fistula as a result of invasion of hepatocellular carcinoma on the computed tomographic scan, water-soluble enema, colonoscopy and biopsy. The patients died a few weeks later without surgery, but, in our case, even though we didn’t suspect, pre-operatively, the hepatocolic fistula in the patient with lower GI bleeding, the cirrhotic patient with hepatocellular carcinoma was treated with operation. The pre-operative diagnosis was considered as a solitary diverticulum or Dieulafoy’s lesion of the sigmoid colon. In another case, extensive hepatocellular carcinoma infiltrated to the hepatic flexure and induced a massive lower gastrointestinal bleeding, which caused the death of the patient [10]. It has been reported that, in hepatocellular carcinoma patients, gastrointestinal bleeding occurs as the hepatocellular carcinoma metastasize or directly infiltrate to the stomach, the duodenum, or rarely to the colon.

In summary, we report a combination case of congenital right side fixation of the sigmoid colon and hepatosigmoidocolic fistula in a patient with hepatocellular carcinoma and cirrhosis, which was treated with surgery successfully. And when a patient with hepatocellular carcinoma develops lower GI bleeding, the possibility of hepatocolic fistula should be considered.

CONFLICTS OF INTEREST

No potential conflict of interest relevant to this article was reported.

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