CASE REPORT

SORAFENIB, RISK OF BLEEDING AND SPONTANEOUS RUPTURE OF HEPATOCELLULAR CARCINOMA. A CLINICAL CASE

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Summary: Spontaneous rupture is a rare and dramatic complication of hepatocellular carcinoma (HCC), burdened by a high mortality. Here we describe a case of a 73-year-old man, who arrived at the ER because of syncope, and acute epigastric and right upper quadrant abdominal pain. He had a history of hepatitis C-related liver cirrhosis and HCC in treatment with sorafenib. The physical examination showed a state of hemorrhagic anemia with the presence of blood in the peritoneal cavity. The patient underwent an urgent liver resection. Thirty days after surgery, he was in good general condition. Sorafenib is a multikinase inhibitor recently introduced in the therapy of patients with advanced HCC. Among the various side effects reported in patients treated with sorafenib, there is a higher risk of bleeding. In conclusion, sorafenib may increase the risk of bleeding and rupture of HCC in susceptible individuals.

Key words: Hepatocellular carcinoma; HCC; Rupture; Bleeding; Sorafenib; Side effects

Introduction

Sorafenib is a multikinase inhibitor of the vascular endothelial growth factor (VEGF) pathway recently introduced in the therapy of advanced hepatocellular carcinoma. Treatment with sorafenib is associated with a significant increase in the risk of bleeding. This report describes the case of a large hepatocellular carcinoma, bleeding from a spontaneous rupture, in a patient taking sorafenib.

Case Report

A 73-year-old Caucasian man came to the emergency room because of syncope and acute epigastric and right upper quadrant abdominal pain. He had a history of hepatitis C-related liver cirrhosis and HCC, in treatment with sorafenib 400 mg orally twice daily. On examination, the patient looked pale and sweaty. The temperature was 36.5 °C, blood pressure 90/70 mmHg, heart rate 110 beats/minute. The abdomen was slightly distended, with tenderness and rebound tenderness in epigastrum and right upper quadrant. The hematocrit was 29.2% (normal range 36–54%), hemoglobin 9.2 g/dl, platelets 371,000 mmc, total bilirubin 1.17 mg/dl, prothrombin time 78%, INR 1.16, albumin 2.37 g/dl, creatinine 1.1 mg/dl. Ultrasound-guided diagnostic paracentesis showed blood in the peritoneal cavity. The patient had a previous abdominal CT scan, performed 5 days before, which revealed a large hepatic tumor of 12.8 cm diameter, and corpuscular fluid around the liver (Fig. 2). The patient, with a Child-Pugh score of B7 and a MELD score of 10, underwent to atypical liver resection, including the tumor mass. The postoperative course was regular and the patient was discharged after 9 days. Thirty days after surgery, he was in good general condition.

Discussion

Each year in the world, there are more than 626,000 new cases of HCC. Its incidence is increasing in Europe and the United States of America (1). The spontaneous rupture of HCC is a rare but dramatic complication. Its incidence is around 7–11.2%, according to the geographic area, and is greater in endemic areas for hepatitis B and C, especially in the East (2–4). It is burdened by high mortality, which in 30 days, range from 32 to 67% (2, 5, 6). Risk factors, with increased rate of mortality, are a high Child-Pugh score, high levels of serum bilirubin, hepatic encephalopathy and hypoalbuminemia (2, 6, 7). Therefore, a poor hepatic functional reserve is associated with a poor prognosis. The causes of spontaneous rupture of HCC are unclear. It has been shown that a large tumor, localized at the surface of the liver, is associated with increased risk of rupture (8). Clinical presentations of HCC rupture is the sudden onset of abdominal pain, hypotension and abdominal distension. The next steps in diagnosis are abdominal ultrasound, diagnostic paracentesis and abdominal CT scan (9). The diagnosis of rupture can be difficult, especially in patients with no history of HCC or liver disease (10). In most cases, the sudden onset of abdominal pain, and hemoperitoneum, in the presence of HCC, can lead to strong diagnostic suspicion. At this
point it is necessary to address the therapeutic approaches. In patients with good liver function (Child-Pugh A–B7), the treatment of choice is liver resection. Otherwise, arterial embolization (TAE) (8, 11, 12), is preferable in patients with advanced liver disease, poor functional hepatic reserve, or unresectable HCC. In this case the patient, before the rupture of HCC, had started therapy with sorafenib, at another hospital. Sorafenib is an inhibitor of the vascular endothelial growth factor (VEGF) pathway. In patients with advanced HCC, it causes an increase in survival and time to radiologic progression (13). Treatment with sorafenib is associated with a significant increase in the risk of bleeding (14). The role of VEGF on vessels and coagulation is not yet known. However, we know that VEGF is important for survival of endothelial cells and for maintaining the architecture and integrity of the vasculature (14). Therefore, inhibition of the VEGF pathway may lead to vascular damage and to the inability of vessels to repair after trauma, with consequent increased risk of bleeding (15). Therefore, sorafenib may increase the risk of bleeding and rupture of HCC in susceptible patients.

References

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