INTRODUCTION

Alveolar echinococcosis (AE) is a larval tapeworm infection that occurs in humans after ingestion of foods contaminated with the larval form of *Echinococcus multilocularis*. AE is one of the most dangerous zoonotic diseases in the world. AE most commonly invades the liver and by acting like a malignant neoplasm in the tissue, it leads to invasive and destructive changes. AE rarely causes Budd-Chiari syndrome (BCS) as a result of occlusion of the hepatic veins and inferior vena cava [1,2]. In this paper, we report a case of AE associated with BCS in Turkey.

CASE DESCRIPTION

A 62-year-old female patient admitted to our clinic because of recent fatigue, shortness of breath, abdominal distention, and pain in the right upper quadrant. In her history, she had a liver wedge resection 3 years ago with AE found in the biopsy material. On physical examination, the liver was 5 cm palpable under the right rib, the spleen was non-palpable, and Traube’s space was clear. Tension ascites and collateral veins in an upward flow direction were noted on the abdomen. Respiratory sounds were decreased at the bases of both lungs and dulness was found with percussion.

Hemoglobin was 8.3 g/dl in the whole blood count. Except for low albumin levels (2.1 g/dl), all other biochemical tests were normal. There was no flow viewed in the left hepatic vein and inferior vena cava on abdominal portal Doppler ultrasound. The portal vein was 16 mm, and the splenic vein was 15 mm in diameter and hepatofugal flow was noted. On contrasted abdominal CT, there was a 54 × 70 × 45 mm sized cystic lesion in left lobe of the liver extending towards the posterior mediastinum and invade the diaphragm, esophagus, and pericardium. The cystic lesion was seen to be occluding the inferior vena cava and left hepatic vein at the level where the hepatic veins poured into the inferior vena cava. Bilateral pleural effusion was also detected. We discussed this secondary Budd-Chiari Syndrome (BCS) case, resulting from the AE occlusion of the left hepatic vein and inferior vena cava, in light of the information in literature.

Key words: *Echinococcus multilocularis*, Budd-Chiari syndrome, alveolar echinococcosis, portal hypertension

A Case of Budd-Chiari Syndrome Associated with Alveolar Echinococcosis

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Abstract: Although alveolar echinococcosis (AE) can cause a serious disease with high mortality and morbidity similar to malign neoplasms. A 62-year-old woman admitted to a hospital located in Sivas, Turkey, with the complaints of fatigue and right upper abdominal pain. On contrast abdominal CT, a 54 × 70 × 45 mm sized cystic lesion was detected in the left lobe of the liver that was seen to extend to the posterior mediastinum and invade the diaphragm, esophagus, and pericardium. The cystic lesion was seen to be occluding the inferior vena cava and left hepatic vein at the level where the hepatic veins poured into the inferior vena cava. Bilateral pleural effusion was also detected. We discussed this secondary Budd-Chiari Syndrome (BCS) case, resulting from the AE occlusion of the left hepatic vein and inferior vena cava, in light of the information in literature.

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A 62-year-old female patient admitted to our clinic because of recent fatigue, shortness of breath, abdominal distention, and pain in the right upper quadrant. In her history, she had a liver wedge resection 3 years ago with AE found in the biopsy material. On physical examination, the liver was 5 cm palpable under the right rib, the spleen was non-palpable, and Traube’s space was clear. Tension ascites and collateral veins in an upward flow direction were noted on the abdomen. Respiratory sounds were decreased at the bases of both lungs and dulness was found with percussion.

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character (serum acid albumin gradient was 0.2 g/dl) and protein was 5.1 g/dl. There was no other pathology found in the Gram staining and culture of the intraperitoneal fluid.

According to these findings, the patient was diagnosed with secondary Budd-Chiari Syndrome (BCS) development due to AE. The patient was given 15 mg/kg albendazole (800 mg/day) for treatment.

**DISCUSSION**

BCS develops as a result of occlusion of the venous system at any location from the small veins to the junction of the inferior vena cava at the right atrium. This occlusion may be due to primary or secondary reasons. Primary reasons mostly include hypercoagulopathy disorders. Secondary reasons are a result of intraluminal invasion or extraluminal compression [3,4]. In our case, secondary BCS developed due to both intraluminal invasion and extraluminal compression of the supra-hepatic inferior vena cava and left hepatic vein. Patients with hepatic hydatid cyst, size and posterior localization of the cyst, invasion of 2 or more segments, and prior surgery or infection are the predisposing factors for development of BCS [4]. In our case, most of the predisposing factors were present.

In a large scale survey conducted with 362 AE cases, BCS rates due to inferior vena cava invasion was reported to be approximately 1.8%. In another study, in BCS patients there were about 25% acute and 75% subacute clinical findings. The predominant symptom was found to be right upper quadrant abdominal pain [4,5]. In our patient, the clinical findings had a subacute chronic course, and the predominant symptom was right upper quadrant abdominal pain.

Current treatments have substantially improved the prognosis of AE patients [6]. However, early diagnosis is important for morbidity and mortality. AE generally remains asymptomatic for 5-10 years and later displays a chronic course. In cases that are not treated or insufficiently treated, mortality rates increase. After a diagnosis, a 5-year mortality rate of the disease is 70%, and a 10-year mortality is 94%. When BCS develops, 1-year mortality is 70%, and 3-year mortality is 90% [7]. Our patient’s life expectancy may be shortened due to the metastatic nature of the disease and the late initiation of treatment after surgery.

The treatment of BCS is medical and surgical. For chemotherapy, albendazole 10-15 mg/kg (in 2 divided doses) or alternatively mebendazole 40-50 mg/kg (in 3 divided doses) is used. Surgery is required for cases where BCS has developed. Other interventional methods (stent placing and angioplasty) or liver transplantation along with vena cava resection can be performed for BCS [7]. Kawamura et al. [8] have obtained good outcomes in advanced cases such as secondary BCS due to AE with chemotherapy followed by surgery. In a case report of the liver hudatid cyst with caval involvement, a combination of hepatic lobectomy and vena cava resection was reported to be successful. In disseminated abdominal hydatidoses, long-term chemotherapy reduces the incidence of recurrence and mortality [9,10].

In conclusion, AE is among the most dangerous zoonotic diseases in the world. In recent years, its frequency has increasing due to transmission of the disease from dogs to humans. In AE disease, the liver is the most commonly invaded organ and due to dissemination of surrounding structures, lymphatic and vessels, it mimics malignant neoplasms. AE is a serious cause of morbidity and mortality. As in our patient, AE can in-
vade the hepatic veins and inferior vena cava leading to BCS. In patients that apply to clinics with BCS, AE, although rare, should be considered among the etiological causes.

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