INTRODUCTION

Malaria is a serious, mosquito-borne, infectious disease that is caused by *Plasmodium* species. Among these species, *Plasmodium vivax* is the most common human malaria parasite and endemic in many regions of Asia, the South Pacific, North Africa, the Middle East, and South and Central America [1].

Malaria can present with various signs and complications on computed tomography (CT) including splenomegaly, hepatomegaly, focal low attenuation in the spleen, gallbladder wall edema, periportal edema, ascites and splenic rupture, or hemorrhage [2]. While an enlarged spleen is considered one of the most common signs, splenic rupture rarely occurs as an important life-threatening complication. Therefore, early diagnosis and appropriate disease management are essential.

No well-established studies on the management of the splenic rupture in malaria patients have been performed. Traditionally, non-operative conservative management was performed for the hemodynamically stable patients. However, for the persistently hemodynamically unstable patients, splenectomy has been recommended as a treatment of choice. To the best of our knowledge, a very limited number of malarial splenic rupture patients have been treated with transcatheter coil embolization. However, splenic preservation is more strongly recommended if possible due to the important role of the spleen in immunity. Here we report a Korean *P. vivax* malaria patient with a spontaneous splenic rupture that was successfully managed by embolization of the splenic artery.

CASE RECORD

A 38-year-old South Korean male visited the Emergency Department of Inha University Hospital with a sudden onset of severe abdominal pain and vomiting. He had suffered from intermittent fever during the previous 10 days. His pulse rate was 132 beats per min and his respiratory rate was 22 per min. The patient also had hypotension (95/63 mmHg) as well as high fever (39°C). He had undergone chemotherapy and radiation therapy for orbital marginal zone B-cell lymphoma in October 2010 and 2011 as well as an anterior orbitotomy and mass excision thereafter. One day before visiting the Emergency Department, he underwent an orbital CT and blood test. No abnormal CT findings or blood test results were obtained. In addition, recurrence of the orbital tumor was not observed.

Given the characteristic intermittent fever, a peripheral blood smear and rapid blood assay for malaria were conduct-
Based on the results, the patient was diagnosed with *P. vivax* (Fig. 1). A prompt examination with an abdominal CT revealed that the patient suffered from hemoperitoneum (36 HU on portal phase). The spleen was enlarged and surrounded by a high attenuated fluid collection (61 HU on portal phase; Fig. 2). The surface of the spleen was scalloped due to the fluid accumulation. Higher attenuation of the perisplenic fluid compared to the other site of hemoperitoneum and splenomegaly led to a diagnosis of spontaneous splenic rupture resulting in a subcapsular hematoma.

The patient's hemoglobin level was 14.4 g/dl 1 day before visiting the Emergency Department and decreased to 11.0 g/dl. The concentration then rapidly dropped to 9.5 g/dl, and the patient received a blood transfusion. The patient also had thrombocytopenia (71 k/μl) with 33 IU/L AST, 44 IU/L ALT, 25.4 mg/dl BUN, and 1.66 mg/dl creatinine. He was admitted to the intensive care unit for a conservative treatment. Despite transfusion, the hemoglobin levels and blood pressure remained unstable. On the second day of hospitalization, a splenic angiogram was performed. No definitive extravasation of contrast media was observed, but contour irregularity was noted in the lateral lower margin (Fig. 3). Embolization was performed with 7 coils (10 × 4 mm, 8 × 4 mm, 6 × 2 mm; Cook, Bloomington, Minneapolis, USA) at the distal splenic artery level (Fig. 4).

A follow-up CT was performed 5 days after the coil emboli-
zation. At this time, a reduced splenic subcapsular hematoma was observed. Several peripheral wedge-shaped low attenuated lesions were also found in the spleen, suggesting splenic infarction resulting from the coil embolization (Fig. 5). Hemoglobin levels were normalized (13.9 g/dl). After discharge, the patient has been closely followed-up at the Outpatient Department of Inha University Hospital. No other complications have developed.

**DISCUSSION**

Malaria is a parasitic disease that occurs in over 100 tropical and subtropical countries. Splenomegaly is one of the most common clinical features of malaria, and is estimated to occur in 70-80% of acute cases [3]. A previous review of 22 cases demonstrated that marked splenic enlargement and rupture are most commonly caused by *P. vivax* among the *Plasmodium* species [4]. There is no difference in clinical presentations according to infection by *Plasmodium* species [5].

The pathophysiology of splenic rupture in malaria patients is unclear. However, 3 mechanisms have been suggested [4,6]. First, increased intrasplenic tension caused by cellular hyperplasia and engorgement may result in splenic rupture. Second, the spleen may be compressed by the abdominal musculature during physical activities. Finally, vascular occlusion due to reticuloendothelial hyperplasia may develop, resulting in thrombosis and infarction. This leads to interstitial and subcapsular hemorrhage along with stripping of the capsule, thus leading to further subcapsular hemorrhage.

No well-established studies on the management of the splenic rupture in malaria patients have been performed. Currently, splenectomy has been accepted as the traditional treatment of choice in most cases of splenic rupture. However, perioperative risks associated with splenectomy and growing concerns about the loss of immunologic functions have promoted the development of nonsurgical management strategies [7,8]. These techniques include careful observation in the hospital, strict bed rest, and maintenance of hemodynamics as well as hemoglobin levels with blood and blood products [9]. Hemodynamic instability despite resuscitation efforts indicates nonsurgical management failure.

Osman et al. [8] formulated a flowchart for the management of malarial splenic rupture after reviewing 252 cases previously reported in the literature. This group emphasized the importance of non-operative management of hemodynamically stable patients but recommended splenectomy as a treatment for persistent hemodynamic instability. Angiography with embolization may also be a potential therapy. However, angiographic embolization was omitted from the flowchart because this modality may not be available as a salvage therapy in many clinical settings.

To the best of our knowledge, a very limited number of malarial splenic rupture patients have been treated with transcatheter coil embolization [10]. However, splenic artery embolization has been reported to have a high success rate in trauma patients and preferable to splenectomy [11,12]. Despite the associated risks such as radiation exposure and more time-consuming, or technically challenging, angiographic embolization may be an appropriate option for avoiding harmful effects of splenectomy for malaria patients.

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**CONFLICT OF INTEREST**

We have no conflict of interest related to this report.

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