Case Report: Scrub Typhus and Cerebrovascular Injury: A Phenomenon of Delayed Treatment?

Jong-Hoon Chung,† Na-Ra Yun,‡ Dong-Min Kim,* Ji-Woon Lee, Sung Ho Yoon, and Seok-Won Kim

Department of Internal Medicine, School of Medicine, Department of Neurosurgery, School of Medicine, Gwang-ju, Republic of Korea

Abstract. Three patients diagnosed with scrub typhus through serology and polymerase chain reaction tests, experienced delayed administration of effective antibiotics after the appearance of symptoms, presented with subdural hemorrhage, intracerebral hemorrhage, or cerebral infarction in the late acute phase. Orientia tsutsugamushi should be considered as a causal or provoking factor for cerebrovascular accidents in regions where scrub typhus is endemic, especially in those who receive delayed treatment.

INTRODUCTION

Orientia tsutsugamushi, the causative agent of scrub typhus, infects principally endothelial cells in all organs.1 Acute infection, such as a systemic respiratory tract infection, may temporarily increase the risk of myocardial infarction or stroke.1 This study presents two cases of cerebral hemorrhage (one case of subdural hemorrhage and one case of intra-cerebral hemorrhage) and one case of cerebral infarction that occurred during antibiotic treatment of three scrub typhus patients, in whom the effective antibiotic therapy was relatively delayed after the initial onset of symptoms.

CASE 1

An 81-year-old man was admitted to Chosun University Hospital, a tertiary hospital in Gwangju City, South Korea. He began to go on walks to pick ginkgo nuts at the beginning of autumn. He had a 2-week history of fever and chills, along with a 1-week history of skin rash. He was initially admitted to a local clinic, where he was diagnosed with scrub typhus and given ceftriaxone and doxycycline for 2 days. His mental status deteriorated, and he was transferred to our hospital. He had no significant medical, social, or family history. He presented with a blood pressure of 100/50 mm of Hg, a respiratory rate of 30/min, a pulse rate of 79/min, and a body temperature of 38°C. The peripheral blood tests performed at presentation indicated a white blood cell count of 5,090/mm^3, a hemoglobin level of 10.9 g/dL, and platelets of 62,000/mm^3; fibrinogen, no coagulation; thrombin time (PT), no coagulation; activated partial prothrombin time, 11.4 sec; international normalized ratio (INR), 0.92; aPTT, 28.6 sec; fibrinogen, 257 mg/dL; and D-dimer, 2,466 ng/mL (normal range, 0–5.0 Ug/mL); and D-dimer, 2,466 ng/mL (normal range, 0–255 ng/mL).

A brain computed tomography (CT) was obtained because of the coagulation defects (Figure 1A). An acute hemorrhage, an infarction or structural abnormalities were not found. Blood cultures showed no growth at presentation. An indirect immunofluorescence test for O. tsutsugamushi, performed at presentation, indicated immunoglobulin M (IgM) 1:2,048, and a nested polymerase chain reaction (PCR) targeting the O. tsutsugamushi 56-kDa protein-encoding gene in the blood buffy coat was positive. The patient was diagnosed with scrub typhus. A comparative analysis of the O. tsutsugamushi DNA sequence from the patient, with those in the GenBank, confirmed that he was infected with the Boryong genotype; a 500-mg azithromycin intravenously was initiated after hospital admission. However, the patient progressed to a semi-coma on the third day of hospitalization. An additional brain CT scan was performed (Figure 1B). Extensive subdural hemorrhage was detected in the left frontotemporoparietal area, along with subfalcine herniation. The patient became comatose and died.

CASE 2

A 53-year-old woman presented at our hospital with a 20-day history of maculopapular skin rashes on the anterior chest and a 5-day history of fever, headache, and nausea. She had worked in an agricultural field twice a week before her admission. Her past medical history, social history, and family history were non-contributory. At presentation, the patient was conscious. Upon the physical examination, non-pruritic erythematous maculopapular rashes were observed on the anterior chest, and a 1 cm × 1 cm eschar was noted on the left axilla.

The blood coagulation test results were as follows: PT, 11.3 sec; international normalized ratio (INR), 0.9; aPTT, 32.6 sec; fibrinogen, 257 mg/dL; FDP, 2.27 µg/mL; and D-dimer, 389 ng/mL. Immunofluorescence assays to detect antibodies against O. tsutsugamushi revealed IgM and IgG titers of 1:40 and 1:512, respectively, at presentation, and increases of at least 4-fold in the IgM titer (1:160) and IgG titer (1:4,096) were observed after 7 days. Nested PCR targeting the O. tsutsugamushi 56 kDa protein-encoding gene was negative, but nested PCR for an eschar was positive. The presence of O. tsutsugamushi Boryong was confirmed by a sequencing test.

After a clinical diagnosis of scrub typhus, 600 mg of rifampin were given, but a severe headache persisted. The CT scans performed on the third hospital day revealed a focal hyperdensity and a small amount of blood in the right cerebral hemisphere (Figure 1C). At that time, the routine blood test results were normal and blood coagulation tests revealed the following: PT, 11.4 sec; INR, 0.92; aPTT, 28.6 sec; fibrinogen, 129 mg/dL; FDP, 5.18 µg/mL; and D-dimer, > 1,050 ng/mL. The serum levels of aspartate aminotransferase (227.6 IU/L) and alanine aminotransferase (417.6 IU/L) had increased,
compared with the levels at the patient’s initial presentation. The patient was discharged from the hospital without any specific sequelae on the ninth day of hospitalization.

**CASE 3**

A 74-year-old man presented at our hospital with high fever and skin rashes. A generalized myalgia had occurred 2 weeks before presentation. He had a history of coronary interventions and took antiplatelet agents and oral hypoglycemic agents. There was no remarkable family history. At presentation, the patient was alert. He had a blood pressure reading of 130/80 mm of Hg, a pulse rate of 85/min, respiratory rate of 24/min, and body temperature of 38°C. The physical examination indicated conjunctival injection and nuchal rigidity. Non-pruritic maculopapular rashes were observed on the chest and abdomen, and a 1 cm × 1.5 cm eschar was noted on the anterior chest. Hematochemical tests revealed the following: white blood count, 11,950/μL; hemoglobin, 12.9 g/dL; platelets, 119,000/μL. Blood coagulation tests indicated the following: PT, 11.9 sec; INR, 0.98; aPTT, 27.9 sec; fibrinogen, 437 mg/dL; FDP 11.8 μg/mL; and D-dimer, >1,050 ng/mL. Immunofluorescence assays to detect antibodies targeting *O. tsutsugamushi* revealed IgM and IgG titers of 1:80 and 1:128, respectively, at presentation, which increased to 1:320 and 1:11,024, respectively, after 13 days. Nested PCR targeting the *O. tsutsugamushi* 56-kDa protein-encoding gene was positive; a comparative analysis of the *O. tsutsugamushi* DNA sequence of the patient and those in the GenBank confirmed that he had the Boryong genotype. Based on the clinical features at presentation, the patient was presumptively diagnosed with scrub typhus and given 600 mg of rifampin, after which his fever subsided. At 3:00 am on the fourth day, the patient fell because of muscle weakness during an attempt to get out of bed, and motor weakness was noted on his left side. The brain MRI indicated recent onset infarction in the right middle cerebral artery territory (Figure 1D–G). This study was approved by the Institutional Ethics Board (2012-12-008) for the Clinical Research of Chosun University Hospital.

**DISCUSSION**

The main pathologic findings of scrub typhus are vasculitis and perivasculitis. Corresponding to this pathologic finding, hemorrhages in various organs, such as atraumatic hemoperitoneum or massive GI bleeding, and acute myocardial infarction or splenic infarctions have been reported as possible complications in scrub typhus patients.3,4 Endothelial cell dysfunction and disseminated intravascular coagulation may occur in response to endothelial cell injury by *O. tsutsugamushi* in scrub typhus patients,5,6 even though actual mechanism is not fully understood. According to the definition of disseminated intravascular coagulation (DIC) based on the criteria presented by the Scientific Subcommittee on DIC of the International Society of Thrombosis and Hemostasis (ISTH), overt DIC and thrombocytopenia were observed in two of the three patients with a cerebral vascular accident (CVA) in this study.7 Microorganisms that cause the spotted fever group rickettsial diseases induce coagulopathies and thrombotic events8,9;
Collazos and others reported that a patient infected with O. tsutsugamushi was the direct cause of their CVA. However, it was difficult to conclude that O. tsutsugamushi was the direct cause of their CVA. How ever, it is very possible that O. tsutsugamushi may have acted as a provoking factor for CVA.

In conclusion, scrub typhus is known to cause widespread endothelial injury, and this injury is also likely to occur in the cerebral endothelium. Patients already predisposed to develop a CVA and who receive delayed treatment of scrub typhus are at an increased risk for developing a CVA.

**REFERENCES**


