A Case of Successful Treatment of Septic Shock Secondary to Scrub Typhus

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Abstract

Scrub typhus is a tropical disease and endemic problem mostly in rural parts of Malaysia. Typical presentation is acute febrile illness with systemic involvement, Multiple Organ failure and death. The non-specific flu like symptoms, low index of suspicion by clinicians and the unavailability of diagnostic test in most healthcare centres results in under-recognized scrub typhus. We report a case of a traveller to rainforest - National Park, Malaysia who presented with acute febrile illness and septicemic shock with pathognomonic eschar and positive serology test for scrub typhus. The patient recovered after appropriate diagnosis and prompt treatment.

Keywords: Scrub Typhus • Eschar • Multiple Organ Failure • Fatal

Introduction

Scrub typhus is a lethal infectious disease caused by Orientia tsutsugamushi, a gram negative intracellular bacterium which is transmitted to humans through the bite of infected Leptotrombidium mites. Typical presentation is acute febrile illness with non specific symptoms, systemic involvement with severe complications and death [1]. The disease is commonly distributed throughout the Asia Pacific regions including Malaysia [2,3]. There were limited data on the prevalence of scrub typhus in Malaysia. However a study on the antibody prevalence Orientia tsutsugamushi to aboriginal population in West Malaysia suggested that scrub typhus remains an important disease amongst various aboriginal subgroups with the highest prevalence being observed for the Pahang subgroup [4,5]. The study concluded environment, socioeconomic and behavioural risk factors have a significant relationship to the risk of exposure to scrub typhus [5]. The study was only included 280 individuals from 7 aboriginal subgroups yet it may not represent the entire native populations [4].

The common clinical manifestations of the disease are fever, headache and myalgia. The diagnosis of scrub typhus can be challenging due to the nature of the disease; non-specific flu like illness which can mimic many other zoonotic diseases. An eschar is a pathognomonic clinical feature of scrub typhus, begins as a papule at the site of chigger feeding and then ulcerates form a black crust like a skin burn from a cigarette [6]. Laboratory methods for diagnosing scrub typhus are mainly based on serological tests and molecular assays with the gold standard test for diagnosis of scrub typhus is the indirect Immunofluorescence Assay (IFA) [8]. The indirect Immunoperoxidase Assay (IIP) serological test is a modification of IFA that provides a comparable sensitivity and specificity in diagnosis of scrub typhus [7]. Scrub typhus can result in severe multiorgan failure with a case fatality rate up to 30% or higher without appropriate treatment [8]. There are limited data on the rate of sepsis in scrub typhus infection however a study done in Northern India emphasized that sepsis with Multiple Organ Dysfunction Syndrome (MODS) is the second most common complications after acute respiratory distress syndrome [9]. We present a case of scrub typhus in a 24-year-old traveller with acute febrile illness and septicemic shock which later found to have pathognomonic eschar and positive serological test for scrub typhus.

Case Report

A 24-years-old Malaysian man presented to the emergency department with a 9-day history of fever, headache, diarrhea and loss of appetite. He denied cough or runny nose. Two weeks before illness, he travelled to National Park in Pahang, Malaysia for jungle trekking. In the ward, patient was ill looking and lethargic, febrile with temperature of 39°C, tachycardic with heart rate of 120 beats per minute, blood pressure of 100/60 mmHg requiring intravenous noradrenalin 0.5 mcg/kg/min with oxygenation saturation of 97% on room air. Physical examination disclosed presence of two eschar measured 7 mm below his right axilla and left forearm (Figure 1). Chest auscultation and abdominal examination were unremarkable. Laboratory test revealed no leukenopenia, thrombocytopenia or anaemia but elevated serum Aspartate Aminotransferase (AST), Alanine Transf erase (ALT) and Lactate Dehydrogenase (Table 1). He was initially given empirical therapy of intravenous ceftriaxone 2 g daily for possible leptospirosis. However in view of the travelling history to rainforest and presence of eschar, we decided to treat the patient for possible scrub typhus and therefore empirical therapy is switched to tablet doxycycline 100 mg twice daily. Patient showed marked clinical improvement in less than 24 h of doxycycline where his temperature and blood pressure return to normal and he felt better. He was discharged without complication. During his clinic visit at day-10 of treatment, his symptoms resolved and the eschars dissapeared. His serology test for scrub typhus antibody IgM test was positive with the titre of 1:100. His leptospirosis serology and blood film for malaria parasite was negative. Blood culture and sensitivity test yielded no growth.

Figure 1. Presence of two eschar measured 7 mm below his right axilla and left forearm.

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**Table 1. Laboratory profile of the patient on admission in Hospital Bentong.**

<table>
<thead>
<tr>
<th>Variable</th>
<th>Value</th>
<th>Range</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hematology</td>
<td></td>
<td></td>
</tr>
<tr>
<td>White blood cell</td>
<td>9.7 x 10^9/UL</td>
<td>4.0-11.0</td>
</tr>
<tr>
<td>Neutrophils</td>
<td>50.20%</td>
<td>45.0-75.0</td>
</tr>
<tr>
<td>Lymphocytes</td>
<td>41.60%</td>
<td>24.0-45.0</td>
</tr>
<tr>
<td>Hemoglobin</td>
<td>14.3 G/DL</td>
<td>12.0-18.0</td>
</tr>
<tr>
<td>Platelet</td>
<td>258 x 10^9/UL</td>
<td>150-400</td>
</tr>
<tr>
<td>Prothrombin time</td>
<td>16.9 sec</td>
<td>14-Dec</td>
</tr>
<tr>
<td>Activated Partial Thrombin Time (APTT)</td>
<td>33.7</td>
<td>&lt;60</td>
</tr>
<tr>
<td>Blood chemistry</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Alanine Transaminase (ALT)</td>
<td>159 U/L</td>
<td>&lt;65</td>
</tr>
<tr>
<td>Aspartate aminotransferase</td>
<td>109 U/L</td>
<td>&lt;35</td>
</tr>
<tr>
<td>Lactate dehydrogenase</td>
<td>603 U/L</td>
<td>133-225</td>
</tr>
<tr>
<td>Creatinine kinase</td>
<td>69 U/L</td>
<td>&lt;171</td>
</tr>
<tr>
<td>Creatinine</td>
<td>101 UMOL/L</td>
<td>59-104</td>
</tr>
<tr>
<td>Enthocyte sedimentation rate</td>
<td>14 MM/HR</td>
<td></td>
</tr>
</tbody>
</table>

**Discussion**

Scrub typhus is a zoonosis caused by *Orientia tsutsugamushi*, a gram-negative intracellular bacterium. Patient often presented with abrupt onset of fever (normally 8 days to 16 days) following bites from infected mite [10]. Other non specific symptoms include myalgia, rashes and headache which also seen in cases of leptospirosis, dengue, and malaria [9]. In view of the clinical similarities in patients infected with zoonotic disease, the diagnosis of scrub typhus remains a challenge to physicians. Mites are very small and therefore patient may not be aware of the mite bite until they develop eschar. Failure to diagnose scrub typhus results in treatment delay and patient will end up with severe complications such as multiple organ failure, disseminated intravascular coagulopathy and even death. In this case, patient initially given empirical therapy for leptospirosis in view of fever with non specific symptoms for more than a week, history of jungle tracking, swimming, and elevated liver enzyme. However the presence of eschar favours the diagnosis of scrub typhus and patient was promptly manage with doxycycline. Eschar is a valuable clue to the diagnosis of scrub typhus however the overall prevalence of eschar ranged widely in patients infected with the disease [11]. An eschar is not observed in every confirmed cases, therefore history of travelling to or reside in an endemic area is also an important clue to favour the diagnosis of scrub typhus [8]. Clinician must examine patient thoroughly to look for presence of eschar and to get proper travel history especially in areas endemic for scrub typhus [8].

Antibody based serological tests are the mainstay of scrub typhus diagnosis [10]. The older test; Weil–FelixOX-K agglutination reaction test is easy to perform but lack of specificity and sensitivity [12]. The Indirect Fluorescent Antibody (IFA) and Indirect Immunoperoxidase (IIP) are the current reference standard and sensitive test for scrub typhus. IFA uses fluorescent anti human antibody to detect specific antibody towards scrub typhus antigen from patient serum whereas the IIP use peroxidase instead of fluorescein for the same manner [13]. Our patient had IIP test for scrub typhus IgM and IgG and the titre was significant for scrub typhus infection. IIP test is more sensitive test for scrub typhus antigen in endemic area due to the rapid clinical resolution manner following this antibiotic [17]. In cases of doxycycline-resistance strain causing scrub typhus, azithromycin has been advocated to be an effective alternative treatment [18]. In this case, early diagnosis and prompt delivery of correct antimicrobial therapy manage to save the patient and prevent further complications.

**Conclusion**

Scrub typhus has appeared as one of the important cause of acute febrile illness which has been neglected for many years [4]. The diagnosis of scrub typhus can be challenging due to the non specific clinical symptoms and signs that can mimic other zoonotic disease. Therefore a thorough history and detailed physical examination are important to aid the diagnosis of scrub typhus. Clinicians must have a high index suspicion of scrub typhus especially to those patients who reside or travel to endemic area. A delay of treatment may result in severe complications and death. A continuous education program and awareness of the disease as one of the differential diagnosis of acute febrile illness in region where the disease are endemic must be advocated.

**Declaration of Patient Consent**

The authors certify that they have obtained all appropriate patient consent forms. In the form the patient(s) has/have given his/her/thier consent for his/her/thier images and other clinical information to be reported in the journal. The patients understand that their names and initials will not be published and due efforts will be made to conceal their identity, but anonymity cannot be guaranteed.

**References**

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