Rickettsia japonica Infections in Humans, Xinyang, China, 2014–2017

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During 2014–2017, we screened for Rickettsia japonica infection in Xinyang, China, and identified 20 cases. The major clinical manifestations of monoinfection were fever, asthenia, myalgia, rash, and anorexia; laboratory findings included thrombocytopenia and elevated hepatic amino-transferase concentrations. Physicians in China should consider R. japonica infection in at-risk patients.

Emerging and reemerging spotted fever group (SFG) rickettsioses are increasingly spreading and being recognized worldwide. Japanese spotted fever (JSF), caused by Rickettsia japonica and first reported in Japan in 1984 (1,2), is an SFG rickettsiosis characterized by fever, malaise, chills, headache, rash, and eschars (3,4). JSF cases have been increasingly reported, but mostly in Japan and the nearby countries of South Korea, the Philippines, and Thailand (4–7). R. japonica has been detected in 3 genera (Haemaphysalis, Dermacentor, Ixodes) and 8 species of ticks (8,9). Because of the wide spectrum of ticks in which this bacterium has been found, R. japonica is postulated to be distributed more widely throughout China. The identification of R. japonica in H. flava and H. hystricis ticks in Wuhan, China, further corroborated this hypothesis (10). Also, serologic responses to R. japonica were detected in patients in Hainan Province in southern China ≈3 decades ago (11). However, laboratory-confirmed JSF cases have been identified in only a few patients in China (12,13). In this study, we aimed to determine if more JSF cases were occurring in China by using hospital-based surveillance.

The Study

During March 2014–June 2017, we screened 2,236 febrile patients seeking treatment at The People’s Liberation Army 990 Hospital in Xinyang, Henan Province, eastern central China (Appendix Figure, https://wwwnc.cdc.gov/EID/article/25/9/17-1421-App1.pdf) with a history of tick bite or field activity within the previous month for infection with R. japonica. We extracted DNA from peripheral blood samples collected during the acute phase of illness and screened them by a nested PCR concurrently targeting the citrate synthase gene (gltA), 17-kDa antigen–encoding gene (htrA), and outer membrane protein A–encoding gene (ompA), as previously described (14). To minimize the risk for contamination, we performed template isolation and PCR setup in separate rooms with specified pipette sets, used certified DNAase- and RNase-free filter barrier tips to prevent aerosol contamination, and performed all PCR assays with appropriate controls. After amplification and sequencing, we found that 20 patients were infected with R. japonica.

The gltA gene sequence (456-bp) obtained from all 20 samples (GenBank accession no. MF693145) showed 100% similarity to the corresponding gene of R. japonica strain YH (GenBank accession no. AP017602). The htrA gene sequence (394-bp) obtained from 18 samples (GenBank accession no. MF693146) was identical to that of R. japonica YH, and the sequence in the remaining 2 samples (GenBank accession no. MF693147) differed by only 3 bp. The ompA gene sequence (311-bp) found in 16 samples (GenBank accession no. MF693148) was identical to that of R. japonica YH, and the remaining 4 samples each differed by just 1 bp (GenBank accession nos. MF693149–52) (Figure 1).

We also assessed other tickborne agents, including Anaplasma phagocytophilum, Anaplasma caprae, Ehrlichia chaffeensis, Borrelia burgdorferi sensu lato, Babesia microti, and severe fever with thrombocytopenia syndrome virus, by PCR or reverse transcription PCR. Six of 20 patients were co-infected with severe fever with thrombocytopenia syndrome virus, so we excluded them from clinical analysis.

We tested the acute serum samples collected ≤7 days after illness onset and the convalescence serum samples collected >14 days after illness onset from the 14 patients with R. japonica monoinfection in parallel using an indirect immunofluorescence assay for Rickettsia heilongjiangensis IgG. We considered an IgG titer >1:64 a positive reaction. Using this definition, we found that 6 patients had a seroconversion, and the other 8 had a 4-fold increased IgG titer; thus, all patients had an acute infection with SFG rickettsiae.
All 14 patients had a history of field activity within the previous month, and 5 had a history of tick bite. Median patient age was 61.5 (range 44–77) years; 9 were men. The median time from tick bite to onset of illness was 4 (range 3–8) days and from onset of illness to physician visit was 5 (range 2–7) days. The median duration of hospitalization was 6 (range 4–10) days.

Fever and asthenia were reported by all 14 patients. Other nonspecific symptoms included myalgia (10/14), lymphadenopathy (4/14), headache (3/14), dizziness (3/14), and chills (2/14). Gastrointestinal manifestations included anorexia (9/14), nausea (7/14), vomiting (3/14), and diarrhea (1/14). Three patients had cough and pneumonia, and we observed rash in 10 patients (Figure 2). The median time from onset of illness to rash was 4 (range 3–5) days, and the median duration of rash was 5.5 (range 4–8) days. Only 3 patients had an eschar. Other signs included splenomegaly (2/14) and facial edema (1/14).

Urinalysis on admission revealed microscopic hematuria in 2 patients and a slight or moderate proteinuria in 8 patients. The most common findings on laboratory tests were thrombocytopenia, elevated hepatic aminotransferase concentrations, elevated serum lactate dehydrogenase, and hypoalbuminemia, followed by hyponatremia, anemia, hyperbilirubinemia, hypopotassiumaemia, leukopenia, and elevated serum creatine kinase (Table). Mild multiple organ dysfunction developed in 3 patients.

Because of the appearance of typical rash, we suspected SFG rickettsioses in 10 patients and treated them with doxycycline (100 mg 2×/d) until their fevers dissipated and clinical disease improved (median 4 [range 3–5] days). The other 4 patients were treated with the antimicrobial drugs...
Multiple organ dysfunction developed in several patients. Moderate, and no deaths have been recorded, although mild resembling those of patients with common SFG rickettsiosis. Bacteremia, hypoalbuminemia, elevated hepatic enzyme levels, and symptoms are rarely seen in patients with spotted fever group rickettsioses. Thrombocytopenia and neurologic signs are commonly seen, and hemorrhagic and neurologic signs are not always easy to identify. In addition, the patients we identified usually had myalgia and gastrointestinal symptoms.

These clinical findings expand the available knowledge of the disease spectrum of *R. japonica*. Compared with endemic rickettsiosis caused by *Candidatus Rickettsia tarasevichiae* infection (reference 16 in Appendix), rash is commonly seen, and hemorrhagic and neurologic signs and symptoms are rarely seen in patients with *R. japonica* infection. These distinctive features could be used to make a differential diagnosis.

We frequently observed in our patient cohort thrombocytopenia, hypoalbuminemia, elevated hepatic enzyme activity, and elevated lactate dehydrogenase levels, findings resembling those of patients with common SFG rickettsioses (reference 17 in Appendix). In general, disease is mild or moderate, and no deaths have been recorded, although mild multiple organ dysfunction developed in several patients.

In conclusion, we identified *R. japonica* as an emerging tickborne pathogen in China. Physicians in areas where *H. longicornis* and other competent vectors for *R. japonica* are endemic should be aware of the risk for infection in humans and prescribe doxycycline to patients in cases of ineffective therapy with other antimicrobial drugs. Surveillance needs to be extended to improve our understanding of the health burden of JSF.

This study was supported by the National Natural Science Foundation of China (81825019, 81722041, 81472005, and 81621005), the China Mega-Project for Infectious Diseases (2018ZX10713002-002), the National Key Research and Development Program of China (2016YFX1201905), and the New Star Plan of Science and Technology of Beijing (Z171100001117089).

### Conclusions

The tick vectors of *R. japonica* (i.e., *H. flava*, *H. hystricis*, *H. cornigera*, *H. longicornis*, *I. ovatus*) are widely distributed throughout China (15), providing *R. japonica* ample opportunity to infect humans. By applying molecular screening techniques to simultaneously amplify 3 genes, we identified *R. japonica* infection in 14 patients.

The clinical signs and symptoms of *R. japonica* infection in our patient cohort differed from those reported in patients in Japan (3, 4). We frequently observed fever, asthenia, and rash but not chills and headache. We also saw fewer eschars in our patient cohort, potentially because they were underreported during clinical examination; eschars are not always easy to identify. In addition, the patients we identified usually had myalgia and gastrointestinal symptoms.

These clinical findings expand the available knowledge of the disease spectrum of *R. japonica*. Compared with endemic rickettsiosis caused by *Candidatus Rickettsia tarasevichiae* infection (reference 16 in Appendix), rash is commonly seen, and hemorrhagic and neurologic signs and symptoms are rarely seen in patients with *R. japonica* infection. These distinctive features could be used to make a differential diagnosis.

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### Table. Laboratory test results of samples from 14 patients with *Rickettsia japonica* infection at different time points, China, 2014–2017*

<table>
<thead>
<tr>
<th>Result</th>
<th>At admission</th>
<th>During hospitalization</th>
<th>At discharge from hospital</th>
</tr>
</thead>
<tbody>
<tr>
<td>Anemia, &lt;3.5 × 10^12 erythrocytes/L</td>
<td>0</td>
<td>5</td>
<td>3</td>
</tr>
<tr>
<td>Leukopenia, &lt;4.0 × 10^3 leukocytes/L</td>
<td>3</td>
<td>3</td>
<td>0</td>
</tr>
<tr>
<td>Thrombocytopenia, &lt;150 × 10^3 platelets/L</td>
<td>9</td>
<td>11</td>
<td>0</td>
</tr>
<tr>
<td>Hyperbilirubinemia, albumin &gt;17.1 μmol/L</td>
<td>4</td>
<td>5</td>
<td>2</td>
</tr>
<tr>
<td>Hypoalbuminemia, albumin &lt;35.0 g/L</td>
<td>3</td>
<td>9</td>
<td>7</td>
</tr>
<tr>
<td>Hyponatremia, sodium &lt;135.0 mmol/L</td>
<td>4</td>
<td>6</td>
<td>2</td>
</tr>
<tr>
<td>Hypopotassemia, potassium &lt;3.5 mmol/L</td>
<td>2</td>
<td>4</td>
<td>2</td>
</tr>
<tr>
<td>Increased ALT level, &gt;0.67 μkat/L</td>
<td>8</td>
<td>10</td>
<td>5</td>
</tr>
<tr>
<td>Increased AST level, &gt;0.67 μkat/L</td>
<td>10</td>
<td>12</td>
<td>5</td>
</tr>
<tr>
<td>Increased LDH level, &gt;4.1 μkat/L</td>
<td>10</td>
<td>11</td>
<td>7</td>
</tr>
<tr>
<td>Increased CK level, &gt;4.1 μkat/L</td>
<td>4</td>
<td>3</td>
<td>0</td>
</tr>
</tbody>
</table>

*ALT, alanine aminotransferase; AST, aspartate aminotransferase; CK, creatine kinase; LDH, lactate dehydrogenase.*

### References


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Appendix

**Appendix Figure.** Location of Xinyang, Henan Province, China, where the patients with *Rickettsia japonica* infection were detected. The triangle indicates the location of Wuhan, Hubei Province, China, where the ticks of *R. japonica* infection were detected in a previous study (reference 10 in main text).
Appendix References


https://doi.org/10.7326/M15-2572