repercussions for human health? Given that 95%–99% of humans possibly exposed to such a reservoir are Duffy negative, and therefore resistant to the parasite, these would appear to be slight. However, as humans encroach more frequently into ape habitats, the chances of humans encountering the parasite will increase. In the short term, the risks are probably limited to Duffy-positive persons who enter areas where apes are present, such as tourists and migrant workers.

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Rickettsia parkeri and Candidatus Rickettsia andeanae in Gulf Coast Ticks, Mississippi, USA

To the Editor: Rickettsia parkeri, a spotted fever group Rickettsia (SFGR) bacterium, is transmitted by Amblyomma maculatum, the Gulf Coast tick (1). The prevalence of R. parkeri in Gulf Coast ticks has been reported as <42% in the United States, which is higher than reported rates of R. rickettsii (the cause of Rocky Mountain spotted fever) in Dermacentor species ticks. Misdiagnosis among SFGR infections is not uncommon, and R. parkeri rickettsiosis can cause symptoms similar to those for mild Rocky Mountain spotted fever (1). We evaluated infection rates of R. parkeri and Candidatus Rickettsia andeanae, a recently identified but incompletely characterized SFGR, in Gulf Coast ticks in Mississippi, USA.

During May–September of 2008–2010, we collected adult Gulf Coast ticks from vegetation at 10 sites in Mississippi. We extracted genomic DNA from the ticks using the illustra tissue and cells genomicPrep Mini Spin Kit (GE Healthcare Life Sciences, Piscataway, NJ, USA). We tested amplifiable tick DNA by PCR of the tick mitochondrial 16S rRNA gene (2). We tested for molecular evidence of any SFGR species by nested PCR of rompA (rickettsial outer membrane protein A gene) (1). Samples positive for SFGR were subsequently tested by using species-specific rompA PCR for R. parkeri (3) and Candidatus R. andeanae (4). All PCRs included 1) a positive control of DNA from cultured R. parkeri—(Tate’s Hell strain) or Candidatus R. andeanae–infected Gulf Coast ticks and 2) a negative control of water (nontemplate). PCR products were purified by using Montage PCR Centrifugal Filter Devices (Millipore, Bedford, MA, USA) and sequenced by using Eurofins MWG Operon (Huntsville, AL, USA). We generated consensus sequences using ClustalW2 (www.ebi.ac.uk/Tools/msa/clustalw2/) alignment and identified the sequences using GenBank BLAST searches (www.ebi.ac.uk/Tools/clustalw2/).

Proportions of ticks infected with SFGR, by region and year, were compared separately by using Fisher exact test followed by pairwise comparisons with a Bonferroni
embrace (PROC FREQ, SAS for Windows, V9.2; SAS Institute, Cary, NC, USA). For all analyses, p<0.05 was considered significant. An index of coinfection was calculated by using the formula IC = (O – E)/N × 100, in which IC is index of coinfection, O is number of coinfections, E is expected occurrence of co-infection caused by chance alone, and N is total number of ticks infected with either Rickettsia species. A χ² test was used to determine statistical significance (5).

A total of 707 adult Gulf Coast ticks were collected during the 3 years (350 in 2008, 194 in 2009, and 163 in 2010). Tick mitochondrial 16S rRNA gene was detected in 698 (98.7%), of which 128 (18.3%) were positive for SFGR DNA, comprising 106 (15.2%) positive only for R. parkeri, 10 (1.4%) positive only for Candidatus R. andeanae, and 12 (1.7%) co-infected with R. parkeri and Candidatus R. andeanae (Table). Positive test results from 22 ticks singly or co-infected with Candidatus R. andeanae were confirmed by sequencing.

Most (94.6%) ticks were from northern (n = 260) and southern (n = 409) Mississippi; this frequency is similar to those reported in other studies of Gulf Coast ticks in Mississippi (9), awareness of R. parkeri rickettsiosis should be increased in this state. We identified Candidatus R. andeanae in ≈3% of Gulf Coast ticks in Mississippi; this frequency is significantly higher than expected from chance alone. The biologic role of co-infections of Gulf Coast ticks with R. parkeri and Candidatus R. andeanae remains to be determined.

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References


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Table. PCR results for adult Rickettsia parkeri– and Candidatus Rickettsia andeanae–infected Gulf Coast ticks (Amblyomma maculatum) collected from 10 sites in Mississippi, USA, 2008–2010

<table>
<thead>
<tr>
<th>Location (no. collection sites)</th>
<th>No. ticks</th>
<th>No. (%)</th>
<th>SFG rompA</th>
<th>R. parkeri only</th>
<th>No. (%)</th>
<th>Expected no. (%)</th>
<th>Co-infected ticks</th>
<th>No. (%)</th>
<th>Co-infected ticks</th>
</tr>
</thead>
<tbody>
<tr>
<td>North (4)</td>
<td>257</td>
<td>49 (19.1; 14.5–24.4)†</td>
<td>48 (18.7; 14.1–24)†</td>
<td>0§</td>
<td>0.19 (0.07)</td>
<td>1 (0.4; 0–2.1)¶</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Central (1)</td>
<td>38</td>
<td>4 (10.5; NA)</td>
<td>1 (2.6; NA)</td>
<td>2 (5.3; NA)</td>
<td>0.16 (0.42)</td>
<td>1 (2.6; NA)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>South (5)</td>
<td>403</td>
<td>75 (18.6; 14.9–22.8)†</td>
<td>57 (14.1; 10.9–17.9)†</td>
<td>8 (2.0; 0.9–3.9)§</td>
<td>2.99 (0.74)</td>
<td>10 (2.5; 1.2–4.5)¶</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total (10)</td>
<td>698</td>
<td>128 (18.3; NA)</td>
<td>106 (15.2; NA)</td>
<td>10 (1.4; NA)</td>
<td>3.65 (0.52)</td>
<td>12 (1.7; NA)</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

*The estimated value of co-infection caused by chance alone (E) was calculated by using the formula E = (a + b)/(a + c) / (a + b + c + d) (5), where a = no. ticks infected with both Rickettsia species, b = no. ticks infected only with R. parkeri, c = no. ticks infected only with Candidatus R. andeanae, and d = no. ticks not infected with either Rickettsia species. SFG rompA, spotted fever group rickettsial outer membrane protein A gene. NA, not applicable. †p = 0.003, §p = 0.06. ¶p = 0.0578 (comparison of prevalence from northern and southern sites only).
Attributing Cause of Death for Patients with Clostridium difficile Infection

To the Editor: Hota et al. report that for deceased patients who had Clostridium difficile infection (CDI), agreement is poor between causes of death reported on death certificates and those categorized by a review panel (1). Our data support the difficulty of attributing cause of death for patients with CDI.

In 2004 in Quebec, Canada, a mandatory CDI surveillance program was implemented. Deaths that occurred within 30 days after CDI diagnosis were classified as 1) directly attributable to CDI (e.g., toxic megacolon, septic shock), 2) having a CDI contribution (e.g., acute decompensation of chronic heart failure), or 3) unrelated to CDI (e.g., terminal cancer) (2). To determine accuracy of the surveillance classifications, we compared cause-of-death classification of 22 deceased CDI patients reported to surveillance by 1 hospital in 2007 with causes of death reported by 13 external reviewers who examined summaries of medical files of the deceased patients. Reviewers were 11 infectious disease and 2 public health physicians involved with CDI surveillance at their respective hospitals but not this hospital. The median (minimal, maximal) κ statistics for comparison of external reviews with surveillance classification were 0.495 (0.252, 0.607) for directly attributable, 0.182 (−0.091, 0.182) for contributed, and 0.321 (0.124, 0.614) for unrelated. Comparison within external reviewers yielded 0.697 (0.394, 1.0), 0.233 (−0.294, 0.703), and 0.542 (0.154, 0.909), respectively. Complete agreement was found for only 6 cases (4 directly attributable and 2 unrelated) (Figure).

Variation among reviewers suggested that categorizations reported to surveillance were inaccurate. Number of deaths among patients with CDI, regardless of the cause of death, seemed to better indicate CDI severity. Since 2008, only the crude numbers of deaths, not subjected to individual interpretation, have been reported to surveillance. A questionnaire addressing concurrent medical conditions, prognosis, level of care, and circumstances of death is being implemented in Quebec hospitals participating in CDI surveillance and should help determine the role of CDI in deaths.
**Rickettsia parkeri** and **Candidatus**
Rickettsia andeanae in Gulf Coast Ticks, Mississippi, USA

Technical Appendix

Technical Appendix Figure. Distribution of Gulf Coast ticks and cases of rickettsiosis. Solid circles indicate confirmed cases of *Rickettsia parkeri* rickettsiosis, and open circles indicate probable cases of *R. parkeri* rickettsiosis noted by Paddock et al. (1) and Cragun et al. (2). A) Gray area indicates the most established distribution of Gulf Coast ticks (*Amblyomma maculatum*); asterisks (*) indicate documented collection of these ticks in incidental reports (may or may not reflect permanent populations). B) Gray shading indicates Mississippi counties where Goddard and Paddock reported the occurrence of Gulf Coast ticks and locations of previously reported cases of *R. parkeri* rickettsiosis in Mississippi (3). X indicates tick collection sites for this study. Map adapted with permission from Teel et al. (4).

References

