The Role of Ranged Horses in Eco-Epidemiology of *Rickettsia raoultii* Infection in China

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*Rickettsia raoultii* is a tick-borne pathogen that infects humans; however, the vertebrate hosts of this pathogen have not been clearly defined. Our molecular examination of *Rickettsia* spp. infecting mammals and ticks in China, identified the *gltA*, *ompA*, and 17KD gene sequences of *R. raoultii* in horses and their ticks. This indicates a role of horses in *R. raoultii* epidemiology.

**Keywords:** horse, infection, *Rickettsia raoultii*, tick-borne pathogens, China

**INTRODUCTION**

Tick-borne rickettsioses are recognized as emerging vector-borne infections, infecting both human and animal hosts worldwide. *Rickettsia raoultii* was initially implicated as the causative agent of human infection in 2006 through the detection of DNA in the blood of a Spanish patient (Ibarra et al., 2006), and has since been reported in human infections in many countries of the world; in particular, China (Jia et al., 2014; Li et al., 2018; Dong et al., 2019). *R. raoultii* was first detected in *Dermacentor nuttalli* and *Rhipicephalus pumilio* ticks in 1999 (Rydkina et al., 1999). Although *R. raoultii* has been detected in bloodsucking insects and other tick species, the dominant vectors are generally considered to be *Dermacentor* spp. (Silaghi et al., 2011; Liu et al., 2016). *Rickettsia raoultii* has been detected in a variety of ticks collected from dogs, cattle, and wildlife (Klitgaard et al., 2017; Chisu et al., 2018; Seo et al., 2020). All of these animals may serve as reservoir hosts for *R. raoultii*; however, experimental evidence is still lacking. Additionally, *R. raoultii* has been detected in squirrels, marbled polecats, red foxes, hedgehogs, yaks, camels and other wild mammals in China (Liu et al., 2018, 2021; Zhao et al., 2019; Li et al., 2020; Fang et al., 2021; Shao et al., 2021). Furthermore, there is a highly significant association of horse contact with tick-borne lymphadenopathy, the pathogens of which are *R. slovaca* and *R. raoultii* (Lakos et al., 2012). Horses also play a role in the epidemiology of Brazilian spotted fever, which is caused by *R. rickettsii*. Horses are considered the most suitable domestic agent for Brazilian spotted fever in some areas (Sangioni et al., 2005), which raises the question of whether horses may also play a role in the disease epidemiology of *R. raoultii*. The findings from this study provide evidence that *Rickettsia* species are in horses and sympatric ticks and that these hosts play a role in the circulation of the spotted fever group of *rickettsiae*. 
TABLE 1 | The status of *Rickettsia raoultii* infection in different samples.

| Sample        | No. tested | No. positive | Prevalence (%) |
|---------------|------------|--------------|----------------|
| Vertebrate    |            |              |                |
| Horse         | 12         | 6            | 50             |
| Cattle        | 16         | 0            | 0              |
| Sheep         | 10         | 0            | 0              |
| Wild mice     | 9          | 0            | 0              |
| Tick (D. Silvarum) |        |              |                |
| Adult (engorged) | 6         | 6            | 100            |
| Adult (free)  | 96         | 39           | 40.63          |
| Egg           | 5          | 5            | 100            |
| Larval        | 7          | 5            | 71.43          |
| Nymph         | 5          | 5            | 100            |

THE STUDY

To identify putative vertebrate hosts of *R. raoultii*, we collected blood samples from horses (*n* = 12), cattle (*n* = 16), sheep (*n* = 10), and brown rats (*Rattus norvegicus*, *n* = 9). All the animals infested by ticks, and all were from the same rangeland in Daqing, northeastern China (46°58′N, 125°03′E), where spotted fever group rickettsiosis was previously reported (Jia et al., 2014).

Free-living ticks (*Dermacentor silvarum*, *n* = 96) were collected by the flagging method and engorged adult ticks (*D. silvarum*, *n* = 6) were collected from horses in the same rangeland. Tick eggs (100 eggs, 5 pools) were laid in our laboratory by engorged wild-caught adult female *D. silvarum* from a PCR-positive horse. Larval and nymphal ticks of the same cohort that fed on mice were collected and pooled (35 larvae/10 nymphs; 12 pools in total). All the samples were tested for *Rickettsia* spp. This study was approved by the Research Ethics Committee of Heilongjiang Bayi Agricultural University, China.

Following the manufacturer’s instructions, DNA was extracted from the blood samples and ticks using a Tissue/Blood DNA Extraction Kit (Tiangen Biotech Inc., Beijing, China). PCR was performed to amplify the rickettsial citrate synthase gene (*gltA*), outer membrane protein A-encoding gene (*ompA*), and the 17-kDa antigen-encoding gene fragments; then, sequencing was performed (Li et al., 2018). The sequences obtained in this study were analyzed by an NCBI BLAST search. *Rickettsia* spp. were not detected in any of the samples from cattle, sheep, or brown rats, but sequences most closely related to *R. raoultii* were detected in 6/12 (50%) horses (Table 1). Additionally, *R. raoultii* was detected in 6/6 (100%) engorged ticks, 39/96 (41%) questing, 5/5 (100%) eggs, 5/7 (71%) larvae, and 5/5 (100%) nymphs (Table 1).

We deposited the sequences from 25 PCR-positive specimens into the GenBank database (accession nos. MH212168–92). For phylogenetic analysis, we used the MegAlign component to perform multiple sequence alignments with the ClustalX1.83 algorithm. A phylogenetic tree was constructed based on the *gltA* and *ompA* sequences identified in this study and other sequences from the GenBank database using the neighbor-joining method.

FIGURE 1 | Molecular phylogenetic analysis of *Rickettsia raoultii* isolates from horse. Two genes were concatenated (*gltA* + *ompA*), a total of (1,047 + 498) positions were tested. GenBank accession numbers of the Sequences of the *Rickettsia* species used to make the concatenated analysis were as follows: *R. rickettsii* (U59729 + U43804), *R. slovaca* (U59725 + U43808), *R. massiliae* (U59719 + U43799), *R. africae* (U59733 + U43790), *R. honei* (U59726 + U43809), *R. conorii* (U59730 + U43808), *R. japonica* (U59732 + U43801), *R. aeschlimanni* (U59722 + U43800), *R. helongiangensis* (AF172943 + AF179364), *R. parkeri* (KJ124257 + KJ271186), *R. rhipicephali* (DQ865208 + DQ865208), *R. monacensis* (LJ794217), *R. sibirica* (DQ365804 + AH015610), *R. raoultii* strain Marne (DQ365803 + AH015609), *R. raoultii* D1 from free-living tick, *R. raoultii* D4 from engorged adult tick on horse, *R. aeschlimanni* (DQ097038 + DQ097032), *R. tamurae* (AF394896 + DQ103256), *R. raoultii* China IM16 (KX745757 + KX74577), *R. raoultii* E1 from horse (MH212183 + MH212190), *R. raoultii* D1 from free-living tick (MH212179 + MH212186), *R. raoultii* E1 from horse (MH212183 + MH212190), *R. raoultii* D4 from engorged adult tick on horse (MH212181 + MH212188).
This revealed that the test samples from a horse, questing tick and an engorged adult tick clustered with other R. raoultii strains and were most closely related to R. raoultii strain MDJ1, which had been isolated from a patient in Mudanjiang, China (Figure 1).

DISCUSSION

Among our samples, R. raoultii was frequently detected in horses, but not in cattle, sheep, or brown rats, despite sharing pasture with the infected horses. A previous study in Xinjiang, China also reported the detection of R. raoultii in horses (Li et al., 2020). In this study, horses and other animals residing in the same environment were tested, but positive samples were only detected in horses. The results indicate that horses may be a vertebrate host of R. raoultii in the Daqing area. Horses are susceptible to a variety of rickettsial pathogens (Sangioni et al., 2005), and the prevalence rate is higher in some areas (Souza et al., 2016), indicating that horses may well be considered hosts. Therefore, horses may serve as a source of R. raoultii infection in ticks. Coincidentally, in a previous study, there were two human cases of R. raoultii infection in the same region (Jia et al., 2014).

In this study, the infection rate of R. raoultii in D. silvarum was 40.63%, which was higher than that (32.25%) in other studies (Wen et al., 2014). Furthermore, all engorged adult ticks and almost all egg/larval/nymphal tick pools were positive for R. raoultii. Our results support the idea that D. silvarum acts as a vector and a reservoir for R. raoultii. Schmuck et al. (2020) confirmed that transovarial transmission might be an efficient way of maintaining the infection cycle of R. raoultii, and both transtadial and transovarial transmission of R. raoultii have been demonstrated in D. nuttalli (Moore et al., 2018).

In conclusion, the main livestock grazing in the area were cattle, horses, and sheep, which were equally likely to be bitten by ticks, but R. raoultii was only detected in horses. Horses may therefore serve as a source of R. raoultii, contributing to the long-term preservation of R. raoultii in this area by promoting the infection of ticks and further increasing the chances of humans becoming infected with R. raoultii via tick bites.

DATA AVAILABILITY STATEMENT

The datasets presented in this study can be found in online repositories. The names of the repository/repositories and accession number(s) can be found below: https://www.ncbi.nlm.nih.gov/genbank/, MH212168-92.

AUTHOR CONTRIBUTIONS

J-Fj, A-QW, and Q-CC designed the project and experiments. YH, T-TW, and X-XM conducted the experiments. B-GJ and NJ analyzed the data. YH and Q-CC drafted the manuscript. All authors corrected, edited, and approved the manuscript.

FUNDING

This work was supported by the National Key Research Development Program of China (2019YFC1200501), the STU Scientific Research Foundation for Talents (NTF21043), and the National Natural Science Foundation of China (32072885).

ACKNOWLEDGMENTS

We thank Mallory Eckstut from Liwen Bianji, Edanz Group China (www.liwenbianji.cn/ac), for editing the English text of a draft of this manuscript.

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