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The First Human Case of *Rickettsia tamurae* Infection in Japan

Kaoru Imaoka^a Sakae Kaneko^a Kenji Tabara^b
Kenji Kusatake^a Eishin Morita^a

^aDepartment of Dermatology, Shimane University Faculty of Medicine, Izumo, and

^bShimane Prefectural Institute of Public Health and Environmental Science, Matsue, Japan

Key Words

Rickettsia tamurae infection · Spotted fever group · *Amblyomma testudinarium* · Local skin inflammation · Human pathogen

Abstract

A case of *Rickettsia tamurae* infection in Japan is reported. A 76-year-old Japanese male had a tick bite which developed to local skin inflammation on his left leg. Anti-rickettsia antibodies were detected in his serum, and *R. tamurae* DNA was identified in his blood, the lesional skin, and the tick.

Introduction

In recent decades, reports of newly described rickettsioses have increased as a result of improved cell culture isolation technique and extensive use of bacterial detection and identification techniques based on molecular biology. *R. tamurae* was isolated from the *Amblyomma testudinarium* tick in 1993 in Japan [1]. Fournier et al. formally identified *R. tamurae* as a novel species by genetic and phylogenetic analyses in 2006 [2]. Although *R. tamurae* has often been isolated from *A. testudinarium* ticks in Japan, the pathogenic role of this rickettsia in humans is yet unknown. We report the first official case of local skin inflammation caused by tick-borne *R. tamurae* in the Shimane prefecture, the western part of central Japan (fig. 1).

Case Report

A 76-year-old man visited Shimane University Hospital with a skin nodule in the left popliteal region and pain in the left leg on June 22, 2009. He had a history of multiple cerebral infarction, and had been suffering from insulin-dependent diabetes mellitus for 10 years. He often took a walk to some fields near his house. Three days before visiting our hospital, he had felt some pain in the left leg. One

day before the visit, a nodule was found in his left popliteal region and the left leg became swollen (fig. 2a). The nodule measured 15 mm in diameter and was identified as a tick body during a subsequent medical examination (fig. 2b). Although diffuse erythema, mild swelling and mild heat with pain were found in the left sural area, none of the typical clinical signs (e.g. high fever, generalized rash, lymphadenopathy) of spotted fever rickettsiosis was observed. The erythema blanched with pressure. Skin hemorrhages were absent. Hematological examination showed a white blood cell count of 9,070/μl and a C-reactive protein level of 1.30 mg/dl (less than 0.5 mg/dl is considered normal) on June 22, 2009. No abnormal results were found in other blood examinations.

The tick was removed with the attached skin by a scalpel under local anesthesia and identified as a female *A. testudinarium* based on morphological characteristics [3]. Further detail classification of the tick was performed with a PCR-based amplification technique of tick-associated DNA. DNA was extracted from the patient's blood, the skin biopsy specimen and the infected tick with a QIAamp DNA Mini Kit (QIAGEN, Courtaboeuf, France) according to the manufacturer's instructions. PCR amplification and sequencing reactions were performed using the following primers in accordance with a previously reported method: primers RpCS.877p/RpCS.1258n for the genus *Rickettsia* which target the citrate synthase-encoding gene (*gltA*) [4], primers Rr17.61p/Rr17.492n for the 17 kDa genus-common antigen gene [5], primers 10/11 for the 56 kDa outer membrane protein gene of *Orientia tsutsugamushi* [6] and primers p3761/p4183 for the 44 kDa outer membrane protein gene of *Anaplasma* sp. (p44 multigene family) [7]. Although the *O. tsutsugamushi* and genus *Anaplasma* genes were not detected, all samples were positive for the *gltA* and 17 kDa genus-common antigen genes. The DNA fragments separated by agarose gel electrophoresis were extracted using the QIAEX Gel Extraction Kit (QIAGEN). DNA sequencing was performed using an ABI PRISM® BigDye™ Terminator v3.1 KIT (Life Technologies, Carlsbad, Calif., USA) on an ABI Prism 3130 Genetic Analyzer (Life Technologies). The nucleotide sequences were compared with the corresponding sequences deposited in the DNA database (GenBank/EMBL/DDBJ) by using the BLAST tool (<http://blast.ddbj.nig.ac.jp/top-j.html>), and aligned using ClustalW on Biomanager by ANGIS (<http://www.angis.org.au>) and Mega 4 (version 1.83). A phylogenetic tree was constructed by the neighbor-joining method. The phylogram analysis of spotted fever group (SFG) rickettsiae derived from *gltA* and 17 kDa genus-common antigen genes revealed 100% nucleotide sequence homology with *R. tamurae* AT-1 (fig. 3a, b). Histological examination was not performed.

Oral minocycline hydrochloride 200 mg/day was administered, whereupon the leukocytosis and the increased serum C-reactive protein level went back to normal limits within 1 week. The medication was continued for 2 weeks until the skin eruption disappeared. The PCR assay for the *gltA* gene and the 17 kDa genus-common antigen gene became negative 1 week after administration of minocycline hydrochloride. Neither fever nor lymphadenopathy was observed throughout the clinical course.

Serum IgM and IgG antibody titers against the following rickettsiae were measured in series by indirect immunofluorescence technique [8]: *R. tamurae* (AT-1 strain), *Rickettsia japonica* (YH strain), *Rickettsia typhi* (Wilmington strain), as well as titers against *O. tsutsugamushi* (Karp, Gilliam, Kuroki, Kawasaki, and Kato strains). The patient showed positive antibody titers against *R. tamurae* and *R. japonica* (table 1). The increased *R. japonica* titers might be serological cross-reactions between *R. tamurae* and *R. japonica*.

Discussion

The clinical features of rickettsiosis can vary in intensity, ranging from the mild symptomatic type, e.g. tick-borne lymphadenopathy [9], or *Rickettsia parkeri* infection [10] to the severe symptomatic type, e.g. Japanese spotted fever. Although there has been no report of a patient associated with *R. tamurae* as a human pathogen, this rickettsia has been suspected to have some pathogenic potential. Here we report the first case of local skin inflammation caused by *R. tamurae* in Japan, which was proved by phylogenetic and serological analysis. There have been several case reports of tick bite due to *A. testudinarium* describing local symptoms, such as papules or pustules, without general symptoms like Japanese spotted fever [11]. These reports might include cases of *R. tamurae* infection, because these patients were not examined for rickettsiae.

The present case exhibited mild local inflammatory signs: swelling, erythema and heat with pain. Although these symptoms were cellulitis-like features and we could not exclude other bacterial infections, increased serum antibody titers against *R. tamurae* strongly suggested that the patient actually had a *R. tamurae* infection.

Japanese spotted fever is well known to clinicians; however, most atypical cases of tick bite remain uninvestigated. Serological testing and PCR analysis play a critical role in the diagnosis of rickettsiosis, including *R. tamurae* infection.

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Table 1. Serum IgM and IgG antibody titers of the rickettsiae and *O. tsutsugamushi* measured by the indirect immunofluorescence technique

Species	June 22, 2009	June 29, 2009	July 6, 2009	July 30, 2009	August 27, 2009
<i>R. tamurae</i>					
IgG	<20	<20	20	160	10,240
IgM	20	20	40	160	2,560
<i>R. japonica</i>					
IgG	<20	<20	<20	160	5,120
IgM	20	20	40	160	640
<i>O. tsutsugamushi</i>					
IgG	<20	<20	<20	<20	<20
IgM	<20	<20	<20	<20	<20
<i>R. typhi</i>					
IgG	<20	<20	<20	<20	<20
IgM	<20	<20	<20	<20	<20

Fig. 1. Geographical position of Shimane in Japan. Star: location of the present case.

Fig. 2. Erythema and mild swelling of the left sural area. There was a 15 mm-sized tick in the left popliteal region (a). The engorged tick, which was an *Amblyomma testudinarium* female, had attached itself to the patient's popliteal region (b).

Fig. 3. Phylogenetic tree of SFG rickettsiae derived from the *gltA* gene (**a**) and phylogenetic tree of SFG rickettsiae derived from the 17 kDa genus-common antigen gene (**b**) by the neighbor-joining method. The numbers at nodes are the bootstrap values obtained from 100 re-samplings. The scale bars represent 0.5% (*glt-A* gene) and 1% (17 kDa genus-common antigen gene) differences in nucleotide sequences (GenBank accession numbers are shown).

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