Streptobacillus moniliformis bacteremia in a rheumatoid arthritis patient without a rat bite: a case report

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Abstract

Background: Rat bite fever is a relatively rare infectious disease due to infection with Streptobacillus moniliformis or Spirillum minus mainly via direct bite by rats, mice, or other rodents. If there is no clear bite history, the diagnosis is difficult or may not be made.

Case presentation: A 72-year-old Asian female with rheumatoid arthritis was admitted for high-grade fever and walking difficulty with severe lumbago. Initially, we suspected lumbar compression fracture with deterioration of rheumatoid arthritis, but Gram-negative bacilli were isolated from blood culture during hospitalization. The isolated organism was identified as S. moniliformis by 16S ribosomal ribonucleic acid (rRNA) sequencing. S. moniliformis is well known to be a primary causative organism of rat bite fever, but this patient had no history of rat bite. Had S. moniliformis bacteremia not been detected, she might have been treated for rheumatic exacerbation.

Conclusion: We emphasize the importance of performing appropriate microbial culture testing for identifying potential infectious diseases. We also conclude that S. moniliformis infection can become established with contaminated vehicle contact alone, not only as a direct result of a bite. We must keep mind that those working in places where rodents breed or are at risk of contact with rats or mice might be at risk for contracting this unusual disease.

Keywords: Streptobacillus moniliformis, Bacteremia, Rheumatoid arthritis, Rat bite, Zoonosis

Background

Rat bite fever is a relatively rare infectious disease due to either Streptobacillus moniliformis or Spirillum minus [1–5]. It presents as an acute or chronic illness with polyarthritis in states such as sepsis and bacteremia [4]. Rat bite fever occurs due to a direct bite by an infected and/or colonized rat, mouse, or other rodent or via intake of contaminated food or milk [4]. Most cases have a history of bites or contact with rodents, but the diagnosis in those without such a history must be based on microbiologic testing. We herewith describe an interesting case of S. moniliformis bacteremia without rat bite. This patient had rheumatoid arthritis. Inexplicable joint and bone symptoms closely mimicked those of S. moniliformis bacteremia. The diagnosis would thus have been very difficult unless microbial culture tests had revealed the isolate to be S. moniliformis.

Case report

A 72-year-old Asian female presented with high-grade fever, 38 °C, and chills which had been present for 8 days prior to admission. She had been followed for rheumatoid arthritis (RA) by the orthopedics department of our hospital for 3 years. At the first visit for febrile episodes, upper respiratory tract infection was diagnosed and she was told to rest and stay in bed. She went home without antipyretics, analgesics or antibiotics. However, she reported worsening lower back pain 3 days after the first visit. Subsequently, she was admitted due to difficulty moving, especially walking, because of severe lower back pain. Furthermore, the presence of a severe inflammatory
disease, such as an infection, was suspected based on blood examination results on admission (Table 1).

Her medical history was unremarkable except for hypertension and RA, reasonably well controlled with low-dose oral steroids and weekly methotrexate. Her familial history was also unremarkable. She neither smoked nor drank alcohol regularly and had no history of caring for animals. Moreover, she had no history of allergies to drugs, foods, or inhaled substances. She ran a dining room in downtown Tokyo. She had undergone dental implant insertions, but had been free of dental caries and periodontal disease for several years.

At the beginning of hospitalization, she was unable to walk because of severe sharp pain, affecting the entire body, including lower back pain which been present prior to admission. She was unable to state definitely where in her body the pain was strongest. She was started on antibiotics (cefazolin 2.0 g/every 8 h) and a nonsteroidal antiphlogistic balm for the pain. Her condition subsequently improved with resolution of the high grade fever spikes, but a low grade fever persisted.

On the 2nd hospital day, highly pleomorphic, filamentous gram-negative bacilli were isolated from anaerobic blood culture of the clinical sample obtained on the day of admission (Fig. 1a). Isolates grew on both Brucella with Hemin and Vitamin K1 (HK) agar (Kyokuto Pharmaceutical Industrial Co., Ltd., Tokyo, Japan) and 5 % sheep blood agar under both aerobic and anaerobic conditions. Those samples could be subcultured at 37 °C on 5 % sheep blood agar plates (Eiken Chemical Co., Ltd.) in a capnophilic atmosphere containing 5 % CO2 (Fig. 1b, c) according to previous reports [6, 7]. We obtained a colony approximately 1 mm in diameter in 3 days using these media. Though the isolate was not identified by biological techniques, we accurately identified the strain with 16S ribosomal ribonucleic acid (rRNA) genotyping, as previously described [8, 9], and a similarity search was conducted using the EzTaxon (http://www.ezbiocloud.net/eztaxon/). The isolate (GenBank accession no. LC062896) showed 100 % (1455/1455) similarity to the \textit{Streptobacillus moniliformis} type strain DSM 12112T (GenBank accession no. CP001779). A phylogenetic tree containing all available 16S rRNA sequences was constructed from a multiple sequence alignment with the neighbor-joining method using Molecular Evolutionary Genetics Analysis (MEGA) software, version 5.1 (Additional file 1: Figure). We performed antimicrobial susceptibility tests on the isolate employing both the microdilution method using MicroScan® Negacombo 3.12 (SIEMENS Inc. Berlin, Germany) and using lysed horse blood broth with microdilution, but we were unable to measure minimum inhibitory concentrations against several antibiotics, because there was no growth of the isolate even in the control well. Thus, though we switched the administered antibiotic to

### Table 1 Laboratory data on admission

| Blood cell counts |  |  |  |
|---|---|---|---|
| White blood cells | 13,300/μL |  |  |
| Neutrophils | 95.0 % |  |  |
| Lymphocytes | 2.3 % |  |  |
| Red blood cells | 367 × 10^4/μL |  |  |
| Hemoglobin | 12.2 g/dL |  |  |
| Platelets | 31.1 × 10^4/μL |  |  |

| Biochemical parameters |  |  |  |
|---|---|---|---|
| Aspartate aminotransaminase | 45 IU/L |  |  |
| Alanine aminotransferase | 34 IU/L |  |  |
| Lactose dehydrogenase | 202 IU/L |  |  |
| Alkaline phosphatase | 1035 IU/L |  |  |
| γ-Glutamyl transferase | 239 IU/L |  |  |
| Sodium | 136 mEq/L |  |  |
| Potassium | 4.5 mEq/L |  |  |
| Chloride | 100 mEq/L |  |  |
| Blood urea nitrogen | 29.9 mg/dL |  |  |
| Creatinine | 1.09 mg/dL |  |  |
| Total protein | 6.0 g/dL |  |  |
| Albumin | 2.4 g/dL |  |  |
| Serology |  |  |  |
| C reactive protein | 26.92 mg/dL |  |  |
sulbactum/ampicillin (3.0 g/every 6 h) on the 2nd hospital day based on interim report of blood culture test, we had initially administered ampicillin (2.0 g/every 6 h) based on previous reports describing antimicrobial susceptibilities of similar isolates [2, 10]. Though overall status including the fever showed improvement with antibiotic administration, the lower back pain persisted. She thus underwent magnetic resonance imaging (MRI) of the lumbar vertebrae. The MRI at the levels of the 3rd and 4th lumbar vertebrae showed a low signal on T1-weighted images (Fig. 2a), whereas signal intensity was high within the intranuclear cleft of the disk on short T1 inversion recovery (STIR) T2 imaging and disc extrusion with a high water content was observed adjacent to the extra-dural area (Fig. 2b). The vertebral endplates at this level were destroyed and high-signal-intensity bone marrow edema was visible. These findings are typical of pyogenic vertebral spondyloïd-spondylodiscitis, but there were no signs of crush injury or transformation in the lesion. We continued the antibiotic administration and confirmed gradual improvement of her lower back pain. Furthermore, exercise capacity, including walking, gradually recovered. Follow-up MRI studies showed no deterioration of the lumbar lesions, and she was discharged on the 71st hospital day.

Discussion

Rat bite fever has been a recognized zoonosis since ancient times in India, more than 2000 years ago. However, unexpectedly, this disease was reported again in the medical literature more recently. In 1840, Wilcox reported rat bite fever for the first time [11]. At the end of the 19th century, Miyake provided the first detailed description in the German literature [12]. In this report, he presented this disease as a definite clinical entity based on a series of 11 cases in his own experiences and prior references in the Japanese literature describing Sodoku, which means rat poisoning in Japanese.

*Streptobacillus moniliformis*, present in some rodent intraoral resident floras, is an aerobic or facultative anaerobic gram-negative, highly pleomorphic, filamentous, nonmotile, and non-acid-fast rod [2]. *S. moniliformis*, as well as *Spirillum minus*, transmitted by bite wounds and scratches from rodents such as mice, can cause systemic rat bite fever. This condition is due to infection by *S. moniliformis* or, less commonly, by *S. minus*. Disease caused by *S. minus* is generally known as Sodoku and occurs primarily in Asia [2]. After the incubation period of 2–10 days, symptoms progress to high fever, arthralgia of multiple joints, muscle ache, and whole body rash [1, 13–16]. In previous reports, arthritis occurred mainly in large joints including the elbows, knees and mid-back, while rashes of the palms or soles and were erythematous [3, 13, 14, 16–19]. Such skin lesions were occasionally accompanied by abscess formation [2]. On the other hand, *S. moniliformis* was reported to be a causative pathogen of infectious endocarditis [1, 20–24] or to be associated with immunocompromised status including human immunodeficiency virus (HIV) infection and hematological malignancies [24–26]. Furthermore, there

![Fig. 2](image-url)
are reports describing occurrence in patients with bacteremia associated with exposure [27, 28], such as working in laboratories [29, 30].

In the present case, fortunately, we were able to identify the causative organism from blood cultures. Thus, we provided appropriate treatment and were able to discharge the patient. An important point is whether she had suffered a rodent bite, but no such bite episode was revealed, despite a detailed interview. There are actually “unusual” case reports without a bite history [31] or febrile episode [5]. However, despite the lack of a clear bite history, we suspected S. moniliformis infection based on other aspects of the patient’s presentation.

She ran a dining room serving Tonkatsu, a Japanese dish which consists of a breaded, deep-fried pork cutlet, in Akihabara, an area of downtown Tokyo, and she had once seen a rat in the dining room kitchen. We considered the possibility of contracting the infection solely by contact with contaminated water and/or food. In a previous report, an outbreak of S. moniliformis blood stream infections occurred in a boarding school and the authors concluded the vehicle of infection to be rats contaminating the water supply [32]. Indeed, S. moniliformis infection is caused not only directly by rodent bites but also via contaminated water.

As for major changes in the global environment, such as deforestation and clearing for farmland as well as global climate change due to the greenhouse effect caused by the rise in carbon dioxide concentration, the risk of zoonoses will increase worldwide. However, epidemiological analysis will be difficult even if zoonosis increases, because accurate diagnosis of this disease is often difficult. The anticoagulant included in the commercial blood culture bottle reportedly inhibits the growth of S. moniliformis, and there are numerous cases lacking an accurate diagnosis [14, 33, 34]. In cases with a bite history, rat bite fever is really suspected, but in those without a bite history, the diagnosis must always be based on microbiological testing alone. An accurate diagnosis cannot be obtained without an isolate. A culture test and the improvement of the device for detecting S. moniliformis are highly anticipated.

Conclusions

We conclude that S. moniliformis infection can become established with contaminated vehicle contact alone, not only as a direct result of a bite. Those working in places where rodents breed or are at risk of contact with rats or mice might be at risk for contracting this unusual disease. The cultural diversification of large urban centers is becoming increasingly complex. Thus, we must keep in mind the possibility of zoonosis.

Consent

Written informed consent was obtained from the patient for publication of this Case Report and any accompanying images.

Additional file

Additional file 1: Figure. A phylogenetic tree containing all available 16S rRNA sequences.

Abbreviations

RA: rheumatoid arthritis; rRNA: ribosomal ribonucleic acid; MRI: magnetic resonance imaging; STIR: short T1 inversion recovery.

Authors’ contributions

Conception and drafting of the report; TN, AS, and KT, medical management; AS, TN, and KT. Pharmacotherapeutic support (including therapeutic drug monitoring); YM, microbial testing support; KS, sequence study of isolate; RS. All authors read and approved the final manuscript.

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Competing interests

The authors declare that they have no competing interests.

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References

1. Madhubashini M, George S, Chandrasekaran S. Streptobacillus moniliformis endocarditis: case report and review of literature. Indian Heart J. 2013;65:442–6.
2. Elliott SP. Rat bite fever and Streptobacillus moniliformis. Clin Microbiol Rev. 2007;20:13–22.
3. Dendle C, Woolley L, Korman TM. Rat-bite fever septic arthritis: Illustrative case and literature review. Eur J Clin Microbiol Infect Dis. 2006;25:7917.
4. Wullenweber M. Streptobacillus moniliformis—a zoonotic pathogen. Taxonomic considerations, host species, diagnosis, therapy, geographical distribution. Lab Anim. 1995;29:1–15.
5. Stehle P, Dubuis O, So A, Dudler J. Rat bite fever without fever. Ann Rheum Dis. 2003;62:894–6.
6. Woo PC, Wu AK, Tsang CC, Leung KW, Ngan AH, Cunneen SQ, Lam KW, Chen JH, Chan JF, Lau SK. Streptobacillus hongkongensis sp. nov., isolated from patients with quinsy and septic arthritis, and emended descriptions of the genus Streptobacillus and Streptobacillus moniliformis. Int J Syst Evol Microbiol. 2014;64(Pt 9):3034–9.
7. Eisenberg T, Gläser SP, Nicklas W, Mauder N, Contzen M, Aledelbi K, Kämpfer P. Streptobacillus felis sp. nov. isolated from a cat with pneumonia. Int J Syst Evol Microbiol. 2015;65:2172–8.
8. Nei T, Akutsu K, Shima A, Tsuboi I, Suzuki H, Yamamoto T, Tanaka K, Shinozawa A, Kojima Y, Washio Y, Okawa S, Sonobe K, Norose Y, Saito R. A case of streptococcal toxic shock syndrome due to Group G streptococci identified as Streptococcus dysgalactiae subsp. equisimilis. J Infect Chemother. 2012;18:915–7.

9. Edwards R, Finch RG. Characterisation and antibiotic susceptibilities of Streptobacillus moniliformis. J Med Microbiol. 1986;21:39–42.

10. Wilcox M. Violent Symptoms from the Rat Bite. Am J Med Sci. 1840;26:245–6.

11. Miyake H. Ueber Die Rattenbisskrankheit. Mittelungen aus den Grenzgebieten der Medizin und Chirurgie. 1899;5:231 (German).

12. Flannery DD, Akinboyo I, Ty JM, Averill LW, Freedman A. Septic arthritis and concern for osteomyelitis in a child with rat bite fever. J Clin Microbiol. 2013;51:1987–9.

13. Hockman DE, Pence CD, Whittler RR, Smith LE. Septic arthritis of the hip secondary to rat bite fever: a case report. Clin Orthop Relat Res. 2000;380:173–6.

14. Budar B, Goswami K, Dhukaram V. Septic arthritis secondary to rat bite fever: a challenging diagnostic course. BMJ Case Rep. 2014. doi:10.1136/bcr-2014-204086.

15. Shanson DC, Pratt J, Greene P. Comparison of media with and without Panmede for the isolation of Streptobacillus moniliformis from blood cultures and observations on the inhibitory effect of sodium polyanethol sulphonate. J Med Microbiol. 1985;19:181–6.

16. Palarasah Y, Skjoedt MO, Vitved L, Andersen TE, Skjødt K, Koch C. Sodium polyanetholate suflonate as an inhibitor of activation of complement function in blood culture systems. J Clin Microbiol. 2010;48:908–14.