CASE REPORT

Non-tuberculosis Mycobacterium Tenosynovitis with Rice Bodies in a Patient with Systemic Lupus Erythematosus

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Abstract:
Infectious disease with various presentations in systemic lupus erythematosus often resembles lupus flare. A 37-year-old woman presented with a swollen left index finger that had not resolved, despite 7 years of immunosuppressive treatment. MRI showed rice-body formation in the flexor tendon sheath and tenosynovectomy demonstrated chronic synovitis with epithelioid granuloma. A mycobacterial culture confirmed invasive mycobacterial tenosynovitis due to Mycobacterium chelonae. The patient was treated with moxifloxacin and clarithromycin and completely recovered.

Key words: tenosynovitis, non-tuberculosis mycobacterium, systemic lupus erythematosus, Mycobacterium chelonae, rice-body formation

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Introduction
Systemic lupus erythematosus (SLE) is an autoimmune disease with diverse clinical manifestations that is characterized by remission and flares. Arthritis is one of the most common disease manifestations in SLE. Infectious arthritis in SLE often resembles a disease flare and hence is difficult to diagnose. Rice-body formations, which could be detected by imaging modalities such as magnetic resonance imaging (MRI), have been reported in non-tuberculosis mycobacterium (NTM) tenosynovitis (1, 2). We herein report a case of refractory arthritis in an SLE patient, who was diagnosed with NTM tenosynovitis after the detection of rice bodies on MRI and the confirmation of Mycobacterium chelonae on a lower temperature tissue culture. The patient was successfully treated with moxifloxacin and clarithromycin.

Case Report
A 37-year-old Japanese woman with a 25-year history of SLE, presented with swelling of the proximal interphalangeal (PIP) joint of the left index finger, which had been present since she was 30 years of age. She has been treated with prednisolone (PSL; 15 mg) and mizoribine (MZB; 150 mg) in recent years. The first manifestations were butterfly rash, arthritis, lupus nephritis and leukopenia, and she was treated with steroid pulse therapy and cyclophosphamide. The swelling of the PIP joint initially improved with immunosuppressive therapy at 33 years of age. However, it relapsed with hypocomplementemia and an elevated anti-DNA antibody titer at 35 years of age. After the addition of tacrolimus, the patient’s laboratory data showed improvement, but the joint swelling remained unchanged. A radiograph of the left index finger was normal, with the exception of swelling. Ultrasonography of the left index finger revealed hypoechoic soft-tissue masses along the flexor tendon.

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(Fig. 1A) with multiple internal hyperechoic ‘flecks’ (Fig. 1B). Fluid-sensitive MRI of the left index finger revealed numerous tiny nodules of low signal intensity (Fig. 1C, D), which had no contrast enhancement effect (Fig. 1E). These findings were compatible with rice body formation in the flexor tendon sheath. Post-contrast enhancement of the flexor tendon sheath and thickened synovium adjacent to the flexor tendon was demonstrated (Fig. 1E). There were no findings of NTM infection in other sites. Synovectomy was performed and rice-body formation was found (Fig. 2A, B). The histopathological examination of the mass lesion demonstrated chronic synovitis complicated by epithelioid granuloma (Fig. 2C) with increased numbers of fibroblast-like synoviocytes (Fig. 2D). Acid-fast staining was negative. However, a colony was detected on a mycobacterial culture incubated at 28-32°C. *M. chelonae* was identified by matrix-assisted laser desorption/ionization time-of-flight mass spectrometry (MALDI-TOF MS) and genome sequencing, pointing to a diagnosis of tenosynovitis of the hand due to *M. chelonae*. A drug sensitivity test was performed using the microdilution method as follows: Imipenem/Cilastatin: S (MIC=4.0), Clarithromycin: S (MIC=0.12), Minocycline: R (MIC>8.0), Levofloxacin: R (MIC=4.0). Moxifloxacin and clarithromycin were administered for 6 months. The swelling disappeared, and the remission has been maintained.

**Discussion**

NTM is a ubiquitous environmental organism found in water and animals, that causes chronic infection in humans (3). A previous report indicated that approximately 7% of patients with NTM infection were immunocompromised, including autoimmune disease patients treated with immunosuppressive drugs (4). Further, a majority of these infections occurred after trauma (5). In a meta-analysis, all patients presented with swelling and pain, and 74% of all cases of hand tenosynovitis involved only one finger (4). Although we need to be aware that the epidemiology of NTM infection varies due to regional differences, a literature review described that the most common organisms of NTM in hand infection is *M. marinum* (82%), followed by *M. chelonae* (5%), *M. kansasii* (3%), and *M. intracellulare* (3%) (4). *M. chelonae* is a rapidly growing mycobacteria (RGM) belonging to Group 4 of the Runyon classification (6). *M. chelonae* has been reported in association with hand infection more than other RGMs, such as *M. abscessus* (7). As acid-fast staining is usually negative in these infections, special Ziehl-Neelsen staining and incubation at 28-32°C in Lowenstein-Jensen medium must be performed (5). *M. chelonae* is iden-
Early surgery and appropriate culture guided antibiotic therapy are crucial in the management of NTM hand tenosynovitis. A combination antibiotic therapy is recommended. The use of amikacin, ciprofloxacin, and clarithromycin was reported in previous studies. Clarithromycin was reported to be the most active agent in inhibiting the growth of *M. chelonae* (9). The duration of treatment should be at least 6 months, but should be no longer than 12 months (4).

A cohort study reported that NTM infection occurred in 1.5% of SLE patients (10). In a review of the relevant literature on NTM infections in SLE patients (11), *M. chelonae* (26.8%) was the most common, followed by *M. kansasii* (21.4%) and *M. avium complex* (14.3%). Extra-pulmonary lesions are more common than tuberculosis infection. Skin and soft tissue infection (55.4%) are the most common locations, followed by the joints (23.2%), and lung (21.4%) (11). However, the frequency of NTM species and the prevalence of infection sites may be different in Japan. Bone marrow infection, which causes pancytopenia has also been reported (12). These manifestations resemble a disease flare. Thus, when the immunosuppressive treatment is ineffective, NTM infection should be considered as a differential diagnosis.

Rice bodies caused in tenosynovitis are reported in rheumatoid arthritis, tuberculosis, and NTM infection (1). The pathogenesis of rice body formation remains unclear, but experts have suggested that they arise from microinfarcted synovium or that the early rice body appears *de novo* and causes progressive enlargement by fibrin deposition (13). MRI is considered the imaging modality of choice for rice bodies, which appear iso- or slightly hyperintense in comparison to skeletal muscle on T1- and T2-weighted imaging (1). In our case, the MRI findings were not typical and required further diagnostic steps.

Based on our experience, we encourage clinicians to suspect NTM infection when patients with SLE present refractory mono-arthritis that is resistant to immunosuppressive treatment. Rice bodies detected on MRI assisted our diagnosis and lower temperature tissue culturing was needed to confirm the diagnosis of *M. chelonae*, which is a common NTM infection in SLE patients.

The authors state that they have no Conflict of Interest (COI).
Yuichiro Fujieda and Keita Ninagawa contributed equally to this work.

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