

Pyogenic tenosynovitis of the wrist due to *Corynebacterium striatum* in a patient with dermatomyositis

A case report

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Abstract

Introduction: Corynebacterium striatum is common contaminant in clinical specimens. Here, we report a rare case of pyogenic tenosynovitis of the wrist caused by C striatum in a dermatomyositis patient taking oral immunosuppressants.

Patient concerns: A 67-year-old Japanese woman with dermatomyositis had a history of multiple intraarticular injections of corticosteroids to the right wrist joint for the treatment of osteoarthritis. She was admitted to our hospital with a painful lump on the right dorsal wrist lasting for three months. MRI revealed cellulitis of the dorsum of the right wrist and hand and fluid collection in the extensor tendon sheath. *C striatum* was detected in the cultures of three samples of synovial fluid taken from the dorsal hand.

Diagnosis: Pyogenic tenosynovitis of the wrist due to *C* striatum.

Interventions: The infection was successfully controlled with synovectomy and adjuvant antibiotic therapy.

Outcomes: There has been no sign of recurrence for 12-months after the surgical treatment.

Lessons: This is the first reported case of pyogenic tenosynovitis due to *C striatum* in a patient with dermatomyositis. Clinicians should be aware that patients undergoing immunosuppressive therapy have a risk of *C striatum* infection.

Abbreviations: CAM = clarithromycin, CRP = C-reactive protein, MRI = magnetic resonance imaging, WBC = white blood cell count, WI = weighted image.

Keywords: corticosteroid injection, Corynebacterium striatum, dermatomyositis, immunosuppressants, pyogenic tenosynovitis

1. Introduction

Corynebacterium species are normal human flora that are common contaminants in clinical specimens. Except for *Corynebacterium diphtheriae*, all *Corynebacterium* spp. are ubiquitous skin commensals.^[1] However, some reports have highlighted the importance of *C striatum* as causes of catheter-related bloodstream

Editor: N/A.

The authors have no funding and conflicts of interests to disclose.

The patient gave written consent for publication of her potentially identifying information (including individual details and images).

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How to cite this article: Kawaguchi Y, Okamoto H, Endo K, Iwata H, Joyo Y, Nozaki M, Tamechika S, Waguri-Nagaya Y, Murakami H. Pyogenic tenosynovitis of the wrist due to Corynebacterium striatum in a patient with dermatomyositis: A case report. Medicine 2020;99:3(e18761).

Received: 10 April 2019 / Received in final form: 14 November 2019 / Accepted: 10 December 2019

http://dx.doi.org/10.1097/MD.000000000018761

infections and various other infections, such as arthritis and endocarditis, in immunosuppressed patients.^[2–5] Here, we report a rare case of pyogenic tenosynovitis of the wrist caused by *C striatum* in a patient with dermatomyositis who was taking oral immunosuppressants. The infection was successfully controlled with synovectomy and adjuvant antibiotic therapy.

2. Case presentation

A 67-year-old Japanese woman who had had dermatomyositis for 4 years was diagnosed with a typical rash (heliotrope rash, Gottron's sign), symmetric proximal muscle weakness, elevated serum skeletal muscle enzymes, and interstitial lung disease. She had a past history of pleurisy caused by Mycobacterium abscessus and mycotic pneumonia. Her dermatomyositis was controlled with prednisolone 5 mg/day and azathioprine 75 mg/ day for 3 years. The patient had received clarithromycin (CAM) (400 mg/day) for the treatment of an *M* abscessus infection and itraconazole (100 mg/day) for the treatment of mycotic pneumonia for 1 year, in addition to trimethoprim/sulfamethoxazole (160/800 mg twice weekly) as a prophylaxis for opportunistic infections for 4 years. She had a history of multiple injections of corticosteroids into the right wrist for arthralgia due to osteoarthritis for 6 months. Prior to admission, the patient had a 3-month history of a painful lump on the dorsal area of the right wrist, that had gradually increased in size and finally measured approximately 3×5 cm (Fig. 1A). The range of motion of the right radiocarpal joint was limited by pain. The patient was



Figure 1. Preoperative appearance and plain radiograph of the right wrist. (A) Lump on the dorsal area of the right wrist and hand (black arrow). (B) Plain radiograph of the right wrist reveals sclerotic changes of the carpal bones and narrowing of the radiocarpal joint space.

not able to extend the right ring and little fingers. *C striatum* was detected in 3 samples of synovial fluid aspirated from the lump. This was resistant to penicillin and ceftriaxone but sensitive to imipenem, amikacin, vancomycin and rifampicin. Blood biochemistry showed slightly elevated C-reactive protein levels (CRP: 0.47 mg/dl) and a normal white blood cell count (WBC: 7200/mm³) (Table 1). Blood cultures showed no evidence of bacteremia. Plain radiographs of the right wrist revealed narrowing of the radiocarpal and midcarpal joints accompanied

Table 1	
Laboratory findings on admissio	on.

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White blood cells	7200/µl	TP	7.9 g/dl	
Neutrophils	94.3%	Alb	3.8 g/dl	
Lymphocytes	3.2%	AST	20 U/I	
Eosinophils	0%	ALT	6 U/I	
Monocytes	2.2%	LDH	273 U/I	
Red blood cells	$359 imes 10^4 / \mu l$	CRP	0.47 mg/dl	
Hemoglobin	11.0 g/dl	CK	29 U/I	
Platelet count	$23.1 \times 10^4 / \mu$ l	BUN	22.4 mg/dl	
<coagulation></coagulation>		Cre	0.59 mg/dl	
PT	120.7%	Na	141 mEq/l	
APTT	28.2 seconds	К	4.5 mEq/l	
Fibrinogen	403 mg/dl	CI	105 mEq/l	
D-dimer	1.0 μg/ml	Anti-MDA5 Ab	Positive	

Alb=Albumin, ALT=Alanine aminotransferase, Anti-MDA5 Ab=anti-melanoma differentiationassociated gene 5 antibody, APTT=Activated partial thromboplastin time, AST=Aspartate aminotransferase, BUN=Blood urea nitrogen, CK=Creatine kinase, CI=Chloride, Cre=Creatinine, CRP=C-reactive protein, K=Potassium, LDH=Lactate dehydrogenase, Na=Sodium, PT= Prothrombin time, TP=Total protein. by sclerotic changes of the carpal bones due to osteoarthritis (Fig. 1B). Magnetic resonance imaging (MRI) revealed cellulitis of the dorsum of the right wrist and hand and fluid collection in the extensor tendon sheath (Fig. 2A). Additionally, MRI demonstrated bone marrow edema with low signal intensity on T1 weighted-images (WI) and high signal intensity on T2 WI in the carpal bones (Fig. 2B and C). Plain radiographs of the wrist revealed no osteolytic lesion of the carpal bones, and MRI demonstrated fluid within the fourth extensor compartment tendon sheath surrounding the tendons, not in the joint. She was diagnosed as pyogenic extensor tenosynovitis of the wrist due to C striatum. Therefore, synovectomy was performed through a dorsal approach. The intraoperative findings revealed massive synovitis along the extensor tendons, with complete rupture of the extensor tendons of the ring and little fingers (Fig. 3). However, there was no intraarticular synovitis. Cultures of synovial tissue was negative, because of intraoperative use of antibiotics. Cultures for tubercle bacillus and polymerase chain reaction for atypical mycobacteriosis were all negative. Histopathology showed synovial tissue with diffuse acute inflammatory infiltrate and the deposition of fibrinous material on the synovial surface, without granulomatous changes. Multidrug chemotherapy including imipenem/cilastatin, daptomycin, and amikacin were initiated. These antibiotics were continued for 2 weeks postoperatively, followed by oral rifampicin 450 mg/day for 3 months. After the 6-month follow-up, MRI showed improved signal intensity in the bone marrow of the carpal bone and extinction of the fluid around the extensor tendons (Fig. 4). There has been no sign of recurrence for 12-months after the surgical treatment. Because the patient feels no disability in daily activities, she does not wish to undergo reconstructive surgery of the extensor tendons at this time.



Figure 2. Preoperative MRI of the right wrist. (A) An MRI T2 weighted contrast transverse image reveals diffuse subcutaneous edema in the dorsum of the wrist and fluid collection in the extensor tendon sheath. Coronal image demonstrates the bone marrow edema with low signal intensity in the T1 WI (B) and high signal intensity in the T2 WI (C) in the bone marrow of the carpal bones. MRI = magnetic resonance imaging, WI = weighted image.

3. Discussion

Pyogenic tenosynovitis frequently occurs subsequent to a puncture wound. Therefore, skin flora such as *Staphylococcus aureus* and *Streptococcus species* are found to be the causative organism in most cases. Atypical organisms including gramnegative rods and mixed flora have been found in immunocompromised patients.^[6]

We presented a case of pyogenic tenosynovitis of the wrist due to *C striatum* in a patient with dermatomyositis who was taking oral immunosuppressants. There have been few reports of pyogenic arthritis and osteomyelitis due to *C striatum*.^[4,7,8] To



Figure 3. Intraoperative findings. Massive synovitis along the extensor tendons and complete rupture of the extensor tendons of the ring and little fingers.

the best of our knowledge, no case of pyogenic tenosynovitis of wrist due to *C striatum* has been reported.

Corynebacterium species are opportunistic human pathogens that are distributed in the skin and mucous membranes of asymptomatic individuals. These organisms are often considered contaminants when isolated in culture because of their low virulence.^[9] However, when isolated repeatedly in pure growth from a normally sterile body site, such as the synovial fluid or blood, in a clinical context consistent with infection, they should be considered clinically relevant, and identification at the species level and antimicrobial susceptibility testing are recommended.^[9] In our case, the first bacterial culture of the fluid was thought to indicate skin contamination; however, *C striatum* was repeatedly detected in the cultures of aspirated material. Therefore, we considered *C striatum* the pathogenic microorganism.

Corticosteroids results in decreased fibrosis and reduced vascular proliferation in peritendinous tissues, along with antiinflammatory effects that include decreased leukocyte migration and phagocytosis.^[10] Saglam et al reported an intratendinous septic abscess of the Achilles tendon after multiple local steroid injections and considered that the abovementioned effect may increase the possibility of infection at the site of local corticosteroid injection.^[11] Even in our case, this effect may have been the basis for the infection.

Previously reported risk factors for *C striatum* infection include a history of collagen disease, diabetes mellitus and immunosuppressants therapy.^[4,7,12,13] In this case, the patient received prednisolone and an immunosuppressant for approximately 3 years for the treatment of dermatomyositis. Additionally, we concluded that the main cause of the infection was the iatrogenic inoculation of *C striatum* from the bacterial flora of the skin into the extensor tendon sheath.

Treatment of *C striatum* infection includes initiation of appropriate antibiotic therapy and involve possible removal of



Figure 4. T2 weighted magnetic resonance imaging images at the 6-month follow-up. (A) Synovitis and fluid collection around the extensor tendons disappeared. (B) Signal intensity in the bone marrow of the carpal bones was improved.

infectious lesion to ensure complete eradication of the nidus of infection. There is growing concern over resistance of *C striatum* to penicillins, and cephalosporines,^[14] as seen in our case, even though previously reported susceptibility to penicillins.^[15] Repeated isolation of *C striatum* from a sterile body site should prompt early laboratory request for drug-susceptibility testing. Susceptibility testing is not routinely performed otherwise by most laboratories as *C striatum* is considered a contaminant in most clinical situations. Early detection and appropriate treatment of this infection is important especially in immunosuppressed patients and likely to lead to improved clinical outcomes.

In summary, we reported the case of a patient with pyogenic tenosynovitis due to *C striatum* with dermatomyositis who was treated with the immunosuppressant azathioprine. Surgical radical debridement and multidrug chemotherapy were useful. Meticulous attention should be paid to aseptic techniques whenever local injections of corticosteroids are administered to patients taking immunosuppressants, and patients should be closely observed during the postinjection period.

Author contributions

Conceptualization: Yohei Kawaguchi.

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- Supervision: Yuko Waguri-Nagaya, Hideki Murakami.
- Writing original draft: Yohei Kawaguchi.
- Writing review & editing: Yohei Kawaguchi, Yuko Waguri-Nagaya.

References

- Yanai M, Ogasawasa M, Hayashi Y, et al. Retrospective evaluation of the clinical characteristics associated with Corynebacterium species bacteremia. Braz J Infect Dis 2018;22:24–9.
- [2] Chen FL, Hsueh PR, Teng SO, et al. Corynebacteriumstriatum bacteremia associated with central venous catheterinfection. J Microbiol Immunol Infect 2012;45:255–8.
- [3] Rozdzinski , Kern W, Schmeiser T, et al. Corynebacterium jeikeium bacteremia at a tertiary care center. Infection 1991;19:201–4.
- [4] Westblade LF, Shams F, Duong S, et al. Septic arthritis of a native knee joint due to Corynebacterium striatum. J Clin Microbiol 2014;52:1786–8.
- [5] Boltin D, Katzir M, Bugoslavsky V, et al. Corynebacterium striatum–a classic pathogen eluding diagnosis. Eur J Intern Med 2009;20:e49–52.
- [6] Draeger RW, Bynum DKJr. Flexor tendon sheath infections of the hand. J Am Acad Orthop Surg 2012;20:373–82.
- [7] Scholle D. A spontaneous joint infection with Corynebacterium striatum. J Clin Microbiol 2007;45:656–8.
- [8] Beltrán-Arroyave C, Díaz-Díaz A, Loaiza-Díaz N. Chronic osteomyelitis due to Corynebacterium striatum in a female adolescent. Rev Chilena Infectol 2016;33:696–9.
- [9] Bernard K. The genus Corynebacterium and other medically relevant coryneform-like bacteria. J Clin Microbiol 2012;50:3152–8.
- [10] Mahler F, Fritschy D. Partial and complete ruptures of the Achilles tendon and local corticosteroid injection. Br J Sports Med 1992;26:7–14.
- [11] Saglam N, Akpinar F. Intratendinous septic abscess of the Achilles tendon after local steroid injection. J Foot Ankle Surg 2009;48:565–8.
- [12] Verma R, Kravitz GR. Corynebacterium striatum empyema and osteomyelitis in a patient with advanced rheumatoid arthritis. BMJ Case Rep 2016;doi: 10.1136/bcr-2016-214691.
- [13] Roy M, Ahmad S. Rare case of Corynebacterium striatum septic arthritis. BMJ Case Rep 2016;doi: 10.1136/bcr-2016-216914.
- [14] Ortiz-Pérez A, Martín-de-Hijas NZ, Esteban J, et al. High frequency of macrolide resistance mechanisms in clinical isolates of Corynebacterium species. Microb Drug Resist 2010;16:273–7.
- [15] Martínez-Martínez L, Pascual A, Bernard K, et al. Antimicrobial susceptibility pattern of Corynebacterium striatum. Antimicrob Agents Chemother 1996;40:2671–2.