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Septic Arthritis of a Native Knee Joint Due to Corynebacterium striatum

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We report a case of septic arthritis of a native knee joint due to *Corynebacterium striatum*, a rare and unusual cause of septic arthritis of native joints. The isolate was identified by a combination of phenotypic, mass spectrometric, and nucleic acid-based assays and exhibited high-level resistance to most antimicrobials.

**CASE REPORT**

A 64-year-old male with a past medical history of poorly controlled diabetes, coronary artery disease, hypertension, deep vein thrombosis, and anticoagulant use presented with right knee pain and fever. A week prior to admission, he had fallen while trying to climb onto a bus.

Four days prior to admission, a right knee arthrocentesis at his primary care doctor’s office, by report, revealed grossly bloody fluid. This was followed by worsening right knee pain upon weight bearing and increasing right knee swelling followed by malaise and subjective fever with chills.

On presentation to the emergency department (ED), he was febrile to 38.5°C. The patient’s right lower extremity was edematous, and examination of the right knee revealed minimal erythema, tenderness to palpation, effusion, and a decreased range of motion secondary to pain. In the ED, his knee was aspirated under sterile conditions and yielded 35 ml of straw-colored cloudy fluid. Analysis of the fluid revealed a small calcium pyrophosphate crystals and a white blood cell count elevated to 52,500/µl with 80% neutrophils. A Gram stain of the specimen was negative for organisms. He was empirically started on vancomycin and ceftazidime and admitted for arthroscopic lavage of the septic knee. He underwent knee lavage in the operating room (OR) 24 h after admission. Two specimens taken from the arthroscopic fluid were cultured on 5% (vol/vol) sheep blood tryptic soy agar plates (Becton, Dickinson and Company, Sparks, MD) incubated at 35°C in 5% carbon dioxide, while one (taken from the ED) was directly inoculated into a blood culture vial (Bactec Plus Aerobic/F culture vial; Becton, Dickinson and Company). All three specimens initially plated to solid media yielded pure cultures of catalase-positive, cream-colored colonies within 48 h, while the blood culture vial was positive within 24 h and, upon subculture, grew a pure culture of catalase-positive, cream-colored organisms. Gram stain of all four isolates revealed pleomorphic, palisading Gram-positive rods.

The isolates were initially identified as *Corynebacterium striatum* with >99% probability using the RapID CB Plus phenotypic system (Remel, Lenexa, KS) (1). Subsequently, the isolates were analyzed by matrix-assisted laser desorption–ionization time of flight mass spectrometry using the recently U.S. Food and Drug Administration (FDA)-cleared Vitek MS v2.0 system (bioMérieux, Durham, NC) (2) and were identified as *C. striatum* with a confidence level of 99.9%. Finally, for one of the isolates, fragments of the 16S rRNA gene (3) and the *rpoB* gene (4) were amplified using the PCR and the PCR products sequenced. However, although they are highly accurate for bacterial identification, neither of these sequence-based methodologies is cleared by the FDA for bacterial identification and both remain restricted to research use only.

The resultant 16S rRNA gene sequence data were analyzed using SmartGene IDNS (Integrated Database Network System) software (SmartGene GmbH, Lausanne, Switzerland) (5), and the results revealed that the best match was *C. striatum* type strain ATCC 6940, with 99.6% similarity (470 bp/472 bp), while the next-best match was *C. xerosis*, with 98.9% similarity (467 bp/472 bp). According to the standards adopted by the Clinical Laboratories Standards Institute (CLSI), the 16S rRNA gene sequence analysis satisfied genus-level but not species-level identification.
requirements (6). The rpoB gene sequence data were queried against GenBank (http://www.ncbi.nlm.nih.gov) and the European Nucleotide Archive (ENA; http://www.ebi.ac.uk/ena/). The best match returned from GenBank was C. striatum type strain CIP 81.15, with 97.3% similarity (434 bp/446 bp), while the next-best match was C. simulans type strain CIP 106488, with 94.0% similarity (419 bp/446 bp). The ENA returned a best match of C. striatum type strains 6940 and CIP 81.15, both with 97.3% similarity (434 bp/446 bp). The next-best match was C. simulans type strain CIP 106488, with 94.0% similarity (419 bp/446 bp). Taken together, the 16S rRNA gene and rpoB gene sequence data strongly support the phenotypic and mass spectrometric identification of C. striatum.

Antimicrobial susceptibility testing results for all four isolates were determined using an Etest (bioMérieux) method that was verified against the published CLSI test conditions (7). A 0.5 McFarland standard was prepared and cultured on Mueller-Hinton agar supplemented with 5% (vol/vol) sheep blood (Remel) for 48 h. Using breakpoints established by the CLSI (7), Etest values revealed that all four isolates were resistant to clindamycin, ciprofloxacin, tetracycline, and ceftriaxone, with MIC values > 256 µg/ml, while all four isolates were susceptible to vancomycin (MIC of 1 µg/ml).

Corynebacterium species are opportunistic human pathogens, and due to their association with skin and mucous membranes in asymptomatic individuals, these organisms are often considered contaminants when isolated in culture (8). However, when isolated repeatedly in pure growth from a normally sterile body site, e.g., synovial fluid or blood, in a clinical context consistent with infection, they should be considered clinically relevant and identification to the species level and antimicrobial susceptibility testing is recommended (8).

C. striatum has been associated with invasive infections, including infective endocarditis, pulmonary infections, and prothetic joint infections (9–12). However, to the best of our knowledge, there are only four cases in the published literature describing septic arthritis of native joints due to C. striatum (PubMed [http://www.ncbi.nlm.nih.gov/pubmed]; terms “corynebacterium,” “striatum,” “septic,” and “arthritis”) (11, 13–15), including two cases of septic arthritis of the shoulder (11, 15), one case of a septic elbow (13), and one case of septic arthritis of the knee (14). Thus, as far as we are aware, the case presented here is only the second case of septic arthritis of a native knee joint due to C. striatum.

The report of a previous case of native knee septic arthritis describes an 87-year-old male with a history of osteoarthritis and advanced heart failure who, following a fall, presented with a swollen knee (14). Aspirated synovial fluid revealed inflammatory cells without crystals, and both synovial fluid and blood cultures were negative. Approximately 3 weeks later, he returned with pneumonia due to Streptococcus pneumoniae and an inability to bear weight. Again, aspirated synovial fluid revealed inflammatory cells without crystals; however, upon two occasions C. striatum was recovered in pure culture within 24 h of culture inoculation. Interestingly, rather than attributing the infection due to direct inoculation of skin-associated C. striatum during aspiration of the joint, it was suggested that the infection was spontaneous and that the offending C. striatum isolate gained access to the patient’s circulation either during the episode of pneumonia or through open venous stasis ulcers.

There are significant similarities between our case and the aforementioned case, namely, the blunt trauma of the knee prior to presentation and the underlying immunosuppression associated with the patients. However, rather than interpreting ours as a second case of spontaneous infection of a joint due to C. striatum, we believe that iatrogenic inoculation of the joint with skin-associated C. striatum during the first knee aspiration likely resulted in the infection described in our case.

The identification of Corynebacterium species to the species level is often difficult or unreliable if phenotypic testing is the sole identification method utilized (8, 9). Therefore, to confirm the phenotypic identification of C. striatum, we utilized mass spectrometric and nucleic acid-based methodologies, with both methodologies convincingly identifying the isolates as C. striatum. Mass spectrometric identification of microbes, including Corynebacterium species, is revolutionizing the fields of clinical microbiology and infectious diseases and has the ability to rapidly identify Corynebacterium species to a level comparable to that achievable with the more labor-, time-, and cost-intensive sequence-based methods (16).

Antimicrobial susceptibility testing of Corynebacterium species should be performed if the isolate is considered clinically relevant, as antimicrobial susceptibility is not predictable on the basis of genus- and species-level identification (8). This is partially due to the fact that, historically, many laboratories were unable to reliably identify coryneform bacteria to the species level. Further, Corynebacterium species, especially C. striatum, demonstrating multidrug resistance have been recovered from clinical specimens, with isolates displaying resistance to several classes of antimicrobials, including beta-lactams, fluoroquinolones, macrolides, lincosamides, and tetracyclines. Typically, these multidrug-resistant isolates are susceptible only to vancomycin, daptomycin, and linezolid (8, 14). The isolates obtained from our patient were multidrug resistant; of all the antimicrobials assayed, vancomycin was the only antimicrobial that tested as susceptible.

This case further highlights the role of C. striatum in native joint infections. Additionally, it emphasizes the importance of identifying Corynebacterium species isolates recovered in multiple cultures to the species level and performing antimicrobial susceptibility testing due to the increased frequency of multidrug resistance in this genus.

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