Authors have no conflicts to disclose.

Multi-focal *Clostridium difficile* Osteomyelitis in a Patient with Sickle Cell Anemia: Case Presentation and Literature Review

Multi-focal *Clostridium difficile* Osteomyelitis

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Abstract

This is the author's manuscript of the article published in final edited form as:


https://doi.org/10.1016/j.diagmicrobio.2019.114915
*Clostridium difficile* infection manifests as intestinal infections, namely pseudomembranous colitis. The occurrence of extra-intestinal disease is thought to be rare with a rate of 1.08% of 2034 isolates of *C. difficile* and an incidence of 4/100,000 admissions. *C. difficile* had been rarely associated with osteomyelitis. Here, we report the occurrence of *C. difficile* infection in a patient with sickle cell disease. The patient had multiple surgeries and a prolonged antimicrobial therapy to achieve a cure. The patient had *C. difficile* infection of native bone and of a prosthetic joint. The patient received prolonged therapy with amoxicillin–clavulanic acid and metronidazole and she remained free of *C. difficile* infection for three years off antibiotics.

Keywords: *Clostridium difficile*; Osteomyelitis; Sickle cell anemia

**Introduction:**

*Clostridium difficile* is usually associated with intestinal infections, namely pseudomembranous colitis. *Clostridium difficile* colitis is an important healthcare associated gastrointestinal infection. There is limited data on the prevalence or incidence of *C. difficile* infection in Saudi Arabia. In one study, the annual incidence rates of *C. difficile* infections were 1.2 and 0.9 per 1000 discharges, and 2.4 and 1.7 per 10,000 patient days in 2007 and 2008, respectively (Al-Tawfiq and Abed 2010). The occurrence of extra-intestinal disease is thought to be rare. In one study, 21 (1.08%) of 2034 isolates were from extra-intestinal sources (García-Lechuz et al. 2001) with an incidence of 4/100,000 admissions (García-Lechuz et al. 2001). Of those cases, five patients had either brain abscess, bacteremia, foot infections, or chronic osteomyelitis (García-Lechuz et al. 2001). Other studies estimated extra-intestinal CDI to represents 0.17-0.6% of all CDI (Mattila et al. 2013; Gupta et al. 2014). In a review article, 33
Extra-intestinal manifestation of *C. difficile* had been reported in the form of bacteremia (Feldman et al. 1995; García-Lechuz et al. 2001; Jacobs et al. 2001; Libby and Bearman 2009; Choi et al. 2013; Shah et al. 2017), abscesses (Durojaiye et al. 2011; Ulger Toprak et al. 2016; Roy et al. 2017), and rarely had also been associated with osteomyelitis (Riley and Karthigasu 1982; Towns et al. 1984; Incavo et al. 1988; Pron et al. 1995; Gaglani et al. 1996; García-Lechuz et al. 2001; Bachmeyer et al. 2008; Al-Najjar et al. 2013; Curtis and Lipp 2013; Ranganath and Midturi 2013). Two previous cases of *C. difficile* chronic osteomyelitis were reported among sickle cell patients (Gaglani et al. 1996; Bachmeyer et al. 2008). Here, we report a case of multifocal osteomyelitis caused by *C. difficile* and review the available literature in this regard.

**Case Presentation:**

The patient is a 28-year-old Saudi female with a history of sickle cell disease and was maintained on Hydroxyurea. She presented with pain in the right arm and right shoulder associated with fever of few days in duration. She did not have any other symptoms and initial temperature was 39.3°C. Laboratory data showed a hemoglobin of 7gm/dl; total white cell count of 6.2 and platelets count was normal. Erythrocyte sedimentation rate (ESR) was 111 mm/hr, and C-reactive protein (CRP) 15 mg/dl. Chest X-ray did not show any infiltrates. She was empirically started on ceftriaxone on admission. Urine and several blood cultures came back negative.

The right shoulder pain got worse and a Magnetic Resonance Imaging (MRI) showed destructive osseous changes involving proximal part of the humerus and a large effusion (Figure
1. Joint aspiration was carried out and synovial fluid analysis showed 200 White Blood Cell (WBC) (80% neutrophil), 4900 Red Blood Cell (RBC), friable and anaerobic culture grew *C. difficile*. She was started on intravenous metronidazole but she developed severe nausea and vomiting and could not tolerate metronidazole despite anti-emetics. She was switched to intravenous vancomycin. She also developed pancytopenia, so hydroxyurea was discontinued pending recovery of the bone marrow. Right shoulder incision and drainage and irrigation was done.

Three weeks after starting vancomycin, she developed severe pain in her right leg with tenderness over the shin, and fever recurred. MRI of the right lower extremity showed sickle cell changes and possible osteomyelitis involving the mid-shaft of the right tibia (Figure 2). Debridement of the right tibia was done with creation of a bone window. The cultures from the right tibia tissue grew *C. difficile*. Metronidazole was added to vancomycin plus anti-emetics. The patient tolerated metronidazole well this time. Two months on a combination of vancomycin and metronidazole, she developed an abscess in the proximal right arm (figure 3); the abscess was evacuated down to the bicepital tendon. There was no communication with the humerus; nevertheless, several drills were made into the humerus and there was no pus or debris. Culture from abscess material grew *C. difficile*.

A month later she developed a right tibial osteomyelitis confirmed by bone curettes cultures to be a recurrence of *C. difficile* infection. She was treated with intravenous (IV) metronidazole 500mg IV and amoxicillin-clavulanic acid 1gm orally twice a day (BID) for 8 weeks. She had multiple debridement and irrigation and subsequent tissue cultures were negative. She was referred for hyperbaric oxygen therapy and had 46 sessions. She required a total of six months of treatment with metronidazole and vancomycin. Although, there is no
specific minimum inhibitory concentration (MIC) cut down for antimicrobial sensitivity, the organism was tested with E-test and the diameter of inhibition is shown in table 1.

Fourth month after completing therapy, the patient went to an out-off Kingdom hospital where she had bilateral total hip replacement for aseptic necrosis. Two weeks after the surgery, she had post-operative course complication in the form of early right total hip prosthetic C. difficile infection. She initially had a debridement of the hip and two weeks later she had hardware removal and a two-stage surgery. Patient was maintained on oral amoxicillin-clavulanic acid and metronidazole. She subsequently returned to our hospital with continue dozing from the right hip. Patient was maintained on oral amoxicillin-clavulanic acid and metronidazole for twelve months. She was followed for three years off antibiotics and she had no evidence of recurrence of C. difficile infection.

Discussion:

We presented a case of Sickle cell disease who had multi-focal osteomyelitis with C. difficile and had a prolonged and protracted illness followed by a complete cure with no evidence of relapse. The pure growth on several occasions and from different sites is strong evidence for the responsibility of C. difficile in the infectious process of this case.

The occurrence of C. difficile osteomyelitis rarely reported (Riley and Karthigasu 1982; Towns et al. 1984; Incavo et al. 1988; Pron et al. 1995; Gaglani et al. 1996; García-Lechuz et al. 2001; Bachmeyer et al. 2008; Al-Najjar et al. 2013; Curtis and Lipp 2013; Ranganath and Midturi 2013) and two previous cases were reported among sickle cell patients (Gaglani et al. 1996; Bachmeyer et al. 2008). In a study of 17 cases of extra-intestinal C. difficile, only 1 (5.9%) had osteomyelitis (García-Lechuz et al. 2001). The development of skin and
bone infections may follow traumatic injury (Jacobs et al. 2001) or secondary to bacteremia. In the current case, the patient had multifocal osteomyelitis and this suggest bacteremia followed by seeding of abnormal bone that may had been traumatized in the course of sickle cell disease. Although, the current patient did not have a documented bacteremia, such occurrence in relation to *C. difficile* had been reported (Libby and Bearman 2009). In another case, vertebral osteomyelitis followed an episode of *C. difficile* diarrhea suggesting a dissemination (Al-Najjar et al. 2013). However, the current case did not have diarrheal illness to suggest this mechanism for the multifocal osteomyelitis. In a case report, a prosthetic devise infection secondary to *C. difficile* occurred two years after diarrheal diseases (Al-Najjar et al. 2013) and thus a remote diarrheal *C. difficile* infection (CDI could not be excluded in the current patient.

In this case, the patient had a chronic relapsing multi-focal osteomyelitis of the tibia, shoulder, arm and of prosthetic hip. The infection was eventually cured after prolonged therapy with oral *amoxicillin*-clavulanic acid and metronidazole for twelve months from the last infection. The exact duration of therapy for *C. difficile* osteomyelitis is not known (Gaglani et al. 1996; Bachmeyer et al. 2008; Al-Najjar et al. 2013).

In conclusion, *C. difficile* osteomyelitis remains rare and this patient had a prolonged and protracted course of infection involving initially native bones and later involved prosthetic hip joint. Such patients may need prolonged antimicrobial therapy coupled with surgical debridement.
References:


Riley TV, Karthigasu KT. Chronic osteomyelitis due to Clostridium difficile. Br Med J (Clin


Figure 1: MRI showing destructive osseous changes involving proximal part of the humerus and a large effusion
Figure 2: MRI of the right lower extremity showing sickle cell changes and possible osteomyelitis involving the mid-shaft of the right tibia
Figure 3: Severe right shoulder changes in keeping with chronic septic arthritis of the shoulder with osteomyelitis of the proximal right humerus with new multifocal proximal right arm fluid collections most likely abscesses.
Table 1: Results of E-test of susceptibility tests of different antibiotics against *Clostridium difficile* (Inhibition zone in millimeter)

<table>
<thead>
<tr>
<th>Antibiotic</th>
<th>Zone of Inhibition in millimeter</th>
</tr>
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<tbody>
<tr>
<td>Ciprofloxacin</td>
<td>0</td>
</tr>
<tr>
<td>Penicillin G</td>
<td>26</td>
</tr>
<tr>
<td>Nitrofurantoin</td>
<td>37</td>
</tr>
<tr>
<td>Cefuroxime</td>
<td>22</td>
</tr>
<tr>
<td>Oxacillin</td>
<td>0</td>
</tr>
<tr>
<td>Cefoxitin</td>
<td>0</td>
</tr>
<tr>
<td>Clindamycin</td>
<td>0</td>
</tr>
<tr>
<td>Vancomycin</td>
<td>35</td>
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