Case Report

Clostridium difficile Colitis Leading to Reactive Arthritis: A Rare Complication Associated With a Common Disease

Asghar Marwat, MD1,2, Hassan Mehmood, MD1,2, Ali Hussain, MD1,2, Muzammil Khan, MD1,2, Asad Ullah MD1,2, and Medha Joshi, MD1,2

Abstract

The relationship between reactive arthritis and enteric infections caused by Yersinia enterocolitica, Campylobacter jejuni, and Salmonella typhimurium is well documented. Clostridium difficile colitis is a less recognized cause of reactive arthritis. We present a case of a 58-year-old woman with Clostridium difficile colitis complicated by reactive arthritis. A 58-year-old woman with no significant past medical history presented to our hospital with complaints of nonbloody watery diarrhea, abdominal pain for the past 1 week, and right knee pain starting 1 day prior. The patient had recently used antibiotics for a respiratory tract infection. On examination, the patient had a swollen and erythematous right knee. While in the hospital the patient also developed a similarly painful and swollen left knee. The patient was found to be positive for C difficile toxin in stool. Synovial fluid analysis of both the knee joints revealed a sterile and inflammatory fluid, negative for crystals and showing no growth on gram stain. We diagnosed the patient with reactive arthritis secondary to C difficile colitis once all other causes of the bilateral knee joint symptoms were ruled out with appropriate laboratory and imaging studies. Treatment with oral vancomycin and an anti-inflammatory was initiated, and the patient had complete resolution of symptoms. This case illustrates the importance of recognizing C difficile colitis as a potential differential for reactive arthritis under the appropriate circumstances. The treatment of reactive arthritis is mainly supportive and treating the underlying cause, which happens to be C difficile in this case.

Keywords

Clostridium difficile infection, reactive arthritis

Introduction

Reactive arthritis typically manifests as an acute aseptic, inflammatory, asymmetric oligoarthritis commonly affecting the large joints of the lower extremities. Associated extra-articular findings including conjunctivitis, uveitis, enthesopathy, urethritis, balanitis, and keratoderma blennorrhagicum may also be observed. Reactive arthritis usually develops after an infection in a distant part of the body, and the causative organism is never isolated from the joint. The relationship between reactive arthritis and enteric infections caused by Yersinia enterocolitica, Campylobacter jejuni, and Salmonella typhimurium is well documented. Clostridium difficile colitis is a less recognized because of reactive arthritis. Only 50 cases have been reported since it was first described in 1976 by Rollins and Moeller. We present a case of a 58-year-old woman with C difficile colitis complicated by reactive arthritis.

Case Report

A 58-year-old woman with no significant past medical history came to our hospital with complaints of abdominal pain, diarrhea, and right knee pain. The patient had an acute onset of severe, nontraumatic right knee pain, with redness and swelling of the joint for the past day. She has nonbloody, watery diarrhea going on for the past 1 week with maximum episodes up to 10 in a day. She recently suffered from an infection. A 58-year-old woman with no significant past medical history presented to our hospital with complaints of nonbloody watery diarrhea, abdominal pain for the past 1 week, and right knee pain starting 1 day prior. The patient had recently used antibiotics for a respiratory tract infection. On examination, the patient had a swollen and erythematous right knee. While in the hospital the patient also developed a similarly painful and swollen left knee. The patient was found to be positive for C difficile toxin in stool. Synovial fluid analysis of both the knee joints revealed a sterile and inflammatory fluid, negative for crystals and showing no growth on gram stain. We diagnosed the patient with reactive arthritis secondary to C difficile colitis once all other causes of the bilateral knee joint symptoms were ruled out with appropriate laboratory and imaging studies. Treatment with oral vancomycin and an anti-inflammatory was initiated, and the patient had complete resolution of symptoms. This case illustrates the importance of recognizing C difficile colitis as a potential differential for reactive arthritis under the appropriate circumstances. The treatment of reactive arthritis is mainly supportive and treating the underlying cause, which happens to be C difficile in this case.

1 Temple University, Philadelphia, PA, USA
2 Conemaugh Memorial Medical Center, Johnstown, PA, USA

Received January 2, 2018. Revised February 23, 2018. Accepted March 4, 2018.

Corresponding Author:
Asghar Marwat, MD, Department of Internal Medicine, Conemaugh Memorial Medical Center, 1086 Franklin Street, Johnstown, PA 15905, USA.
Email: asghardjk@gmail.com

Creative Commons CC BY: This article is distributed under the terms of the Creative Commons Attribution 4.0 License (http://creativecommons.org/licenses/by/4.0/) which permits any use, reproduction and distribution of the work without further permission provided the original work is attributed as specified on the SAGE and Open Access pages (https://us.sagepub.com/en-us/nam/open-access-at-sage).
Discussion

**Clostridium difficile** carries a significant disease burden in the United States. According to the Center for Diseases Control and Prevention, *C difficile* was responsible for almost half a million infections, resulting in roughly 29,000 deaths in 2011.1 *C difficile* infection most commonly leads to pseudomembranous colitis, which presents itself as fever and diarrhea starting from 4 to 9 days after starting antibiotic treatment.2 Extracolonic manifestations of *C difficile* including bacteremia, osteomyelitis, visceral abscess, empyema, small bowel disease, and reactive arthritis are less frequent and rarely reported.3 Our case adds to the growing body of literature on this subject. The hypothesized pathogenesis of reactive arthritis following an enteric infection with *C difficile* is postulated to be an autoimmune response to bacterial antigens in joints and other tissues that gain access into the bloodstream via the intestinal mucosa.4 Our patient fulfilled the criteria for the diagnosis of *C difficile* reactive arthritis as established by Putterman and Rubinow in 19935: a sterile inflammatory arthritis with preceding diarrhea and prior antibiotic exposure6; stool test positive for *C difficile* toxin7; and no alternative diagnosis for arthritis or diarrhea.8 With the increased use of antibiotics in recent times and the associated increased incidence of *C difficile* colitis, many rare manifestations of this disease process that were previously unrecognized are coming to the forefront. It would be prudent for physicians to consider reactive arthritis secondary to *C difficile* colitis as a differential diagnosis in a patient with otherwise unexplained acute inflammatory arthritis in the right clinical setting of recent antibiotic use and diarrhea. Early diagnosis and appropriate treatment can improve patient outcomes and prevent unnecessary diagnostic procedures. Reactive arthritis secondary to *C difficile* colitis is managed conservatively with treatment primarily focused on eradicating the *C difficile* infection and has an excellent long-term prognosis.

Conclusion

Given the expected rise in the incidence of *C difficile* infection, both internist and rheumatologist should include this pathogen in the differential diagnosis of the enteric organism responsible for reactive arthritis. We suspect *C difficile* reactive arthritis may be underrecognized and recommend raising awareness in the health care profession to test *C difficile* toxin in undifferentiated arthritis patients.

Declaration of Conflicting Interests

The author(s) declared no potential conflicts of interest with respect to the research, authorship, and/or publication of this article.

Funding

The author(s) received no financial support for the research, authorship, and/or publication of this article.

Ethics Approval

Ethical approval to report this case was obtained from the Institutional Review Board Approval Committee, Conemaugh Memorial Medical Center.

Informed Consent

Verbal informed consent was obtained from the patient(s) for their anonymized information to be published in this article.

ORCID iDs

Hassan Mehmood https://orcid.org/0000-0002-6754-4913
Asad Ullah https://orcid.org/0000-0001-7400-8724

References

1. Flores D, Marquez J, Garza M, Espinoza LR. Reactive arthritis: newer developments. *Rheum Dis Clin North Am.* 2003;29: 37-59.
2. Hill Gaston JS, Lillicrap MS. Arthritis associated with enteric infection. *Best Pract Res Clin Rheumatol*. 2003;17:219-239.
3. Ahvonen P, Sievers K, Aho K. Arthritis associated with *Yersinia enterocolitica* infection. *Acta Rheumatol Scand*. 1969;15:232.
4. Keating RM, Vyas AS. Reactive arthritis following *Clostridium difficile* colitis. *West J Med*. 1995;162:61-63.
5. Legendre P, Lalande V, Eckert C, et al. *Clostridium difficile* associated reactive arthritis: case report and literature review. *Anaerobe*. 2016;38:76-80.
6. Rollins DE, Moeller D. Acute migratory polyarthritis associated with antibiotic-induced pseudomembranous colitis. *Am J Gastroenterol*. 1976;65:353-356.
7. Lessa FC, Mu Y, Bamberg WM, et al. Burden of *Clostridium difficile* infection in the United States. *N Engl J Med*. 2015;372:825-834.
8. Jacobs A, Bamard K, Fishel R, Gradon JD. Extracolonic manifestations of *Clostridium difficile* infections. Presentation of 2 cases and review of the literature. *Medicine (Baltimore)*. 2001;80:88-101.
9. Putterman C, Rubinow A. Reactive arthritis associated with *Clostridium difficile* pseudomembranous colitis. *Semin Arthritis Rheum*. 1993;22:420-426.
10. McClusky J, Riley TV, Owen ET, Langlands DR. Reactive arthritis associated with *Clostridium difficile*. *Aust N Z J Med*. 1982;12:535-537.
11. Atkinson MH, Mcleod BD. Reactive arthritis with *Clostridium difficile* enteritis. *J Rheumatol*. 1988;15:520-522.