Cureus

Received 02/01/2023 Review began 02/11/2023 Review ended 02/22/2023 Published 02/28/2023

© Copyright 2023

Stolzenberg et al. This is an open access article distributed under the terms of the Creative Commons Attribution License CC-BY 4.0., which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Rheumatoid Arthritis-Linked Artificial Joint Infections Leading to Amputations

Laurence Stolzenberg 1 , Austin Huang 2 , Mohammad Usman 3 , Alexis Koch 4 , John Stevenson 5 , Colby Kihara 6 , Jason Seale 7

1. Orthopedic Surgery, Alabama College of Osteopathic Medicine, Dothan, USA 2. Neurology, Alabama College of Osteopathic Medicine, Dothan, USA 3. Anesthesiology, Alabama College of Osteopathic Medicine, Dothan, USA 4. Internal Medicine, Alabama College of Osteopathic Medicine, Dothan, USA 5. Gastroenterology, Alabama College of Osteopathic Medicine, Dothan, USA 7. General Surgery, Decatur Morgan Hospital, Decatur, USA

Corresponding author: Laurence Stolzenberg, stolzenbergl@acom.edu

Abstract

Rheumatoid arthritis (RA) is a common autoimmune condition that can rarely cause more serious complications, such as permanent joint damage or infection, and may pose a significant additional risk during certain routine procedures. One major consequence of RA is that it can lead to serious and permanent joint damage requiring arthroplasty. Additionally, RA is a known cause of infection, with orthopedic prosthetic joint infections (PJIs) being documented. We explore one such serious case of a patient with long-term RA and a left knee joint replacement who presented to the emergency room with a serious PJI. History revealed that he repeatedly was affected by infections and had an extensive and severe clinical course, including nine revision surgeries. After a physical examination, imaging was performed, which further supported the diagnosis of joint infection. Considering the extensive number of attempts to salvage the joint, clinicians decided an above-knee amputation was necessary. This case highlights the fact that RA both increases the need for orthopedic arthroplasties and the risk of complications from these procedures, complicating clinical decision-making for physicians. Additionally, this patient had other underlying medical conditions and social habits that may have contributed to his severe clinical course, and we hope to explore these, discuss possible methods of modifying them, and assist clinicians in not only better treating similar patients but also emphasizing the importance of further developing standardized predictive algorithms and scoring tools.

Categories: Infectious Disease, Orthopedics, Rheumatology

Keywords: sulfasalazine, amputation, joint infections, immune suppressing medications, orthopedics, orthopedic surgery, prosthetic joint infection, rheumatoid arthritis

Introduction

Rheumatoid arthritis (RA) is an autoimmune disease that causes chronic, systemic inflammation, which can lead to inflammatory arthritis as well as extra-articular involvement [1]. The etiology of RA is not fully understood; however, various alleles have been identified that have a strong association with developing the condition [2-4]. RA primarily affects synovial joints, and these affected joints are often in a symmetric distribution. Progression of joint involvement often begins in small peripheral joints and then proceeds to involve large joints, such as knees [1]. RA patients will often present with swelling, arthralgias, erythema, decreased range of motion, and stiffness that is worse in the morning upon waking up and classically improved by evening [5,6]. Appropriate management of RA is essential as those affected can have a significantly decreased quality of life and increased risk for infections, cardiovascular disease, cancer, osteoporosis, respiratory disease, and mortality than the general population [5]. There is no known cure for RA: however, management is aimed at suppressing the activity and severity of the disease. Treatment with short-term nonsteroidal anti-inflammatory drugs (NSAIDs) and corticosteroids has been shown to decrease pain and stiffness in those affected by RA [7]. It is important to note, however, that these analgesic and antiinflammatory medications do not improve or slow down disease progression, but instead improve the overall quality of life [6]. Other various long-term disease-modifying antirheumatic drugs (DMARDs) are other options that, as the name implies, modify the disease course.

Infections are of major concern for those afflicted with RA as they contribute to increased overall mortality [8-12]. Associations between RA and infections have been well documented in the literature, including various cohort studies that have found an increased risk of serious infections in RA patients [8,13]. However, there is no one specific etiology responsible for the pathogenesis of these infections, rather it is likely to occur due to a multifactorial set of issues. Infections can also be the result of the disease itself due to immobility and immunological dysfunction [14]. The hypothesis behind the immunological dysfunction resulting from RA is that premature aging of the immune system may result in a weakened ability to fight off pathogens [15].

Other major contributors to immunological dysfunction in patients with RA are chronic comorbid

How to cite this article

Stolzenberg L, Huang A, Usman M, et al. (February 28, 2023) Rheumatoid Arthritis-Linked Artificial Joint Infections Leading to Amputations. Cureus 15(2): e35622. DOI 10.7759/cureus.35622

conditions, such as diabetes mellitus (DM) and kidney disease, as well as social factors such as smoking [11,15]. Smoking has been shown to have a significant impact on the immune system, which promotes infection. Inhalation of the chemicals in cigarette smoke results in the formation of oxidative moieties that activate immune cells such as macrophages to secrete proinflammatory cytokines [16]. Additionally, cigarette smoke can affect immune system signaling pathways and modify transcription, which can result in immunosuppression and anti-inflammatory effects [16]. The result of this spectrum of possible effects from cigarette smoke determines the overall modification of the immune system [16].

Finally, the increased risk of infection may be due to treatments such as joint surgery or pharmacological treatments such as DMARDs, immunosuppressants, and steroids [14,15,17]. Joint surgery in RA has been shown to increase the development of septic arthritis [18,19]. Glucocorticoids are the mainstay of RA treatment and have been well researched and have been linked to increased risk of major infections in several rheumatic diseases [20].

As RA progresses, joints can often become severely deformed, thus having a significant impact on the patient's quality of life. One option available to patients is joint replacement. Joint replacement is a procedure to improve the patient's quality of life with overall benefits such as reducing pain and restoring independence [21]. Rates of joint surgeries, including joint arthroplasties, have been rising and are expected to continue to increase [21,22]. Whether or not a prosthetic joint is involved, joint infection is an inherent complication of any joint surgery. Specifically, with prosthetic joint infections (PJIs), comorbid conditions such as obesity and DM can compound this risk for infections [23-31]. Of note, RA has been found in multiple studies to be associated with an increased risk of PJIs [23,27,28,32-37]. Other risk factors have also been documented such as male gender, smoking, arthroplasty revision surgery, and antecedent septic arthritis of the specific joint [22,26,27,30-32,36-44].

The pathogenesis of postoperative joint infections is thought to be due to the introduction of microorganisms during surgery. However, the contiguous and hematogenous spread of microorganisms can also result in joint seeding. The most common infectious organisms are gram-positive cocci, most of which are Staphylococcus aureus and coagulase-negative staphylococci [30,32,45-56]. Patients with joint infections often present with pain, joint swelling or effusion, erythema, warmth, fever, drainage, and/or the presence of a sinus tract communicating with the arthroplasty [45,57-59]. The diagnosis is made with consideration of clinical, laboratory, and radiological findings with intraoperative inspection and tissue evaluation [21]. Treatment is often a combination of both antibiotics and surgical intervention [21].

Case Presentation

This patient was a 72-year-old male with multiple underlying medical and social conditions, including RA, tobacco abuse, hypertension, and a history of opioid abuse who presented to the emergency department with a complaint of left lower extremity (LLE) swelling, redness, and purulent drainage for the last four weeks. He had been taking oral doxycycline as prescribed by his family physician for five days, but his symptoms had continued to worsen. The patient was admitted to integumentary abnormalities, swelling, and an ulcerated lesion on his LLE but denied any other symptoms. Family history was significant for RA in both parents, and the patient endorsed an allergy to intravenous (IV) contrast. A review of home medications was notable for the previously mentioned doxycycline, sulfasalazine 1,500 mg twice daily, ibuprofen 800 mg thrice daily, and hydroxychloroquine 200 mg twice daily. Physical examination revealed unremarkable respiratory, cardiovascular, and gastrointestinal examinations. When attention was turned to the LLE, swelling, fluctuance, warmth, and tenderness over the left knee were noted. The left dorsalis pedis pulse was noted as diminished, being graded as 1+. The patient was noted to have a normal gait. The patient was fully alert, oriented, and in no acute distress, although he was described as ill-appearing. Vitals were within normal ranges with a temperature of 97.5 °F, a pulse of 85 beats per minute, a body mass index (BMI) of 20.5 kg/m², a respiratory rate of 20 breaths per minute, and a blood pressure of 154/86 mmHg.

Initial laboratory investigations included a complete metabolic panel (CMP), which was unremarkable, liver function tests within normal limits, and renal function tests (RFTs), which were equally unremarkable with a blood urea nitrogen (BUN) of 21 mg/dL and serum creatinine (sCr) of 0.8 mg/dL. Glucose was 91 mg/dL, and serum calcium was at the upper end of normal, revealing a value of 10.2 mg/dL. Complete blood count (CBC) was remarkable for a significantly elevated white cell count of 28,110 cells/µL, a low hemoglobin level of 12.6 mg/dL, and a low hematocrit of 37%. A urinalysis was also completed and found to be normal as well. Imaging performed included computed tomography (CT) of the lower extremities, which showed a left knee arthroplasty with a long stem prosthesis. These images also revealed a collection of fluid with internal droplets of gas anterior to the knee measuring 8.1 cm × 4.5 cm × 12.3 cm, suggesting an abscess (Figures *1-2*). The patient was empirically placed on vancomycin, piperacillin-tazobactam, and morphine.



FIGURE 1: CT of the left lower extremity, transverse cut, showing gas bubbles (green arrow) in the fluid collection, suggestive of an abscess. CT, computed tomography



FIGURE 2: CT of the left lower extremity, sagittal cut, showing gas bubbles (green arrow) in the fluid collection, suggestive of an abscess.

CT, computed tomography

The patient was then admitted to the hospital where surgery and orthopedics were consulted for further management. The hospitalist obtained further history, which revealed an extensive number of surgeries on the LLE, beginning with a total left knee arthroplasty in 2014. He said it got infected and replaced in 2015 with a Teflon implant. In 2017, there was a second infection, which was treated with two separate instances of antibiotic spacer placement and another replacement of the prosthetic joint. He endorsed a further infection in 2018 where surgery was performed to cut nonviable muscle and place a graft. In total, the patient endorsed having had nine separate surgical procedures on his LLE, with the most recent one in 2018. The last surgeon installed an antibiotic spacer, which was considered a final attempt at saving the limb before considering above-knee amputation. Repeat physical examination revealed the possibility to express grossly purulent drainage from a small ulcer in the lateral portion of the left knee as well as a large soft tissue fluid collection anterior to the joint. Vancomycin was continued, but piperacillin-tazobactam was discontinued and replaced with cefepime based on hospital protocol.

The general and orthopedic surgeons decided an urgent above-knee amputation was necessary, and it was scheduled for the following day. During surgery, a large pocket of pus was encountered anterior to the joint, as delineated by imaging, while separating the soft tissue and collected for culture. The bone had a *motheaten* appearance, and debridement of the medullary cavity of the femur was required with a curette. The surgery was performed by both the general and orthopedic surgeons with no complications other than an estimated blood loss of 50 mL.

The patient recovered from surgery relatively well, although treating pain while minimizing opioid use was challenging. The patient was also affected by severe anxiety and anorexia, which were treated successfully with dronabinol. Multiple cultures from the fluid collected during surgery were inconclusive, but empiric antibiotic therapy was continued. On postsurgery day 9, the patient was discharged home with outpatient follow-up with infectious disease, orthopedics, general surgery, and home health. He was set up with a mechanical wheelchair and appropriate pain measures with minimal opioids, considering his history of opioid dependence and personal preference to minimize opioids and oral antibiotics. He is expected to do well as of the time of this paper.

Discussion

This patient underwent multiple extensive surgeries in an attempt to resolve the recurrent severe infection in his prosthetic knee, and as such, it is reasonable to assume that this patient was affected greatly. Not only did he have to deal with significant levels of pain, which might have contributed to his opioid abuse and dependence, but his condition significantly affected his quality of life and activities of daily living. As such, the considerable morbidity experienced by this patient implores us to further explore the underlying causes of his recurrent infections and discover if any other treatments could have been considered to improve his condition. It is unfortunately quite difficult to pinpoint the exact origin of his multiple subsequent infections, but we will attempt to go over the possible etiologies one by one. The first condition to consider is his RA. While it is difficult to isolate the specific level of risk due to that disease process alone, multiple studies have cited this condition as a significant risk factor [23,27,28,32-37]. Aside from RA, it is also important to explore other social habits, risk factors, and medications that likely contributed to the development and progression of his infections.

Our patient was a heavy smoker, with a history of smoking half a pack a day for 63 years. Literature has established a link between smoking and generally increased risk of infection and thus likely played a role in his clinical course [15,27,60,61]. Although it is nearly certain that many physicians and healthcare professionals suggested and counseled them to quit smoking, it would perhaps be worth considering if anyone pointed out the well-supported link between these infections and tobacco use to the patient [15,27,60,61]. During his stay in the hospital, he had been able to temporarily refrain from smoking with help from a nicotine patch, so it is reasonable to suppose further pharmacological therapy could have at least assisted him in this difficult endeavor.

Considering our patient's severe clinical course, we need to review comorbid conditions that may have contributed to increased infection risk. This patient denied any history of DM, which was supported by his blood glucose levels being consistently within normal limits. His estimated glomerular filtration rate (eGFR) also precluded any form of chronic kidney disease. A commonly cited risk factor for the development of PJIs is obesity. In this specific case, our patient was on the other end of the spectrum, with a BMI of 20.5 kg/m². One study found that a low BMI (<25 kg/m²) was associated with an increased risk of PJIs, possibly due to a low nutritional reserve and immunosuppression [62]. Interestingly, treating malnutrition as a possible intervention for PJI prevention has been a recent area of research interest, with multiple pharmaceutical interventions showing promise in treating this risk factor [61]. In this case, dronabinol successfully improved this patient's anorexia. Dronabinol use is an area of research that has received increasing attention, and there have been some promising data showing that its use may benefit patients with chronic malnutrition [63,64]. However, more research is certainly needed to conclude whether this patient or similar patients would benefit from regular outpatient dronabinol use.

Nonmodifiable risk factors, such as male sex and advanced age likely also played a role in the pathogenesis of this patient's recurrent infections. It is widely documented in the existing literature that advanced age can increase the risk of specific kinds of infections, and indeed skin and soft tissue infection rates are elevated in the elderly. Further complicating the management of this patient is the fact that nosocomial infections are a major risk factor in the elderly. In fact, not only are elderly patients more likely to spend lengthy amounts of time in a hospital setting, putting them at risk, but they are also more likely to become infected each day they stay in the hospital [65]. Another nonmodifiable intrinsic risk factor in this patient is that he is of the male sex, which has specifically been shown to increase the risk of PJIs [26,37-41].

An additional risk factor for this patient is his multiple revision surgeries. Research has shown that the incidence of PJIs is higher in patients that have undergone an arthroplasty revision surgery when compared to primary implantation [30-32,36,42-44,66]. This is likely due to prolonged operating time. Our patient had undergone extensive repair surgeries following multiple joint infections, with each surgery increasing his risk for a future infection. There are possibly other reasons for the increased risk associated with revision surgeries that could be covered in further work.

Medications are important to review as many medications are known to cause immunosuppression or increase the risk of infection, and they can be modified to mitigate risk. The patient took ibuprofen regularly. Regular ibuprofen use does not seem linked to any increase in the risk of infection as a review of available literature demonstrates no strong or significant association between NSAIDs and PJIs. Multiple publications support the benefits of NSAIDs during joint infections, prosthetic or not [67,68]. Next, this patient took

sulfasalazine daily. While an increased risk of infection is noted in the documentation associated with this medication [69], no published case reports specifically linking this medication to infections were found when using multiple keywords. Additionally, the information available on this medication from the American Academy of Rheumatologists has no mention of infections [70]. Finally, hydroxychloroquine revealed a similar pattern of contraindication and reported infections in existing literature as sulfasalazine [71]. Existing literature specifically mentions these are the safest agents for use in the perioperative management of an RA patient. A review of the literature shows many other treatments for RA are associated with higher risks of infection [72], although these were not included in his treatment plan. Based on current guidelines and an in-depth review of medication, we can state that the management of this patient's pharmaceutical regimen followed all the latest guidelines and literature, and it is doubtful that using the other available agents would have prevented his complications.

When considering this patient's severe and prolonged clinical course with PJIs, discussed risk factors, and active medications, it is important to consider if an initial arthroplasty should have been performed at all. While clinical acumen is important in these decisions, the numerous factors described above and the overall complex clinical picture show the need for a standardized method to aid clinical decision-making. Recently, there has been a push to develop a tool to aid in the assessment of the risk of developing PJIs. Of note, the Mayo PJI risk score (Mayo Clinic, Scottsdale, AZ, USA) is one such tool; however, it is not yet fully validated [62]. The Mayo PJI risk score accounts for multiple risk factors such as BMI, prior joint surgery, immunosuppression, American Society of Anesthesiology (ASA) score, and procedure duration to assist in risk stratification and consider additional preventative measures. We believe further work on this scoring tool would be of great assistance to patients and may help prevent severe clinical courses as seen in this patient.

Conclusions

We hope this case will help future clinicians in managing similar patients affected by both autoimmune disorders such as RA and repeated infections to improve clinical outcomes for those undergoing arthroplasties. As the rate of procedures such as arthroplasties rises in the elderly population, so does the risk of complications, notably infections. As such, clinicians should consider all underlying conditions that could cause significant complications when planning such procedures and during the postoperative care environment. Clinicians should potentially consider interventions such as further or more potent prophylactic antibiotics, more aggressive treatment of the underlying RA, or more radical interventions such as above-knee amputations and the use of a prosthetic limb in the future if presented with a similar patient affected by repeated infections. It is also important to analyze the risk and benefits before planning potentially complicated procedures. In addition, if it is decided that further arthroplasties are indicated, potentially using a different surgical technique or different replacement parts intraoperatively should be considered. We also hope this case will assist other clinicians working on predictive models and algorithms to help score the likelihood of PJIs before arthroplasty.

Additional Information

Disclosures

Human subjects: Consent was obtained or waived by all participants in this study. Conflicts of interest: In compliance with the ICMJE uniform disclosure form, all authors declare the following: Payment/services info: All authors have declared that no financial support was received from any organization for the submitted work. Financial relationships: All authors have declared that they have no financial relationships at present or within the previous three years with any organizations that might have an interest in the submitted work. Other relationships: All authors have declared that there are no other relationships or activities that could appear to have influenced the submitted work.

References

- 1. Chauhan K, Jandu JS, Brent LH, Al-Dhahir MA: Rheumatoid Arthritis. StatPearls, Treasure Island, FL; 2022.
- Wu H, Liao W, Li Q, et al.: Pathogenic role of tissue-resident memory T cells in autoimmune diseases . Autoimmun Rev. 2018, 17:906-11. 10.1016/j.autrev.2018.03.014
- du Teil Espina M, Gabarrini G, Harmsen HJ, Westra J, van Winkelhoff AJ, van Dijl JM: Talk to your gut: the oral-gut microbiome axis and its immunomodulatory role in the etiology of rheumatoid arthritis. FEMS Microbiol Rev. 2019, 43:1-18. 10.1093/femsre/fuy035
- Kłodziński Ł, Wisłowska M: Comorbidities in rheumatic arthritis. Reumatologia. 2018, 56:228-33. 10.5114/reum.2018.77974
- 5. Sparks JA: Rheumatoid arthritis. Ann Intern Med. 2019, 170:ITC1-16. 10.7326/AITC201901010
- Guo Q, Wang Y, Xu D, Nossent J, Pavlos NJ, Xu J: Rheumatoid arthritis: pathological mechanisms and modern pharmacologic therapies. Bone Res. 2018, 6:15. 10.1038/s41413-018-0016-9
- Fraenkel L, Bathon JM, England BR, et al.: 2021 American college of rheumatology guideline for the treatment of rheumatoid arthritis. Arthritis Care Res (Hoboken). 2021, 73:924-39. 10.1002/acr.24596
- Doran MF, Crowson CS, Pond GR, O'Fallon WM, Gabriel SE: Frequency of infection in patients with rheumatoid arthritis compared with controls: a population-based study. Arthritis Rheum. 2002, 46:2287-93. 10.1002/art.10524
- 9. Cobb S, Anderson F, Bauer W: Length of life and cause of death in rheumatoid arthritis . N Engl J Med. 1953,

249:553-6. 10.1056/NEJM195310012491402

- Björnådal L, Baecklund E, Yin L, Granath F, Klareskog L, Ekbom A: Decreasing mortality in patients with rheumatoid arthritis: results from a large population based cohort in Sweden, 1964-95. J Rheumatol. 2002, 29:906-12.
- Wolfe F, Mitchell DM, Sibley JT, et al.: The mortality of rheumatoid arthritis. Arthritis Rheum. 1994, 37:481-94. 10.1002/art.1780370408
- 12. Prior P, Symmons DP, Scott DL, Brown R, Hawkins CF: Cause of death in rheumatoid arthritis. Br J Rheumatol. 1984, 23:92-9. 10.1093/rheumatology/23.2.92
- Widdifield J, Bernatsky S, Paterson JM, et al.: Serious infections in a population-based cohort of 86,039 seniors with rheumatoid arthritis. Arthritis Care Res (Hoboken). 2013, 65:353-61. 10.1002/acr.21812
- 14. Atzeni F, Masala IF, di Franco M, Sarzi-Puttini P: Infections in rheumatoid arthritis. Curr Opin Rheumatol. 2017, 29:323-30. 10.1097/BOR.00000000000389
- 15. Listing J, Gerhold K, Zink A: The risk of infections associated with rheumatoid arthritis, with its comorbidity and treatment. Rheumatology (Oxford). 2013, 52:53-61. 10.1093/rheumatology/kes305
- Stämpfli MR, Anderson GP: How cigarette smoke skews immune responses to promote infection, lung disease and cancer. Nat Rev Immunol. 2009, 9:377-84. 10.1038/nri2530
- Pérez-Sola MJ, Torre-Cisneros J, Pérez-Zafrilla B, Carmona L, Descalzo MA, Gómez-Reino JJ: Infections in patients treated with tumor necrosis factor antagonists: incidence, etiology and mortality in the BIOBADASER registry. Med Clin (Barc). 2011, 137:533-40. 10.1016/j.medcli.2010.11.032
- Goldenberg DL: Infectious arthritis complicating rheumatoid arthritis and other chronic rheumatic disorders. Arthritis Rheum. 1989, 32:496-502. 10.1002/anr.1780320422
- Rimoin DL, Wennberg JE: Acute septic arthritis complicating chronic rheumatoid arthritis . JAMA. 1966, 196:617-21. 10.1001/jama.1966.03100200057018
- Dixon WG, Suissa S, Hudson M: The association between systemic glucocorticoid therapy and the risk of infection in patients with rheumatoid arthritis: systematic review and meta-analyses. Arthritis Res Ther. 2011, 13:R139. 10.1186/ar3453
- 21. Tande AJ, Patel R: Prosthetic joint infection. Clin Microbiol Rev. 2014, 27:302-45. 10.1128/CMR.00111-13
- Singh JA, Yu S, Chen L, Cleveland JD: Rates of total joint replacement in the United States: Future projections to 2020-2040 using the National Inpatient Sample. J Rheumatol. 2019, 46:1134-40. 10.3899/irheum.170990
- Pulido L, Ghanem E, Joshi A, Purtill JJ, Parvizi J: Periprosthetic joint infection: the incidence, timing, and predisposing factors. Clin Orthop Relat Res. 2008, 466:1710-5. 10.1007/s11999-008-0209-4
- Namba RS, Inacio MC, Paxton EW: Risk factors associated with surgical site infection in 30,491 primary total hip replacements. J Bone Joint Surg Br. 2012, 94:1330-8. 10.1302/0301-620X.94B10.29184
- Malinzak RA, Ritter MA, Berend ME, Meding JB, Olberding EM, Davis KE: Morbidly obese, diabetic, younger, and unilateral joint arthroplasty patients have elevated total joint arthroplasty infection rates. J Arthroplasty. 2009, 24:84-8. 10.1016/j.arth.2009.05.016
- Dowsey MM, Choong PF: Obese diabetic patients are at substantial risk for deep infection after primary TKA. Clin Orthop Relat Res. 2009, 467:1577-81. 10.1007/s11999-008-0551-6
- 27. Peersman G, Laskin R, Davis J, Peterson M: Infection in total knee replacement: a retrospective review of 6489 total knee replacements. Clin Orthop Relat Res. 2001, 15-23.
- Peel TN, Dowsey MM, Daffy JR, Stanley PA, Choong PF, Buising KL: Risk factors for prosthetic hip and knee infections according to arthroplasty site. J Hosp Infect. 2011, 79:129-33. 10.1016/j.jhin.2011.06.001
- Dowsey MM, Choong PF: Obesity is a major risk factor for prosthetic infection after primary hip arthroplasty. Clin Orthop Relat Res. 2008, 466:153-8. 10.1007/s11999-007-0016-3
- Berbari EF, Osmon DR, Carr A, et al.: Dental procedures as risk factors for prosthetic hip or knee infection: a hospital-based prospective case-control study. Clin Infect Dis. 2010, 50:8-16. 10.1086/648676
- Uçkay I, Lübbeke A, Emonet S, et al.: Low incidence of haematogenous seeding to total hip and knee prostheses in patients with remote infections. J Infect. 2009, 59:337-45. 10.1016/j.jinf.2009.08.015
- 32. Berbari EF, Hanssen AD, Duffy MC, Steckelberg JM, Ilstrup DM, Harmsen WS, Osmon DR: Risk factors for prosthetic joint infection: case-control study. Clin Infect Dis. 1998, 27:1247-54. 10.1086/514991
- Cipriano CA, Brown NM, Michael AM, Moric M, Sporer SM, Della Valle CJ: Serum and synovial fluid analysis for diagnosing chronic periprosthetic infection in patients with inflammatory arthritis. J Bone Joint Surg Am. 2012, 94:594-600. 10.2106/JBJS.J.01318
- Bongartz T, Halligan CS, Osmon DR, et al.: Incidence and risk factors of prosthetic joint infection after total hip or knee replacement in patients with rheumatoid arthritis. Arthritis Rheum. 2008, 59:1713-20. 10.1002/art.24060
- Ravi B, Escott B, Shah PS, et al.: A systematic review and meta-analysis comparing complications following total joint arthroplasty for rheumatoid arthritis versus for osteoarthritis. Arthritis Rheum. 2012, 64:3839-49. 10.1002/art.37690
- Poss R, Thornhill TS, Ewald FC, Thomas WH, Batte NJ, Sledge CB: Factors influencing the incidence and outcome of infection following total joint arthroplasty. Clin Orthop Relat Res. 1984, 117-26.
- Jämsen E, Huhtala H, Puolakka T, Moilanen T: Risk factors for infection after knee arthroplasty. A registerbased analysis of 43,149 cases. J Bone Joint Surg Am. 2009, 91:38-47. 10.2106/JBJS.G.01686
- Dale H, Fenstad AM, Hallan G, et al.: Increasing risk of prosthetic joint infection after total hip arthroplasty. Acta Orthop. 2012, 83:449-58. 10.3109/17453674.2012.733918
- Kurtz SM, Ong KL, Lau E, Bozic KJ, Berry D, Parvizi J: Prosthetic joint infection risk after TKA in the Medicare population. Clin Orthop Relat Res. 2010, 468:52-6. 10.1007/s11999-009-1013-5
- 40. Ong KL, Kurtz SM, Lau E, Bozic KJ, Berry DJ, Parvizi J: Prosthetic joint infection risk after total hip arthroplasty in the Medicare population. J Arthroplasty. 2009, 24:105-9. 10.1016/j.arth.2009.04.027
- Hinarejos P, Guirro P, Leal J, et al.: The use of erythromycin and colistin-loaded cement in total knee arthroplasty does not reduce the incidence of infection: a prospective randomized study in 3000 knees. J Bone Joint Surg Am. 2013, 95:769-74. 10.2106/JBJS.L.00901
- 42. Cazanave C, Greenwood-Quaintance KE, Hanssen AD, et al.: Rapid molecular microbiologic diagnosis of

prosthetic joint infection. J Clin Microbiol. 2013, 51:2280-7. 10.1128/JCM.00335-13

- 43. Fitzgerald RH Jr, Nolan DR, Ilstrup DM, Van Scoy RE, Washington JA 2nd, Coventry MB: Deep wound sepsis following total hip arthroplasty. J Bone Joint Surg Am. 1977, 59:847-55.
- 44. Bozic KJ, Katz P, Cisternas M, Ono L, Ries MD, Showstack J: Hospital resource utilization for primary and revision total hip arthroplasty. J Bone Joint Surg Am. 2005, 87:570-6. 10.2106/JBJS.D.02121
- Peel TN, Cheng AC, Buising KL, Choong PF: Microbiological aetiology, epidemiology, and clinical profile of prosthetic joint infections: are current antibiotic prophylaxis guidelines effective?. Antimicrob Agents Chemother. 2012, 56:2386-91. 10.1128/AAC.06246-11
- Lee J, Kang CI, Lee JH, et al.: Risk factors for treatment failure in patients with prosthetic joint infections. J Hosp Infect. 2010, 75:273-6.
- 47. Raut VV, Siney PD, Wroblewski BM: One-stage revision of total hip arthroplasty for deep infection. Longterm followup. Clin Orthop Relat Res. 1995, 202-7.
- Schäfer P, Fink B, Sandow D, Margull A, Berger I, Frommelt L: Prolonged bacterial culture to identify late periprosthetic joint infection: a promising strategy. Clin Infect Dis. 2008, 47:1403-9. 10.1086/592973
- Shukla SK, Ward JP, Jacofsky MC, Sporer SM, Paprosky WG, Della Valle CJ: Perioperative testing for persistent sepsis following resection arthroplasty of the hip for periprosthetic infection. J Arthroplasty. 2010, 25:87-91. 10.1016/j.arth.2010.05.006
- Biring GS, Kostamo T, Garbuz DS, Masri BA, Duncan CP: Two-stage revision arthroplasty of the hip for infection using an interim articulated Prostalac hip spacer: a 10- to 15-year follow-up study. J Bone Joint Surg Br. 2009, 91:1431-7. 10.1302/0301-620X.91B11.22026
- Bengtson S, Knutson K: The infected knee arthroplasty. A 6-year follow-up of 357 cases . Acta Orthop Scand. 1991, 62:301-11. 10.3109/17453679108994458
- 52. Kim YH, Choi Y, Kim JS: Treatment based on the type of infected TKA improves infection control . Clin Orthop Relat Res. 2011, 469:977-84. 10.1007/s11999-010-1425-2
- Klouche S, Leonard P, Zeller V, et al.: Infected total hip arthroplasty revision: one- or two-stage procedure? . Orthop Traumatol Surg Res. 2012, 98:144-50. 10.1016/j.otsr.2011.08.018
- Kusuma SK, Ward J, Jacofsky M, Sporer SM, Della Valle CJ: What is the role of serological testing between stages of two-stage reconstruction of the infected prosthetic knee?. Clin Orthop Relat Res. 2011, 469:1002-8. 10.1007/s11999-010-1619-7
- Mahmud T, Lyons MC, Naudie DD, Macdonald SJ, McCalden RW: Assessing the gold standard: a review of 253 two-stage revisions for infected TKA. Clin Orthop Relat Res. 2012, 470:2730-6. 10.1007/s11999-012-2358-8
- Marculescu CE, Berbari EF, Hanssen AD, Steckelberg JM, Harmsen SW, Mandrekar JN, Osmon DR: Outcome of prosthetic joint infections treated with debridement and retention of components. Clin Infect Dis. 2006, 42:471-8. 10.1086/499234
- Tsaras G, Osmon DR, Mabry T, et al.: Incidence, secular trends, and outcomes of prosthetic joint infection: a population-based study, Olmsted county, Minnesota, 1969-2007. Infect Control Hosp Epidemiol. 2012, 33:1207-12. 10.1086/668421
- Duff GP, Lachiewicz PF, Kelley SS: Aspiration of the knee joint before revision arthroplasty. Clin Orthop Relat Res. 1996, 132-9. 10.1097/00003086-199610000-00018
- Sendi P, Banderet F, Graber P, Zimmerli W: Clinical comparison between exogenous and haematogenous periprosthetic joint infections caused by Staphylococcus aureus. Clin Microbiol Infect. 2011, 17:1098-100. 10.1111/j.1469-0691.2011.03510.x
- Kunutsor SK, Whitehouse MR, Blom AW, Beswick AD: Patient-related risk factors for periprosthetic joint infection after total joint arthroplasty: a systematic review and meta-analysis. PLoS One. 2016, 11:e0150866. 10.1371/journal.pone.0150866
- Batty LM, Lanting B: Contemporary strategies to prevent infection in hip and knee arthroplasty. Curr Rev Musculoskelet Med. 2020, 13:400-8. 10.1007/s12178-020-09653-9
- Berbari EF, Osmon DR, Lahr B, et al.: The Mayo prosthetic joint infection risk score: implication for surgical site infection reporting and risk stratification. Infect Control Hosp Epidemiol. 2012, 33:774-81. 10.1086/666641
- 63. Wilson MM, Philpot C, Morley JE: Anorexia of aging in long term care: is dronabinol an effective appetite stimulant? a pilot study. J Nutr Health Aging. 2007, 11:195-8.
- 64. Struwe M, Kaempfer SH, Geiger CJ, et al.: Effect of dronabinol on nutritional status in HIV infection . Ann Pharmacother. 1993, 27:827-31. 10.1177/106002809302700701
- Gavazzi G, Krause KH: Ageing and infection. Lancet Infect Dis. 2002, 2:659-66. 10.1016/S1473-3099(02)00437-1
- Singh JA, Sperling JW, Schleck C, Harmsen W, Cofield RH: Periprosthetic infections after shoulder hemiarthroplasty. J Shoulder Elbow Surg. 2012, 21:1304-9. 10.1016/j.jse.2011.08.067
- 67. Tahami SM, Aminian A, Azarpira N: Experimental study on protective role of NSAID on articular cartilage destruction in septic arthritis. Arch Bone Jt Surg. 2020, 8:89-93. 10.22038/abjs.2019.37045.2001
- Wei YP, Chien JC, Hsiang WH, Yang SW, Chen CY: Aspirin administration might accelerate the subsidence of periprosthetic joint infection. Sci Rep. 2020, 10:15967. 10.1038/s41598-020-72731-y
- 69. Sulfasalazine: MedlinePlus Drug Information. Accessed: November 26, 2022. (2022). Accessed: November 26, 2022: https://medlineplus.gov/druginfo/meds/a682204.html.
- Sulfasalazine (Azulfidine). Accessed: November 26, 2022. (2022). Accessed: 11/26/2022: https://www.rheumatology.org/I-Am-A/Patient-Caregiver/Treatments/Sulfasalazine-Azulfidine..
 Hydroxychloroquine (Oral Route) Side Effects - Mayo Clinic . (2022). Accessed: November 26, 2022:
- hydroxychiorodune (Ora Koure) side Enects Mayo Chinic . (2022). Accessed: November 26, 2022. https://www.mayoclinic.org/drugs-supplements/hydroxychloroquine-oral-route/side-effects/drg-20064216.
- 72. Goodman SM: Rheumatoid arthritis: perioperative management of biologics and DMARDs . Semin Arthritis Rheum. 2015, 44:627-32. 10.1016/j.semarthrit.2015.01.008