

Introduction to Musculoskeletal Infection

In-person training for Distributor Sales Representatives



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Training objectives

At the end of this 30 minute training session you will:

- Understand the fundamentals of musculoskeletal infection; its cause, impact on patients and cost to the healthcare system
- Be aware of the most common types of musculoskeletal infection and the treatment options available to your surgeons
- Have gained a working knowledge of the key terms used in discussions about musculoskeletal infection



Agenda

- Key terms
- Overview of musculoskeletal infection
- Surgical site infections
- Types of bacteria
- Biofilm
- Common orthopaedic infections
- Impact of musculoskeletal infection
- Treatment options
- Summary of key points





Key terms

Acute infection: An infection that is present for less than 30 days

Antibiotic: A drug that kills or inhibits the growth of bacteria but not viruses

Antimicrobial: An agent that kills microorganisms or inhibits their growth

Bacteria: A large group of unicellular microorganisms

Biofilm: A complex community of bacterial cells enclosed in a self-produced extracellular matrix (ECM) that adhere to an inert or living surface

Chronic infection: A prolonged or persistent invasion of the body by pathogens, which indicates that biofilm is present

Debridement: A surgical technique where dead, necrotic and infected tissue is surgically excised

Drug-resistant infections: Infections that are resistant to antibiotics commonly used to kill infections caused by resistant strains of bacteria

ECM: Extracellular matrix is a polysaccharide coating formed by bacteria that contains host proteins acquired by the bacterial network



Key terms

Gram Stain: A method used to differentiate two large groups of bacteria based on their cell wall constituents

Gram-negative bacteria: Have a cell wall composed of a thin layer of peptidoglycan that is located between an inner cell membrane and a bacterial outer membrane. Gram-negative bacteria take on the color of red/pink counterstain in the Gram's staining method

Gram-positive bacteria: Have a cell wall composed of a thick layer of peptidoglycan. Stain Gram-positive in Gram's method of staining because they retain the color of the crystal violet strain

HAI: Hospital acquired infection, which is a localized or systemic condition resulting from an adverse reaction to the presence of infectious agents or toxins that occurs in a healthcare setting

Inoculation: The introduction of an antigenic substance or vaccine into the body to produce immunity to a specific disease

MIC: Minimum inhibitory concentration is the lowest concentration of an antimicrobial that will inhibit the visible growth of a bacterium

Micro-organism: A microscopic organism that may exist as a single cell or in a colony of cells

Musculoskeletal infection: Invasion and multiplication of pathogenic microorganisms in the body



Key terms

Non-union: A fracture that will not heal

Osteomyelitis: An infection of bone and bone marrow caused by bacteria

Peptidoglycan: A polymer made of amino acids and sugars that makes up the cell wall of bacteria

PJI: Periprosthetic Joint Infection is a postoperative complication that can occur following total hip or knee arthroplasty by bacterial inoculation at the time of surgery or through open draining wounds

Pathogen: Infectious microorganisms such as bacteria, viruses and fungi

Persister cells: Dormant variants of regular cells in microbial populations that are highly tolerant to antibiotics

Planktonic bacteria: Free-floating bacteria behave as unicellular organisms and can be easily identified

Polymicrobial: Disease state involving multiple species of multiple organisms

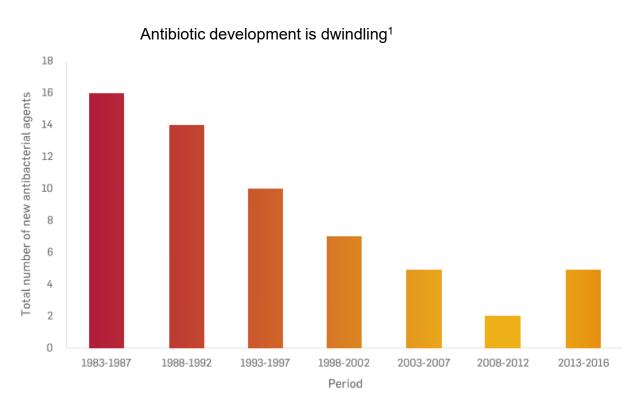
SSI: Surgical site infections, which are directly related to an operative procedure



Overview of musculoskeletal infection

Musculoskeletal infections are one of the biggest healthcare challenges in the 21st century

 Rapid growth of antibiotic-resistant strains and biofilms continues to outpace the development of new antibiotics and antibiotic strategies^{1,2}



- 50-60% of all hospital acquired infections are caused by antibiotic-resistant bacteria³
- In the USA, the cost of antibioticresistant infections to the healthcare system is \$21 billion to \$34 billion each year and more than 8 million additional hospital days⁴



Occurrence of musculoskeletal infection

Surgical site infections (SSI)

- Significant risk associated with surgery
- Present a huge burden on hospital resources and annual expenditure
- 1.7 million patients per year acquire an infection while in the hospital³
- Account for approximately 23% of all hospital acquired infections⁵
- Cost up to \$10 billion annually in direct medical expenses⁵





The cause of musculoskeletal infection

Bacteria

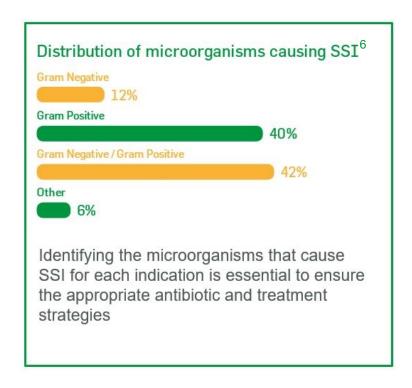
- Invasion of pathogenic unicellular micro-organisms
- Most common source for bacteria in hospitals is people⁷
- Differentiated into two distinct types

Gram-positive bacteria⁸

- Found in skin particles and spread through the air or by direct contact with coated objects
- More receptive to antibiotics than Gram-negative bacteria

Gram-negative bacteria^{9,10}

- Found and transferred through direct contact with wet objects
- Resistant to most available antibiotics





Gram's method of staining

Gram Stain

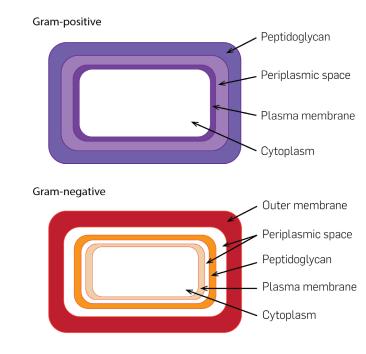
- A technique used to differentiate bacteria based on the difference in their cell wall properties
 - Distinguishes between Gram-positive and Gram-negative bacteria by coloring their cells violet or red

Gram-positive bacteria:

- Cell wall composed of a thick layer of peptidoglycan
- Stain Gram-positive: the cell wall retains the color of the crystal violet stain

Gram-negative bacteria:

- Cell wall composed of a thin layer of peptidoglycan, located between an inner cell membrane and a bacterial outer membrane
- Stain Gram-negative: the thin cell wall does not retain the crystal violet stain, and takes on the color of the red counterstain





Minimum Inhibitory Concentration (MIC)

Overview

- MIC is the lowest concentration of an antimicrobial that will inhibit the visible growth of a bacterium
- MIC is used by clinicians to choose which antibiotics to administer for specific infections and to identify the clinically effective dose of antibiotic
 - Different antibiotics are used for Gram-positive, Gram-negative and polymicrobial infections
 - Polymicrobial infections often require more than one antibiotic to target all bacterial pathogens
- This is important because populations of bacteria exposed to an insufficient concentration of a particular drug or to a broad-spectrum antibiotic, can evolve resistance to those drugs
- MIC scores aid in improving outcomes for patients and preventing drug-resistant microbial strains evolving



Planktonic bacteria vs. biofilm

Forms of bacteria in the body¹¹

Planktonic bacteria

- Free-floating unicellular organisms
- Adhere to tissue or implant surfaces
- Easily identified
- Cleared by host's immune system or antibiotics

Biofilm

- Complex community of bacterial cells enclosed in a self-secreted extracellular matrix (ECM)
- Can firmly attach to inert or living surfaces
- Difficult to detect
- Resistant to immune system and antibiotics
- Cause of chronic disease



Biofilm lifecycle

Timeline for biofilm development¹²

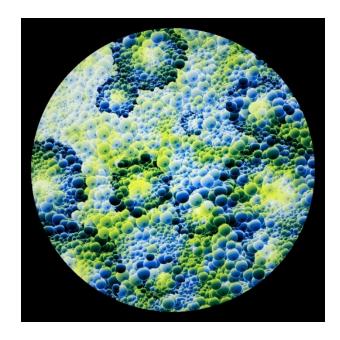
Biofilm stage	Timeline
Free-floating planktonic bacteria attach to a surface	minutes
Firmly attached microcolonies form	2-4 hours
Extracellular matrix develops	6-12 hours
Fully mature biofilm entering a dormant "persister" state	2-4 days
Biofilm re-forms after dispersion	24 hours



Biofilm

Properties of biofilm that lead to infection

- Limits the penetration of antimicrobials¹³
- pH differences allow some bacteria to become dormant while others remain active¹³
- Persister cells can withstand antibiotic attack and develop resistance to antibiotics¹⁴
- Difficult to detect and eradicate¹³
- Ability to avoid body's immune defenses¹⁵
- No methods or chemicals exist to dissolve a biofilm¹³

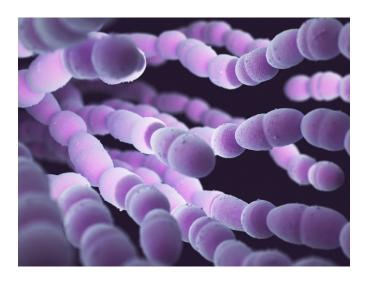




Types of musculoskeletal infection

Common infections in orthopaedics

- Periprosthetic joint infection
- Osteomyelitis
- Infected non-unions
- Diabetic foot ulcers





Periprosthetic joint infection (PJI)

Overview

- Postoperative complication following arthroplasty procedures
 - Occurs in 2.1% of total hip and 2.3% of total knee arthroplasty procedures¹⁶
 - 15% of hip revisions are carried out due to PJI¹⁶
 - 25% of knee revisions are carried out due to PJI¹⁶
- Acute PJI:
 - Present for less than 30 days
 - Biofilm is not established and treatment is focused on implant preservation
- Chronic PJI:
 - A prolonged or persistent invasion of the body by pathogens, which indicates that biofilm is present
 - Two stage procedures are often required for the treatment of the infection and for revision arthroplasty

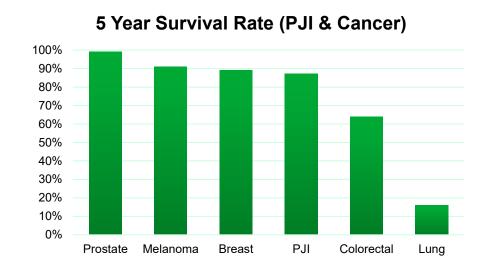




Periprosthetic joint infection (PJI)

Impact

- The annual hospital cost for hip and knee PJI is expected to reach \$1.85 billion by 2030¹⁶
- Revisions for PJI are associated with a fivefold increase in mortality compared to revisions for aseptic failure¹⁷
- PJI has a lower 5 year survival rate than the three most common cancers¹⁷
 - 2018 data: The 5 year survival rate for Medicare patients with PJI was 67% after total hip arthroplasty and 72% after total knee arthroplasty¹⁸
 - 2013 data: The relative 5 year survival rate for patients with PJI was 87.3%¹⁷





Acute PJI

Treatment example

- Debridement (used when joint components are not loose):
 - Incision and arthrotomy
 - Irrigation and debridement
 - Polyethylene insert exchange
 - Antibiotic therapy
- One-stage resection arthroplasty:
 - Incision and arthrotomy
 - Irrigation and debridement
 - Removal of cement and infected total knee components
 - Immediate implantation of new total knee components
 - Lavage and periprosthetic antibiotic therapy



Chronic PJI

Treatment example – 2 stage resection arthroplasty

- Stage 1 Focused on eradicating the infection:
 - Incision and arthrotomy
 - Resection of implant components
 - Irrigation and debridement
 - Use of antibiotic impregnated spacers to maintain space and provide stability
- Stage 2 Occurs after infection is eradicated:
 - Incision and arthrotomy
 - Removal of antibiotic spacers
 - Irrigation and debridement
 - Implantation of revision arthroplasty components
 - Dead space and soft tissue management





Osteomyelitis

Overview

- Osteomyelitis is an infection of bone and bone marrow
 - The incidence of osteomyelitis in the U.S. is reported to be 50,000 cases annually¹⁹
- Common in infected non-unions and diabetic foot ulcers
- Chronic osteomyelitis occurs in 5-50% of open fractures and is associated with significant morbidity²¹
- The recurrence rate of chronic osteomyelitis in adults is 30% within 12 months²⁰





Infected non-unions

Overview

- Post-operative fracture that will not heal
 - 10% of patients who sustain fractures (790,000) have impaired bone healing, resulting in delayed union or non-union²²
- Caused by contamination by bacteria and failure of fixation
- Risk of infection depends on the type of fracture:
 - Closed fractures: Infection rates ≤ 5%²³
 - Open fractures: Infection rates ≤ 50%²³
- Complications:
 - Bone and soft tissue loss
 - Internal fixation loosening or breakage
 - Poor vascularity of the bony fragments
 - Osteomyelitis

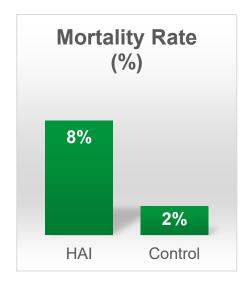




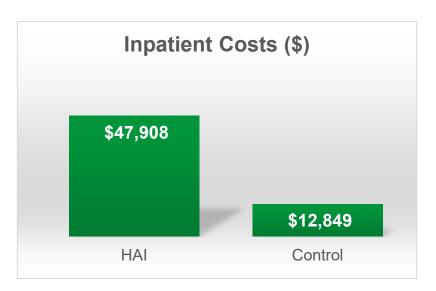
Infected non-unions

A study was conducted to explore the clinical impact and economic burden of hospital acquired infections (HAIs) in trauma patients²⁴

- Increased mortality rates for patients with HAIs
- Longer hospitalization periods for patients with HAIs
- Higher costs for patients with HAIs









Infected non-unions

Treatment example

- Removal of original hardware
- Debridement
- Revision stabilization of the fracture
- Supplemental bone grafting
 - Unmanaged spaces may contribute to infection
- Treatment of infection with antibiotics
- Two stage treatments may also be required to treat the infection and stabilize the non-union





Diabetic foot ulcers

Overview

- A foot ulcer is an open sore or wound that is commonly located on the lower legs or bottom of the feet
 - 7.3 million people affected by diabetes in the U.S. have a lifetime risk of developing a foot ulcer^{25,26}
- If untreated, a diabetic foot ulcer can become infected and lead to osteomyelitis
 - The prevalence of osteomyelitis ranges from 10-20%²⁷
- Osteomyelitis is the leading cause of non-traumatic lower extremity amputations
 - 60% of lower extremity amputations are caused by osteomyelitis²⁷

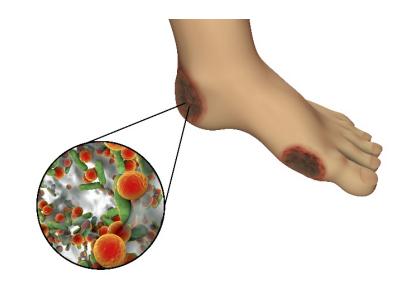




Diabetic foot ulcers with osteomyelitis

Impact

- Diabetic foot ulcers impose a tremendous medical and financial burden on our healthcare system
 - Patients require frequent emergency room visits, have increased hospital readmissions and require longer lengths of stay
 - Treatment costs are estimated to be ~\$45,000 per patient²⁸
 - Ulcer care adds \$9-\$13 billion to the direct yearly costs associated with diabetes²⁹
- Remission rates for osteomyelitis can range up to 88%²⁹
 - Approximately 50% of patients who have foot amputations due to diabetes die within five years³⁰
 - 40% of amputations could be prevented with appropriate wound care²⁶





Diabetic foot ulcers with osteomyelitis

Treatment example

- Diagnosis of osteomyelitis
- Remove load from the ulcer site to redistribute force
- Debridement of infected bone
- Resection of compromised soft tissue
- Antimicrobial therapy
- Wound dressings with regular changes
- Increased healing rates and decreased length of antibiotic treatment have been reported with early surgical intervention for osteomyelitis²⁹

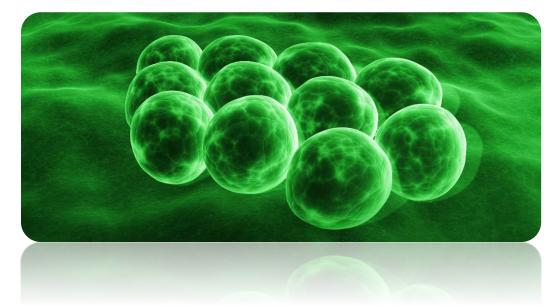




Musculoskeletal infection

Key points

- One of the biggest healthcare challenges of the 21st century
- Present a tremendous medical and economic burden on the healthcare system
- Represents a major cause of patient morbidity and mortality
- Care delivery is becoming increasingly challenging and complicated





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- Improving outcomes
- Reducing costs



Musculoskeletal infection

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Patents granted: EP 1390086 B1, US 8632796, CN ZL02809194.9, US 8496955, GB2367552, EP 1204599 B1, US 6780391, EP 1436019 B1, US 8563024, CN ZL02825134.2, EP 2594231 B1, US 8883063, CN ZL201210466117.X, GB2496710, EP 3058899 B1, US 10390954, US 10,588,748, CN ZL201610089710.5

Patents Pending: GB1502655.2, GB1704688.9, EP 18275044.8, US 15/933936, CN 108619579A

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References

- 1. Spellberg, B., et al., The Epidemic of Antibiotic-Resistant Infections: A Call to Action for the Medical Community from the Infectious Diseases Society of America. Clinical Infectious Diseases, 2008. 46(2): p. 155-164.
- 2. Gauland, C., Managing lower-extremity osteomyelitis locally with surgical debridement and synthetic calcium sulfate antibiotic tablets. Adv Skin Wound Care, 2011. 24(11): p. 515-23.
- 3. Reportlinker, 2013.
- 4. Golkar, Z., Bagasra, O., Pace, D..Bacteriophage therapy: a potential solution for the antibiotic resistance crisis. The Journal of Infection in Developing Countries, North America, 8, Feb. 2014.
- 5. Shea KG et al. Surgical Site Infection Reduction Program: Challenges and Opportunities. J Pediatr Orthop. 2015. Jul-Aug; 35 (5 Suppl 1): S51-4.
- 6. Public Health England. Surveillance of surgical site infections in NHS hospitals in England, 2017 to 2018, December 2018. Available at: https://www.gov.uk/government/publications/surgical-siteinfections-ssi-surveillance-nhs-hospitals-in-england Last accessed: December 2019.
- 7. Collins, Amy S. "chapter 41: Preventing Health Care Associated Infections." *Patient Safety and Quality: An Evidence-Based Handbook for Nurses*. https://www.ncbi.nlm.nih.gov/books/NBK2683/
- 8. Muilwijk J., et al., Random Effect Modeling of Patient-Related Risk Factors in Orthopaedic Procedures: Results from the Dutch Nosocomial Infection Surveillance Network 'PREZIES'. J Hosp Infect. 2006 Mar; 62(3): 319-26. PudMed PMID: 16406851.
- 9. Rodriguez-Pardo D., et al., Gram-negative Prosthetic Joint Infection: Outcome of a Debridement, Antibiotics and Implant Retention Approach. A large multicentre study. Clin Microbiol Infect. 2014 Nov; 20(11):0911-9. doi: 10.1111/1469-0691.12649. PubMed PMID: 24766536.
- 10. Jamei O., *et al.*, Which Orthopaedic Patients are Infected with Gram-negative Non-fermenting Rods?". J Bone Joint Infect 2017; 2(2):73-76. doi: 10.7150/jbji.17171. Available from http://www.jbji.net/v02p0073.htm.
- 11. McConoughey SJ, Howlin R, Granger JF, et al. Biofilms in periprosthetic orthopedic infections. Future Microbiol. 2014;9(8):987-1007.
- 12. Fehring TK, Odum SM, Berend KR, et al. Failure of irrigation and debridement for early postoperative periprosthetic infection. Clin Orthop Relat Res. 2013;471(1):250-257.
- 13. McPherson E.J., Peters, C.L, *Musculoskeletal Infection*, Orthopaedic Knowledge Update 10, American Academy of Orthopaedic Surgeons, 2011, 20: p. 239-258.
- 14. https://phys.org/news/2014-06-bacteria-dormancy-antibiotic-drugs.html
- 15. Shah SR, Tarata AM, D'Souza RN, Mikos AG, Kasper FK. Evolving strategies for preventing biofilm on implantable materials. *Materials Today*. 2013;16(5):177-182.
- 16. Premkumar A, Kolin DA, Farley KX, Wilson JM, McLawhorn AS, Cross MB, Sculco PK. Projected Economic Burden of Periprosthetic Joint Infection of the Hip and Knee in the United States. J Arthroplasty. 2020 Dec 9:S0883-5403(20)31244-4. doi: 10.1016/j.arth.2020.12.005. Epub ahead of print. PMID: 33422392.
- 17. Zmistowski, Benjamin; Karam, M.D., Joseph A.; Durinka, Joel B; Casper, MD, David S; and Parvizi, Javad MD, Periprosthetic Joint Infection Increases the Risk of One-year Mortality. 2013. Rothman Institute. Paper 44.
- 18. Fehring TK, Fehring KA, Hewlett A, Higuera C, Otero J, Tande, A. What's new in musculoskeletal infection. J Bone Joint Surg AM. 2019;101:1237-44.
- 19. Momodu II, Savaliya V. Osteomyelitis. [Updated 2019 Dec 8]. In: StatPearls [Internet]. Treasure Island (FL): StatPearls Publishing; 2020 Jan-. Available from: https://www.ncbi.nlm.nih.gov/books/NBK532250/



References

- 20. Hatzenbuehler J, Pulling, TJ. Diagnosis and Management of Osteomyelitis. Am Fam Physician. 2011;84(9):2017-1033.
- 21. Lima A.L., Oliveira P.R., Carvalho V.C., et al. Recommendations for the treatment of osteomyelitis. Braz J Infect Dis. 2014;18(5):526-534.
- 22. Wu N., Lee Y., Segina D., Murray H., Wilcox T., Boulanger L. Economic burden of illness among US patients experiencing fracture nonunion. Orthopedic Research and Reviews. March 2013. 2013(5):21-23.
- 23. Ruedi, T.P., Buckley, R.E., Moran, C.G. AO Principles of Fracture Management. 2015. https://www2.aofoundation.org/wps/portal/surgerymobile?contentUrl=/srg/popup/further_reading/PFxM2/45_Antibio_prophyl.jsp&soloState=precomp&title=&Language=en.
- 24. Glance, L.G., et al., Increases in mortality, length of stay, and cost associated with hospital-acquired infections in trauma patients. Arch Surg, 2011. 146(7): p. 794-801.
- 25. National Center for Chronic Disease Prevention and Health Promotion, C.f.D.C.a.P. *National Diabetes Statistics Report. Estimates of Diabetes and Its Burden in the United States*, 2014. Atlanta, GA: US Department of Health and Human Services; 2014.
- 26. Clayton, W. and T.A. Elasy, A Review of the Pathophysiology, Classification, and Treatment of Foot Ulcers in Diabetic Patients. Clinical Diabetes, 2009. 27(2): p. 52-58.
- 27. Thomas-Ramoutar C., E.T., DPM., Robert Frykberg, DPM, MPH, *Osteomyelitis and Lower Extremity Amputations in the Diabetic Population*. The Journal of Diabetic Foot Complications, 2010. **2**(1): p. 18-27.
- 28. Wu, S.C., et al., Foot Ulcers in the Diabetic Patient, Prevention and Treatment. Vasc Health Risk Manag, 2007. 3(1): p. 65-76.
- 29. Rice, J.B., et al., Burden of Diabetic foot Ulcers for Medicare and Private Insurers. Diabetes Care, 2013. http://care.diabetesjournals.org/content/early/2013/10/29/dc13-2176.abstract.
- 30. https://bmcsurg.biomedcentral.com/articles/10.1186/1471-2482-14-83

