# Microbiology of Bone and Joint Infections (Osteomyelitis & Septic Arthritis)

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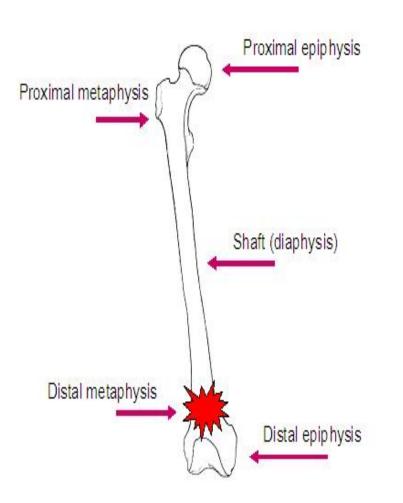
# **Objectives**

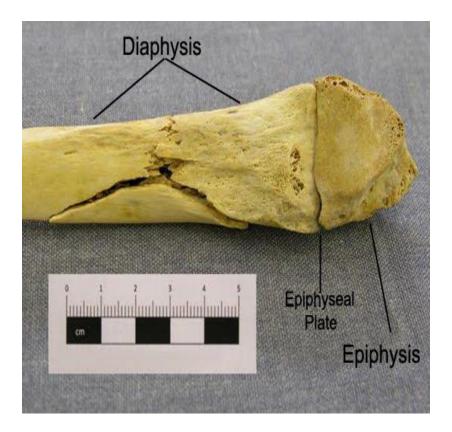
- Define osteomyelitis and arthritis
- Know that the two conditions can happen together or separately.
- Differentiate between acute and chronic osteomyelitis and arthritis
- Know the pathogenesis and risk factors of both osteomyelitis and arthritis
- Realize that bone and joint infections can be acquired through blood or directly from adjacent affected organs and tissues.

- Know the commonest causative agents of arthritis and osteomyelitis.
- Know the laboratory diagnosis and investigation of both conditions.
- know the management and treatment of both osteomyelitis and arthritis.

# Introduction

- Bone & joint infections may exist separately or together.
- Both are more common in infants and children.
- Usually caused by blood borne spread ,but can result from local trauma or spread from contiguous soft tissue infection.
- Often associated with **foreign body** at the primary wound site.
- If not treated lead to devastating effect.





# **Acute Osteomyelitis**

- Acute osteomyelitis is an acute infectious process of the bone and bone marrow .
- How the pathogen reach the bone ?
- 1- Hematogenous route
- 2- Contiguous soft tissue focus ( post operative infection, contaminated open fracture, soft tissue infection , puncture wounds)
- **3- In association with peripheral vascular disease** (diabetes mellitus ,severe atherosclerosis, vasculitis)
- May have a short duration (few days for hematogenously acquired infection) or may last several weeks to months (if secondary to contiguous focus of infection).

# **Etiology, Epidemiology & Risk Factors**

• **Primary hematogenous** is most common in infants & children.

Infants: S.aureus, group B streptococci, E.coli.

Children: S.aureus, group A streptococci, H.influenzae.

Site : Metaphysis of long bones (femur,tibia& humerus )

Adults: Hematogenous cases less common, but may occur due to reactivation of a quiescent focus of infection from infancy or childhood. Most cases are due to *S.aureus*.

Septic arthritis is common as the infection begins in the Diaphysis.

### **Other causes -special clinical situations**

- Streptococci and anaerobes in fist injuries, diabetic foot and decubitus ulcers.
- Salmonella or Streptococcus pneumoniae in sickle cell patients.
- *Mycobacterium tuberculosis* (MTB) or *Mycobacterium avium* in AIDS patients.

## Common causes of acute Osteomyelitis

#### Age /special conditions

- Infants
- Children
- Adults
- Sickle cell disease
- Infection after trauma ,injury or surgery
- AIDS patients

#### Common causative organism

- S.aureus, group B Streptococcus,Gram negative rods (eg. E.coli, Klebsiella).
- S. aureus, group A Streptococcus & H. influenzae
- S.aureus
- S.aureus, S. pneumoniae, Salmonella species
- S.aureus, group A Streptococcus,Gram negative rods, anaerobes.
- MTB or M.avium.

# **Clinical presentation & blood tests**

- Acute osteomyelitis usually of abrupt onset
- fever
- localized pain , heat , swelling
- Tenderness of affected site ( one or more bones or joints affected in hematogenous spread).
- May be local tissue infection( abscess or wound) .
- **Blood test**: leukocytosis, high ESR and C-reactive protein.

# Diagnosis

### • Blood culture

- Blood culture **or aspiration** of overlying abscess if blood cultures are negative.
- Leukocytosis ( high WBCs) may or may not occur.
- Erythrocyte sedimentation rate ( ESR) elevated or normal.

### • Imaging:

1. X-RAY, MRI, CT-SCAN

### Treatment

- MSSA ( methicillin sensitive *S.aureus*) : Cloxacillin, or Clindamycin .
- MRSA( methicillin resistant *S.aureus*): Vancomycin followed by Clindamycin, Linezolid, or TMP-SMX.
- **Polymicrobial infection**: Piperacillin-Tazobactam or Quinolone with Metronidazole.

# **Chronic Osteomyelitis**

- A chronic infection of the bone and bone marrow usually secondary to inadequately treated or relapse of acute osteomyelitis.
- Management difficult , prognosis poor.
- Infection may not completely cured.
- May recur many years or decades after initial episode.
- Most infections are secondary to a contiguous focus or peripheral vascular disease.
- Chronic infection due to hematological spread is rare.

# **Chronic Osteomyelitis**

- S.aureus is the most common pathogen
- Other microorganisms: S.epidermidis, Enterococci, streptococci, Enterobactericae, Pseudomonas and anaerobes.
- Polymicrobial infection common with decubitus ulcers and diabetic foot infections.
- **TB and fungal osteomyelitis** clinically have indolent "chronic" course

# **Chronic Osteomyelitis**

- Mycobacteria and fungi may be seen in immunosuppressed patients.
- *MTB* osteomyelitis primarily results from haemtogenous spread from lung foci or as an extension from a caseating lymph bone ( 50% in spine). It resembles *Brucella* oesteomyelitis .
  - TB & Brucella are common in KSA.
- Haematogenous osteomyelitis due to fungi eg. Candida species, Aspergillus species and other fungi may occur.

# Diagnosis

- Blood culture is not very helpful because bacteremia is rare.
- WBC usually normal, ESR elevated but not specific.
- Radiological changes are complicated by the presence of bony abnormalities.
- MRI helpful for diagnosis and evaluation of the extent of disease.

## **Blood Culture & Osteomyelitis Images**







Fig. 7. Recompanyous of latitized. The process photons of the town, digit down with anyonizat processed thickening.





## **Management & Treatment**

- Extensive surgical debridement with antibiotic therapy. Parenteral antibiotics for 3-6 weeks followed by long term oral suppressive therapy.
- Some patients may require life long antibiotic ,others for acute exacerbations.
- MSSA: Cloxacillin
- *MRSA & S.epidermidis*: Vancomycin then oral Clindamycin or TMP-SMX.
- Other bacteria: treat as acute osteomyelitis.
- **MTB**: 4 drugs : INH,RIF ,Pyrazinamide & Ethambutol for 2 months followed by RIF + INH for additional 4 months. *Brucella* is treated with Tetracycline and Rifampicin for 2 to 3 months.



# **Septic Arthritis**

**Septic (Infectious) Arthritis** is inflammation of the joint space secondary to infection.

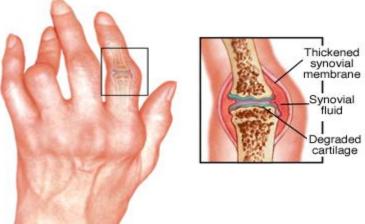
Generally affects a single joint and result in suppurative inflammation.

### Haematogenous seeding of joint is most common.

- Common symptoms :pain, swelling, limitation of movement.
- Diagnosis by **Arthrocentesis** to obtain synovial fluid for analysis; Gram stain, culture & sensitivity
- Drainage & antimicrobial therapy important management.

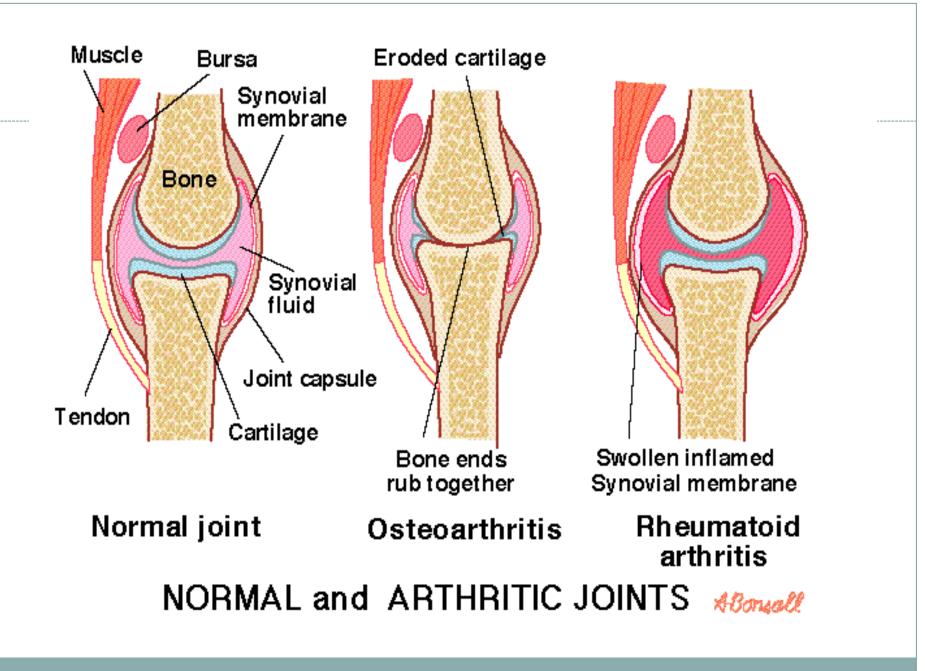
# **Septic Arthritis**







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### Common causes of septic arthritis

### **Age/special conditions**

- Neonates
- Infants /children
- Adults
- Sickle cell disease
- Trauma /surgical procedure
- Chronic arthritis
- Prosthetic arthritis

#### **Common organism**

- S.aureus, group B streptococcus, Gram negative rods.
- S.aureus, group A streptococcus,
  S.pneumoniae, H. influenzae type b
- S.aureus, Neisseria gonorrheae
- Salmonella species, S.aureus
- S.aureus
- MTB, Fungi
- Skin flora

# **Risk factors & Pathogenesis**

- Gonococcal infection most common cause in young, sexually active adults caused by *Neisseria* gonorrheae. Leads to disseminated infection secondary to urethritis/cervicitis. Initially present with polyarthralgia, tenosynovitis, fever, skin lesions. If untreated leads to suppurative monoarthritis.
- **Nongonococcal** arthritis occurs in older adults. Results from introduction of organisms into joint space as a results of bacteremia or fungaemia from infection at other body sites.

# **Risk factors**

Occasionally results from direct trauma, procedures (arthroscopy) or from contiguous soft tissue infection.

- *S.aureus* is most common cause. Other organisms : streptococci and aerobic Gram negative bacilli.
- Lyme disease due to tick bite in endemic areas. Uncommon in KSA.
- In sickle cell disease patients , arthritis may be caused by *Salmonella* species.
- Chronic arthritis may be due to MTB or fungi.

# **Diagnosis of Septic Arthritis**

- History/examination to exclude systemic illness. Note history of tick exposure in endemic areas
- Arthrocentesis should be done as soon as possible; 1-Synovial fluid is cloudy and purulent
  - 2- Leukocyte count generally > 50,000/mm3,with > 75 % neutrophils.
  - 3- Gram stain and culture are positive in >90% of cases.
  - 4-Exclude crystal deposition arthritis or noninfectious inflammatory arthritis.

### Blood cultures indicated

- If gonococcal infection suspected, take specimen from cervix, urethra, rectum & pharynx for culture or DNA testing for *N.gonorrheae*.
- Culture of joint fluid and skin lesions .

### **Treatment & Management**

- Arthrocentesis with drainage of infected synovial fluid.
- Repeated therapeutic Arthrocentesis often needed
- Occasionally, arthroscopic or surgical drainage/debridement
- Antimicrobial therapy should be directed at the suspected organism and susceptibility results:
- Gonococcal arthritis: IV Ceftriaxone (or Ciprofloxacin or Ofloxacin) then switch to oral Quinolone or Cefixime for 7-10 days.

### • <u>Nongonococcal infectiuos arthritis</u>:

- 1. MSSA: Cloxacillin or Cefazolin
- 2. MRSA: Vancomycin
- 3. Streptococci: Penicillin or Ceftriaxone or Cefazolin
- *4. Enterobacetriacae*: Ceftriaxone or Fluroquinolone
- 5. *Pseudomonas:* Piperacillin and Aminoglycoside
- 6. Animal bite : Ampicillin-Sulbactam
- Lyme disease arthritis: Doxycycline for 1 month.

# **Prognosis & Complications**

- Gonococcal arthritis has an excellent outcome .
- Non-gonococcal arthritis: can result in scarring with limitation of movement, ambulation is affected in 50% of cases.
- **Risk factors** for long term adverse sequelae include:

Age, prior rheumatoid arthritis, polyarticular joint involvement, hip or shoulder involvement, virulent pathogens and delayed initiation or response to therapy.

# **Infections of Joint Prosthesis**

- Occur in 1 5 % of total joint replacement.
- Most infections occur within 5 years of joint replacement.
- Often caused by skin flora.
- Diagnostic aspiration of joint fluid necessary .
- Result in significant morbidity and health care costs.
- Successful outcomes results from multidisciplinary approach.

## **Joint Prosthesis**





Before





\*ADAM

# **Diagnosis of Prosthetic Arthritis**

- Aspiration & surgical exploration to obtain specimen for culture , sensitivity testing & histopathology.
- Skin flora regarded as pathogens if isolated from multiple deep tissue cultures.
- **Plain X-ray** may not be helpful.
- Arthrography may help define sinus tracts.
- Bone scan-not specific for infection.
- ESR and C-reactive protein( CRP ) may be high.

## **Management & Treatment**

- Surgical debridement and prolonged antimicrobial therapy
- Surgery: removal of prosthesis
- Antibiotic –impregnated cement during reimplantation
- Antimicrobial for 6 weeks:
- Begin empiric IV antibiotic to cover MRSA and Gram negative rods (Vancomycin+ Cefepime, Ciprofloxacin, or Aminoglycoside)
- Chronic therapy with oral drug if removal of prosthesis not possible.