

## 1.Osteomyelitis

### Osteomyelitis - Key Points

Definition: Bone infection, usually bacterial

Causes:

Hematogenous spread (children)

Direct inoculation (trauma, surgery)

Contiguous spread (diabetes, ulcers)

Common organism: Staphylococcus aureus

Symptoms:

Fever, localized pain, swelling

Limited movement, erythema

Chronic: draining sinus

Diagnosis:

↑ WBC, ESR, CRP

Blood cultures

X-ray, MRI (best test), Bone biopsy

Treatment:

IV antibiotics (6 weeks)

Surgical debridement (if needed)

Chronic cases: bone graft, amputation (rare)

## 2.Duchenne Muscular Dystrophy (DMD)

X-linked recessive (DMD gene mutation)

Dystrophin deficiency → muscle degeneration



Onset: 2-5 years, affects males

Symptoms:

Proximal muscle weakness (pelvic & shoulder)

Gower's sign, waddling gait

Calf pseudohypertrophy

Frequent falls, delayed milestones

Cardiac (DCM) & respiratory failure

Diagnosis:

↑ CK levels, genetic testing

Muscle biopsy (absent dystrophin)

EMG, ECG, ECHO

Treatment:

Steroids (prednisolone/deflazacort)

Physiotherapy, bracing, wheelchair

Cardiac & respiratory support

Gene therapy (research phase)

Complications:

Loss of ambulation (by 12 years)

Respiratory & cardiac failure (~20-30 years)

Genetic counseling & prenatal testing

3. Basal cell carcinoma

Definition: Most common, slow-growing skin cancer

Risk Factors:



UV exposure, fair skin, chronic sun exposure

Immunosuppression, genetic predisposition

Common Sites: Face, nose, ears, neck

Appearance:

Pearly, translucent nodule with telangiectasia

Central ulceration (rodent ulcer)

Slow growth, rarely metastasizes

Diagnosis: Clinical + Biopsy

Treatment-

Surgical excision (gold standard)

Mohs micrographic surgery (for cosmetically sensitive areas)

Cryotherapy, radiotherapy (if surgery not possible)

#### 4.Warts

Warts - Key Points

Definition: Benign skin growths caused by human papillomavirus (HPV)

Types:

Common warts (Verruca vulgaris) – rough, raised, on hands/fingers

Plantar warts – soles of feet, painful

Flat warts (Verruca plana) – smooth, face/hands

Genital warts (Condyloma acuminata) – sexually transmitted

Transmission: Direct contact, autoinoculation

Diagnosis: Clinical examination

Treatment:



Salicylic acid, cryotherapy (first-line)

Electrocautery, laser therapy, surgical removal

Immunotherapy (imiquimod, intralesional bleomycin) for resistant cases

Prevention: HPV vaccine, avoid direct contact

## 5. Laparoscopic and robotic surgery

Laparoscopic Surgery

Minimally invasive

Hand-held instruments, direct surgeon control

2D camera view

Limited dexterity & range of motion

Robotic Surgery

Advanced form of laparoscopy

Robotic arms controlled by the surgeon

3D HD visualization

Greater precision, flexibility, and dexterity

Less surgeon fatigue

## 6. Stevens-Johnson Syndrome (SJS) & Toxic Epidermal Necrolysis (TEN)

Severe mucocutaneous reactions (drug-induced)

SJS: <10% body surface area (BSA) detachment

TEN: >30% BSA detachment

Causes: Drugs (Sulfa, NSAIDs, Anticonvulsants, Antibiotics)



Symptoms: Fever, malaise → painful blisters, skin sloughing

Nikolsky sign: Positive

Treatment:

Stop offending drug immediately

ICU/burn unit care, IV fluids, wound care

IVIg, cyclosporine (severe cases)

No steroids in TEN

## 7. iron def anemia

Iron Deficiency Anemia (IDA) - Key Points

Cause: Low iron intake, blood loss, malabsorption

Common in: Women (menstruation, pregnancy), children, elderly

Symptoms: Fatigue, pallor, pica, glossitis, koilonychia (spoon nails)

Labs:

↓ Hemoglobin, ↓ MCV (microcytic anemia)

↓ Serum iron, ↓ Ferritin, ↑ TIBC

Treatment:

Oral iron (ferrous sulfate) + vitamin C

IV iron (if severe or malabsorption)

Treat underlying cause

## 8. Electrocautery

Definition: Use of heat from electrical current to cut, coagulate, or destroy tissue



Types:

Monopolar (needs grounding pad)

Bipolar (no grounding pad needed)

Uses:

Surgical hemostasis (control bleeding)

Tissue destruction (warts, tumors)

Incision & dissection

Advantages: Minimal blood loss, precise tissue control

Complications: Burns, thermal injury, smoke inhalation risks

## 9. Osteoarthritis (OA)

Definition: Degenerative joint disease with cartilage loss

Risk Factors: Age, obesity, joint overuse, trauma

Joints Affected: Knees, hips, hands (DIP, PIP), spine

Symptoms: Joint pain, stiffness (morning <30 min), crepitus

X-ray Findings: Joint space narrowing, osteophytes, subchondral sclerosis

Treatment:

Lifestyle: Weight loss, exercise, physiotherapy

Medications: NSAIDs, acetaminophen, intra-articular steroids

Advanced cases: Joint replacement surgery

## 10. Aplastic anemia

Definition: Bone marrow failure → pancytopenia



Causes:

Idiopathic (most common)

Drugs (NSAIDs, chemotherapy, antibiotics)

Toxins (benzene, radiation)

Viral infections (EBV, Hepatitis, HIV)

Autoimmune diseases

Symptoms: Fatigue, infections, bleeding/bruising

Labs: ↓ WBC, ↓ RBC, ↓ Platelets, hypocellular bone marrow

Treatment:

Supportive: Transfusions, infection control

Immunosuppressive therapy (ATG, cyclosporine)

Bone marrow transplant (severe cases)

## 11. Chronic Myeloid Leukemia (CML)

Definition: Clonal myeloproliferative disorder of granulocytes

Cause: Philadelphia chromosome (t[9;22], BCR-ABL fusion gene)

Phases:

Chronic (asymptomatic, leukocytosis)

Accelerated (worsening symptoms, blasts 10-19%)

Blast crisis (resembles acute leukemia, blasts ≥20%)

Symptoms: Fatigue, weight loss, splenomegaly, night sweats

Labs: ↑ WBC, ↓ LAP score, basophilia, anemia

Diagnosis: PCR/FISH for BCR-ABL, bone marrow biopsy

Treatment:



Tyrosine kinase inhibitors (Imatinib, Dasatinib, Nilotinib)

Bone marrow transplant (for resistant cases)

## 12. Malaria

Cause: Plasmodium spp. (*P. falciparum*, *P. vivax*, *P. ovale*, *P. malariae*)

Transmission: Anopheles mosquito bite

Symptoms: Fever, chills, sweating (cyclical), headache, anemia, splenomegaly

Severe Malaria (*P. falciparum*): Cerebral malaria, shock, multi-organ failure

Diagnosis: Peripheral blood smear (thick & thin), rapid antigen test

Treatment:

Uncomplicated: Chloroquine (if sensitive), Artemisinin-based therapy (ACT)

Severe: IV Artesunate, supportive care

Prevention: Mosquito control, bed nets, chemoprophylaxis (for travelers)

## 13. Enhanced Recovery After Surgery (ERAS) - Six Key Components

1. Preoperative Optimization – Patient education, smoking cessation, nutritional support
2. Preoperative Fasting & Carbohydrate Loading – Minimize fasting, clear fluids up to 2 hours before surgery
3. Minimally Invasive Surgery – Laparoscopic/robotic approach when possible
4. Multimodal Analgesia – Reduce opioids, use regional anesthesia/NSAIDs
5. Early Mobilization – Encourage movement soon after surgery
6. Early Oral Nutrition – Resume feeding as soon as tolerated



#### 14. Thalassemia –

Definition: Genetic disorder causing defective hemoglobin synthesis

Types:

Alpha-thalassemia ( ↓  $\alpha$ -globin chain production)

Beta-thalassemia ( ↓  $\beta$ -globin chain production)

Clinical Forms:

Minor (trait): Mild anemia, asymptomatic

Intermedia: Moderate anemia, splenomegaly

Major (Cooley's anemia): Severe anemia, transfusion-dependent, growth delay

Symptoms: Pallor, fatigue, hepatosplenomegaly, skeletal deformities

Diagnosis:

Microcytic hypochromic anemia

Hemoglobin electrophoresis ( ↑ HbA<sub>2</sub>, HbF in beta-thalassemia)

Genetic testing

Treatment:

Regular blood transfusions (major cases)

Iron chelation therapy (to prevent iron overload)

Splenectomy (if needed)

Bone marrow transplant (curative in severe cases)

#### 15. Henoch-Schö nlein Purpura (HSP) - Key Points

Definition: IgA-mediated small vessel vasculitis

Common in: Children (3-10 years), post-infection

Triad:



Palpable purpura (buttocks, legs)

Arthritis/arthralgia (knees, ankles)

Abdominal pain (colicky, GI bleeding)

Renal Involvement: IgA nephropathy (hematuria, proteinuria)

Diagnosis: Clinical, ↑ IgA, kidney biopsy (if severe)

Treatment:

Supportive (hydration, NSAIDs for pain)

Steroids (for severe GI or renal involvement)

Monitor renal function

## 16. Acute Respiratory Distress Syndrome (ARDS)

Definition: Severe lung inflammation → non-cardiogenic pulmonary edema

Causes: Sepsis (most common), pneumonia, trauma, aspiration, pancreatitis

Symptoms: Severe dyspnea, hypoxia, respiratory failure

Diagnosis (Berlin Criteria):

Acute onset (<1 week of insult)

Bilateral lung opacities (CXR/CT)

PaO<sub>2</sub>/FiO<sub>2</sub> ratio ≤ 300 mmHg (hypoxemia severity)

Not due to heart failure (rule out cardiogenic edema)

Management:

Low tidal volume ventilation (lung-protective strategy)

PEEP to improve oxygenation

Prone positioning (severe cases)

Treat underlying cause



Supportive care (fluids, vasopressors if needed)

17.DMARDS SIDE EFFECTS



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## Anti-Rheumatic Drugs

## Agents

	Agents	Side Effects
1	<p><b>Methotrexate (MTX):</b> <i>once/wk: complete 8 days</i></p> <ul style="list-style-type: none"> <li>It is the best initial DMARD.</li> <li>It takes 1 – 2 months to take its effect.</li> <li>A 6 – month course should be given, before concluding that it has been ineffective</li> <li>Monitor = CBC &amp; LFTs monthly, then every 3 mo.</li> <li><b>Supplement with folic acid (5 mg/week)</b></li> </ul>	<ol style="list-style-type: none"> <li>Mouth ulcers</li> <li>Hepatotoxicity.</li> <li>Bone marrow suppression <i>AB</i></li> <li>Alopecia</li> <li>Pulmonary fibrosis</li> </ol>
2	<p><b>Anti-Tumor Necrosis Factor (TNF) Therapy:</b></p> <ul style="list-style-type: none"> <li>Agents: Etanercept, Adalimumab, Infliximab</li> <li>These are the first-line biologic agents for RA.</li> <li>These agents are often used in combination with MTX to prevent disease progression.</li> <li>These agents can be used as monotherapy, except for infliximab, which must be prescribed with MTX (reduces the risk of developing neutralizing antibodies)</li> <li><b>Always screen for TB (PPD and Chest x-ray) prior to starting treatment.</b></li> </ul>	<ol style="list-style-type: none"> <li>Reactivation of latent TB</li> <li>Reversible lupus-like syndrome</li> <li>Injection site reactions</li> <li>Heart failure</li> <li>Demyelination</li> <li>Increased risk of malignancy (basal cell carcinoma of skin)</li> </ol>
3	<p><b>Sulfasalazine</b></p> <ul style="list-style-type: none"> <li>It is a DMARD used in combination with MTX &amp; other agents.</li> <li>Monitor = CBC &amp; LFTs initially monthly, then every 3 months</li> </ul>	<ol style="list-style-type: none"> <li>Nausea &amp; GI upset, Rash</li> <li>Hepatitis</li> <li>Neutropenia, Pancytopenia</li> <li>Orange staining (urine, contact lenses)</li> </ol>
4	<p><b>Hydroxychloroquine</b></p> <ul style="list-style-type: none"> <li>It is a DMARD</li> <li><b>It can be used as monotherapy in seronegative &amp; mild disease.</b></li> <li>It is also used in combination with other DMARDs.</li> <li>Monitor = visual acuity &amp; fundoscopy before starting, then annually</li> </ul>	<ol style="list-style-type: none"> <li>Maculopapular rash</li> <li>Retinal toxicity</li> <li>Corneal deposits</li> </ol>
5	<p><b>Penicillamine:</b></p> <ul style="list-style-type: none"> <li>It is a DMARD</li> <li>It is less commonly used now.</li> <li>Monitor = CBC &amp; Urine for protein</li> </ul>	<ol style="list-style-type: none"> <li>Mouth ulcer</li> <li>Metallic taste</li> <li>Proteinuria</li> <li>Thrombocytopenia</li> </ol>
6	<p><b>Gold:</b></p> <ul style="list-style-type: none"> <li>It is a DMARD (sodium aurothiomalate)</li> <li>It is less commonly used now.</li> <li>Monitor = CBC &amp; Urine for protein</li> </ul>	<ol style="list-style-type: none"> <li>Mouth ulcer</li> <li>Alopecia</li> <li>Proteinuria</li> <li>Myelosuppression</li> </ol>

## 18. Consent & Types of Consent

Definition: Legal & ethical permission for medical intervention

Key Elements: Capacity, Voluntariness, Information, Understanding

Types of Consent:

1. Informed Consent – Patient fully understands risks, benefits, and alternatives
2. Implied Consent – Assumed in routine procedures (e.g., taking blood)
3. Verbal Consent – Spoken agreement for minor procedures
4. Written Consent – Signed document for major procedures/surgeries
5. Express Consent – Directly given (verbal or written)
6. Proxy Consent – Given by legal guardian (children, incapacitated adults)
7. Advance Directive – Pre-stated consent (living will, DNR orders)

## 19. Tinea Versicolor - Diagnostic Tests

Clinical Diagnosis: Hypo/hyperpigmented macules with fine scaling

Wood's Lamp Examination: Yellow-green fluorescence

KOH Mount (Microscopy): "Spaghetti and meatballs" appearance (hyphae & spores)

Fungal Culture: Rarely needed but confirms *Malassezia* spp.

Dermoscopy: Fine scales with yellowish hue

## 20. Gibbus Deformity with Osteoporosis

Gibbus Deformity: Sharp, angular kyphotic deformity of the spine

Causes in Osteoporosis:



Vertebral compression fractures (due to weak bones)

Multiple wedge fractures leading to kyphosis

Senile osteoporosis, postmenopausal osteoporosis

Symptoms: Back pain, height loss, hunched posture

Diagnosis: X-ray (wedge fractures), DEXA scan ( ↓ bone density)

Treatment:

Bisphosphonates, calcium, vitamin D

Pain management, physiotherapy

Bracing (if needed), surgical correction (severe cases)

