Diagnosis

Clinical features In general:

- Morning stiffness
- Easy fatigability in early afternoon
- Joint pain later in the day
- Joint swelling
- Involved joints often warm
- lacks full range of movement
- Occasionally painful on motion
- Usually not erythematous

CLINICAL MANIFESTATIONS of JRA







Investigations: No single test diagnostic of JIA

<u>Laboratory</u>

Haematology:

1.Ful blood examination may reveal normochromic normocytic anaemia, neutrophil leukocytosis or thrombocytosis.

2.ESR and CRP : often raised; sometimes normal.

- 3. Serum ferritin: Disproportionately high in Systemic JIA
- 4. Serum albumin: Hypoalbuminemia is common in SOJIA

5. Liver enzymes: high in Systemic JIA

6. Renal functions

7.Uric acid and LDH: as a screening test for malignancy in children with fever and joint symptoms.

Serology:

1.IgM rheumatoid factor: 10% seropositive, high IgM RF titre carry a worse prognosis.

2.ANA :positive in 50% . positive ANA is a risk factor for silent uveitis in oligoarthritis JIA.

3-Anti-eep: positive

Immunology:

- **1.Immunoglobins:** IgG may be raised; occasionally IgAmay be low or absent .
- **2.Compliment:** C3 often elevated ; a low C2 may occasionally be found ; raised alpha-2-globulin.

HLA typing:

- 1. B27 : positive in enthesitis-related arthritis in older males, psoriatric arthritis, arthritis associated with bowel disorders such as IBD.
- 2. DR4: positive in RF positive polyarthritis.
- **3. DR8** : positive in oligoarthritis , ANA positive younger female.

Imaging:

- A) Plain radiography: Plain radiographs are useful in excluding other differential diagnosis such as osteomyelitis, septic arthritis, trauma, and malignancy.
 - **Common findings on plain X-ray include:**
 - 1.Soft tissue swelling.
 - 2. Joint-space narrowing.
- 3. Periarticular osteoporosis.
- (The above three are the most common findings in RFnegative JIA)
- 4. Joint erosions .
- **5.** Leg-length discrepancy (accelerated maturation due to hyperaemia around the joint and low grade inflammation)

Radiographic changes(in Plain radiographs) in JIA

- Early: soft tissue swelling, e.g.blurring of infrapatellar fat pad on lateral knee radiograph and periarticular osteopenia.
- Intermediate: cortical erosions, joint space narrowing and subchondral sclerosis.
- Late:destructive joint changes with ankylosis, joint contractures, metaphyseal and diaphyseal changes and growth anomalies.



B) Ultrasound:

- Ultrasound can be useful in:
 - 1.Effusions (especialy useful in hips , shoulders.)
 - 2. Synovitis/tenosinovitis (increased echogenicity with inflamed tissues).
 - 3. Guiding intra-articular therapy.

C) Magnetic resonance imaging (MRI):

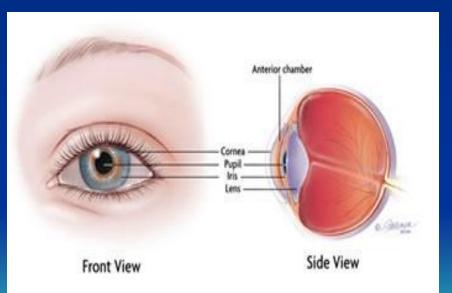
- MRI is useful in demonstrating anatomy,
- 1. Early joint damage.
- **2.** Positive effects of treatment before they are clinically apparent.
- **3.** Amount of inflamed synovium(when performed with gadolinium enhancement)

 Long-term effects in difficult-to-monitor joints(cervical spine, TM joints , hips.)

D) Bone scan: to identify unrecognized sites of inflammation

E) Slit lamp examination of eye to exclude uveitis .





Differential diagnosis

- 1. SOJIA should be differentiated from malignancy and infection by bone marrow examination
- 2. Systemic lupus erythematosus
- 3. Reactive arthritis
- 4. Septic arthritis

Complications..

- Joint destruction
- Leg length discrepancy,gait abnormalities
- Severe growth retardation
- Osteoporosis or osteopenia
- Muscle wasting
- Malnutrition,
- Anaemia of chronic disease,
- Chronic uveitis-higher in persistent pauciarthritis and ANA +ve patient.
- Untreated eye disease result in glaucoma,cataract,band keratopathy,blindness.
- Amyloidosis-in systemic arthritis.
- Macrophage activation syndrome: It is a rare but potentially life threatening complication of systemic JIA.

Complications Cont...

- Micrognathia
- Atlantoaxial sublaxation in poly arthritis RF+ve
- Mandibular asymmetry
- Complications of therapy



Macrophage activation syndrome. Actiology: it may be episode of viral infecton ,may arise denovo or associated with SOJIA

Symptoms:

- High unremitting fever
- Lymphadenopathy and hepatosplenomegaly
- DIC, encephalopathy Investigations:
- Pancytopenia, elevated liver enzymes and coagulopathy raises the suspicion of MAS.
- Hypertriglyceridemia and hyperferritinemia
- Decrease albumin, increase prothrombin time and partial thromboplastin
 - Paradoxically normal ESR (secondary to drop in fibrinogen as a consumptive coagulopathy)