#### **MANAGEMENT**

- Optimal management of JIA is delivered by an experienced multidisciplinary team.
  - Members of multidisciplinary team are as follows:
- 1. Medical: Rheumatologist, Opthalmologist, Radiologist.
- 2. Community: Family, Friends, School teacher.
- 3 .Professional allied to medicine:
   Physiotherapist, occupational therapist,
   Psychologist

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#### THE GOALS OF MANAGEMENT OF JIA

- The goals of JIA management are the following:
- 1. Provide analgesia.
- 2. Control inflammation.
- 3. Maintain joint function.
- 4. Prevent deformities.
- 5. Treat complications and extra-articular manifestations.
- 6. Ensure optimal nutrition.
- 7. Rehabilitition.
- 8. Ensure optimal psycosocial health.
- 9. Educate parents/ patients regarding disease.

#### MEDICAL MANAGEMENT

• First line drugs: -NSAIDs are usually given all types of JIA for 4 to 8 weeks.

 Commonly used are: Naproxen, Ibuprofen, diclofenac and Indomethacin. Usually well tolerated. Naproxen may be associated with scarring pseudoporphyria affecting face.

#### MEDICAL MANAGEMENT Cont...

Second line Drugs: `Disease Modifying Antirheumatic Drugs`-: include Methotrexate,
 Hydroxycholroquine, Sulphasalazine, Leflunamide,
 Cyclosporine, Cyclophosphamide and Azathioprine.

- Methotraxate:
  - a). Parenteral(o.5-1mg/kg/week): for polyarthritis, extended oligoarthritis and systemic onset JIA.

- b). Subcutaneous route
  (0.5-1mg/kg/week): is reserved for those who fails to tolerate parenteral therapy
- NOTE: immunization with live vaccine is contraindicated during treatment with MTX. It is better to vaccinate the child with varicella zoster vaccine 2 weeks prior to start therapy to those who are known to susceptible.

#### Leflunamide:

#### oral route:

#### Loading dose:

- < 20 Kg -> 100 mg once
- 20-40 kg -> 100 mg daily for 2 days
- > 40 kg -> as adults

#### Maintenance dose:

- < 20 Kg -> 10 mg every other day
- 20-40 kg → 10 mg daily or 20 mg every other day
- > 40 kg > as adults

- Hydrochloroquine:
- Oral route: 200 mg, 1-2 times daily
- Sulfasalazine:
- -Children 6 yrs and older
- Oral route: 40 60 mg/kg/day divided into 3 to 6 doses.

#### Management cont...

#### Use of corticosteroid in treatment of JIA:

- **# Oral corticosteroid-:**
- Oral use in large doses (up to 60 mg) prior to DMARDS is to avoid MAS in SOJIA.
- Use of oral corticosteroid in other subtypes is while awaiting the desired effect of

DMARDs therapy.

#### Management cont...

- # Intra-articular therapy: Intra-articular corticosteroid (IATH) is well established for mono or oligoarthritis.
- # Parenteral therapy: Parenteral high dose used intermittently in 'Pulse fashion' is a useful and very effective in SOJIA or polyarticular JIA.

### Management cont...

- **# New approach in treatment:** 
  - 1. Biological agent :
    - Eternacept, Infliximab Anti –TNF.
    - Anakinra-anti IL-I. Rituximab-B cell depletor.
    - 2. Early initiation of DMARDs and combination DMARDs
    - 3. Intravenous Ig.
    - 4. Autologous stem cell transplantation.

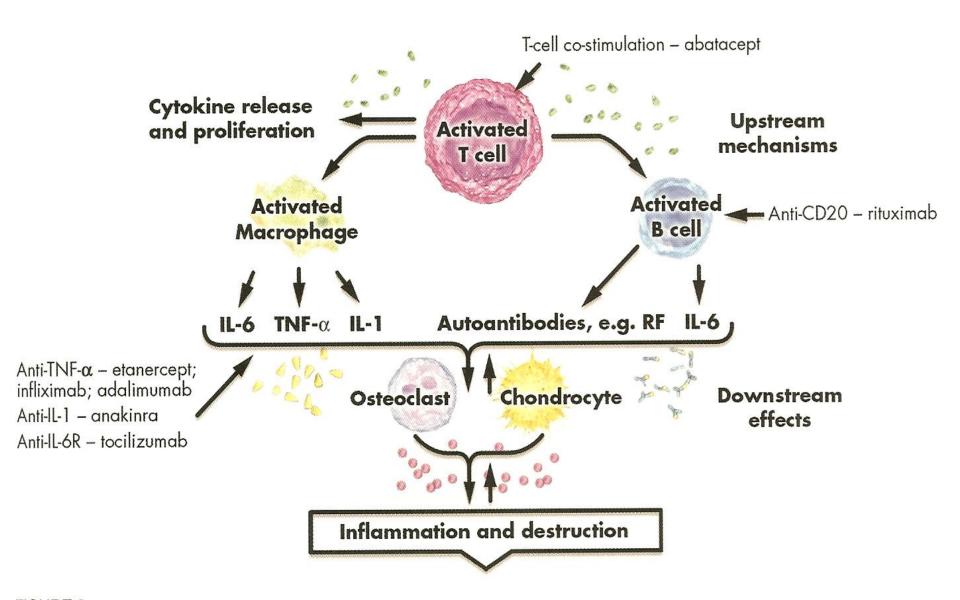
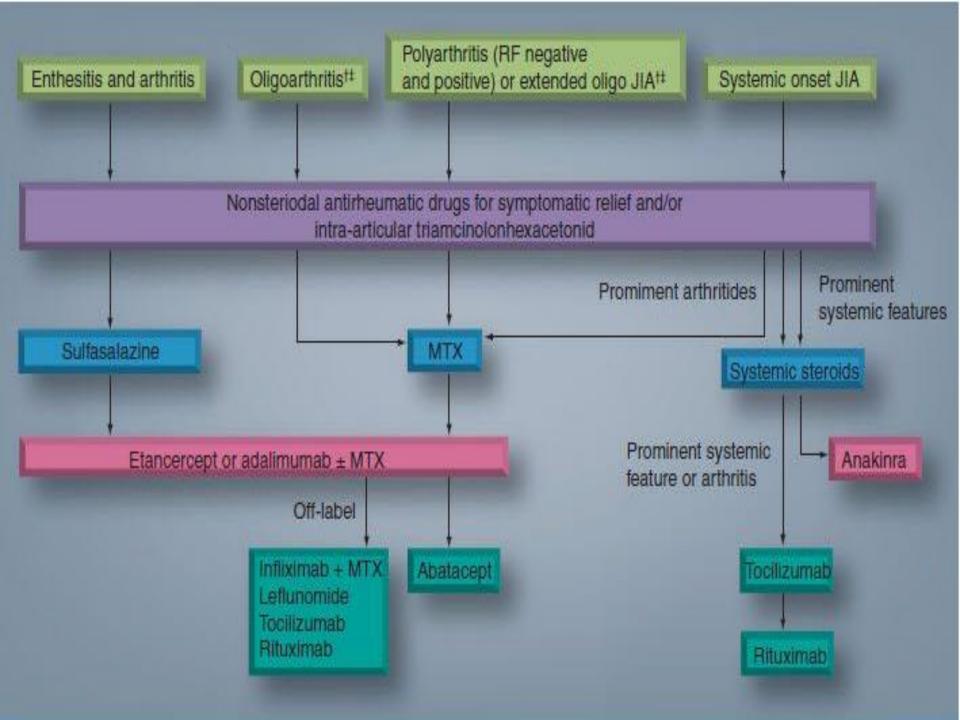


FIGURE 3. Current and emerging biologic therapies that selectively target rheumatoid arthritis immunopathology. IL, interleukin; TNF, tumor necrosis factor; RF, rheumatoid factor; IL-6R, interleukin-6 receptor.



#### Rx of Oligoarthritis JIA

- NSAID: for 4 to 6 weeks and/or intraarticular steroid.
- If improvement: observe, may be in remission or flare and become extended oligoarthritis or become polyarthritis or remain as persistant oligoarthritis
- # If extended oligoarthritis: Treat as polyarthritis JIA.

# Rx of Oligoarthritis JIA cont..

- # May progress to polyarthritis: Treat as Polyarthritis or
- # Remain as persistant oligoarthritis:
  Intermittent Intrarticular steroid
  and/ or Methotraxate or Sulfasalazine may
  be given.
- If no response: Antitumor necrosis factor medication may be given.

## Rx of Polyarthritis JIA

- NSAID for up to 6 weeks
- DMARDS

Similar to treatment of adult rheumatoid arthritis: Step up or step down strategy

Step up strategy:

Start with one drug in minimum dose (MTX OR Leflunamide) then gradually increase the dose until maximum dose if no response use combination therapy with Hydroquine or salazopyrine.

## Rx of Polyarthritis JIA cont...

Step down strategy:

We start with combination therapy

- Double combination therapy
- # MTX+ Hydroquine OR Salazopyrine
- #Leflunamide+ Hydroquine OR Salazopyrine
- # MTX+ Leflunamide
- Tripple combination therapy
- # MTX + Hydroquine + Salazopyrine
- # Leflunamide + Hydroquine + Salazopyrine

## Rx of Polyarthritis JIA cont...

- Steroid therapy as bridging medication or during serious disease flare
- If inadequate response: Antitumor necrosis factor medication, consider oral steroid

# Patient with Systemic onset JIA If systemic features are prominent

Steroid

We start with steroid therapy intially prior to DMARDS to avoid MAS

#Oral route: high dose of steriod up to 60 mg|day

#Parenteral route: consider I.V in 1st week or intermittent I.V pulses.

DMARDS

When fever and other systemic features improve we can use DMARDS as in polyarthritis subtype

NSAIDS: Full dose For 6-8 weeks

If arthritis is prominent

Treatment as in polyarticular

# Patient with Systemic onset JIA cont..

- If inadequate response: Steroid sparing medication such as **#I.V** immunoglobin #Antitumor necrosis factor #Interleukin 1 receptor antagonist #cyclosporin A #Interleukin 6 antibody.
- If inadequate response with very severe disease: consider autologous stem cell transplantation.

#### **Rx of complications**

- Uveitis: Topical steroid,
   Mydriatics,
   MTX(second line).
- If not controlled with oral MTX, then parentral MTX or MTX with ciclosporin should be considered.
- MAS: use of high dose corticosteroids and ciclosporin with vigorous treatment of sepsis.

#### CRITERIA OF REMISSION

- Morning stiffness<15 min</li>
- No fatigue
- No arthralgia.
- No joint tenderness or pain on motion.
- No soft tissue swelling in joints or tendon.
- ESR<30 in female and <20 mm in lst hr in male.
  - \*Five of the above criteria for at least 2 consecutive months.

- Bad prognostic factor:
- Rheumatoid factor positive.
- Systemic onset disease.
- More than a year without remission.
- Early development of nodules and erosions.
- Severe functional impairment.
- Involvement of hip joint.

#### **PROGNOSIS**

- JIA is some times self limiting, may persists months to yrs. Prognosis depends on subtypes of JIA.
- Oligoarticular JIA: is most benign.
- Seropositive poly articular and systemic onset JIA: have worst prognosis.
- SOJIA:
- 1. 30% complete remission by 5yrs.
- 2. 30% cyclic periods of resmission with exacerbation.
- 3. 30% still have disease.

Psoriatic arthritis: outcome is variable, with both oligo and polyarticular courses described.

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