# COMMON CAUSES OF ARTHRALGIAS

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## OBJECTIVES

Recognize common causes of joint pain

## TOPICS

- Growing Pains
- Benign Joint Hypermobility Syndrome
- Patellofemoral Pain syndrome or Chondromalacia Patellae syndrome
- Ehler's Danlos Syndrome
- Pain amplification syndrome
- Juvenile Idiopathic Arthritis

## GROWING PAINS

- Age of onset: 4 to 12 years
- Affects males and females equally
- 10-20% of school age children will complain of growing pains.

## SYMPTOMS OF GROWING PAINS

- Deep, aching, crampy pain
  - Usually bilateral
  - Thigh and calf
  - May be precipitated by excersice
- Evening to late night
- Never presents with morning pain
- Never presents with a limp

## WORK-UP FOR GROWING PAINS

- Laboratory studies should be normal
- X-ray studies should be normal

#### TREATMENT OF GROWING PAINS

#### Family education

Benign course with no significant sequlae

#### Supportive care

- Pain usually releived with massage
- If frequent attacks, tylenol/motrin prior to sleep
- Passive stretching

## BENIGN JOINT HYPERMOBILITY SYNDROME (BJHS)

- Frequency in general population
  - 8-20% in Caucasian
  - Higher frequency in Asians, African and Middle Eastern Descent
- Female to male ratio: 2:1
- Family history of hypermobility is common
  - Appears to have a strong genetic component with an autosomal dominant pattern
- Common age range: 3-10 years of age
  - Prevalence decreases with age

## SYMPTOMS OF BJHS

- Intermittent pain
  - May affect one or multiple joints
  - May be generalized or symmetric
- Mostly in late afternoon and night
  - Often wakes children up at night
  - Morning stiffness is uncommon
- Precipitated by certain activities
  - Physical activity or repetitive use
  - Jumping, running, gymnastics and ballet
- Mild joint effusions can be present

## INITIAL PROPOSED CRITERIA FOR BJHS

- Three of five are required to establish a diagnosis of hypermobility
  - Touch thumb to volar forearm
  - Hyperextend metacarpophalangeal joints so fingers parallel forearm or passive dorsiflexion of the MCP joint to 90 degrees
  - > 10 degree hyperextension of elbows
  - >10 degree hyperextension of knees
  - Touch palms to floor with knees straight

## BEIGHTON SCALE

- Requires four or more of nine tests (one point for each side)
  - Passive dorsiflexion of 5<sup>th</sup> MCP joint beyond 90 degrees with forearm flat
  - Passive apposition of the thumb to flexor aspect of the forearm
  - Passive hyperextension of the elbows >10 degrees
  - Passive hyperextension of the knees > 10 degrees
  - Forward flexion of the trunk, with knees straight, so that the palms rest easily on the floor

## REVISED BEIGHTON

#### • Requires:

- Two major criteria
- One major criteria and two minor criteria
- Four minor criteria

## CRITERIA FOR BJHS

#### Major

- Beighton score of at 4/9
- Arthralgias for >3 months in four or more joints

## CRITERIA CONTINUED

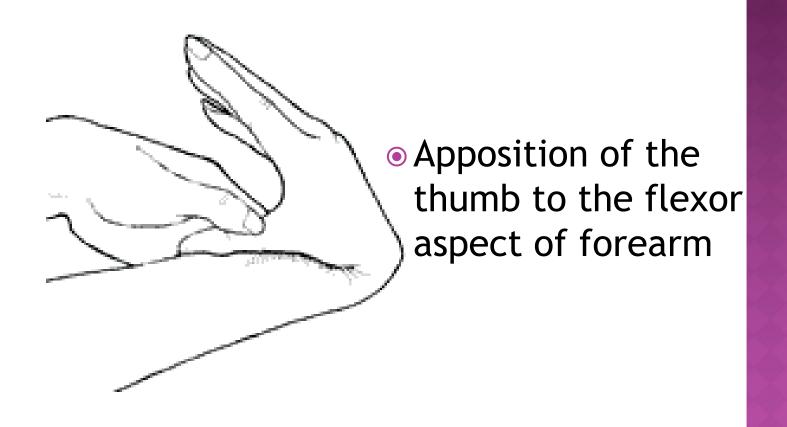
#### Minor Criteria

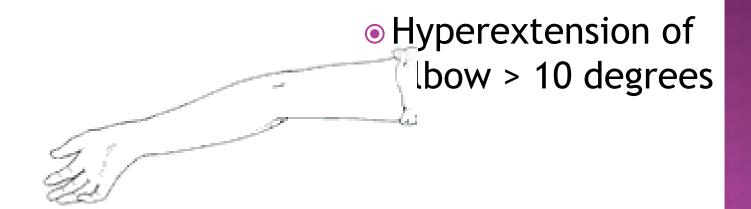
- Beighton score of less that 4/9
- Arthralgias > 3 months in 1-3 joints or back pain
- Dislocation/subluxation in more than one joint
- Soft-tissue rheumatism with three or more lesion
  - Bursitis or tenosynovitis
- Marfanoid habitus
- Abnormal skin
- Eyes signs
- Varicose veins or uterine/rectal prolapse

## OTHER NONCRITERIA FEATURES OF BJHS

- Put heel behind head
- Excessive internal rotation of hip
- Extensive ankle dorsiflexion
- Excessive eversion of the foot
- Passively touch elbows behind back

## DIAGRAMS OF BJHS







Hyperextension of Knee > 10 degrees



Ability to touch palms flat to floor with knees straight



Passive dorsiflexion of the MCP joint to 90 degrees

### TREATMENT

- Reassurance
  - Nonprogressive, noninflammatory condition
- For acute symptoms
  - NSAIDS such as motrin and acetaminophen
  - Message or warm baths
- Physical therapy for strengthening
- Avoidance of activities such as ballet or gymnastics

## **PROGNOSIS**

- Symptoms improve with age
  - Joints become less lax over time
- Increased frequency of:
  - Acute ligament and soft tissue injury
  - Overuse injury
  - Joint instability
  - Uterine and rectal prolapse
  - Possible increased risk of fracture or scoliosis

## PATELLOFEMORAL PAIN SYNDROME OR CHONDROMALACIA PATELLAE

- Age at onset: Adolescence to young adulthood
- Sex ratio: Girls>Boys
  - Most common in teenage girls

## PATELLOFEMORAL PAIN SYNDROME OR CHONDROMALACIA PATELLAE

- Insidious development of retropatellar knee pain
- Pain initially occurs with activity that stresses the quadriceps
  - Deep knee bends
  - Climbing or descending stairs
  - Prolonged sitting

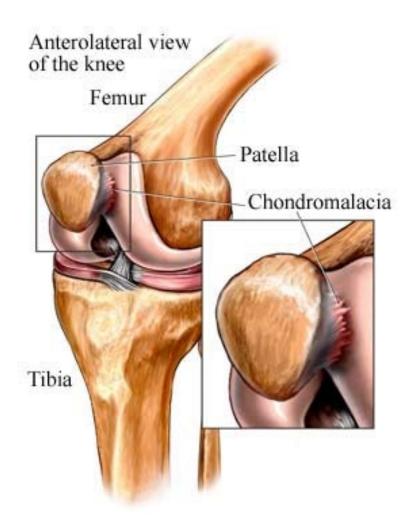
#### • Reproducing pain:

- Compression of the patella and by palpitation along its inferomedial edge
- Upward movement of the patella is restrained while the quads are being contracted

## DIAGNOSIS

- Physical exam
  - Knee flexion is accompanied by crepitus
- Laboratory studies are normal
- Plain x-rays films are usually normal
  - May have small effusion
- MRI of knee
  - Decreased signal intensity
  - Irregularity with focal thickening of the articular cartilage

## DIAGRAM



### TREATMENT

- Can be chronic and difficult to treat
- Initially activities that provoke pain should be avoided
  - Once pain improved may gradually increase activity
- PT to strengthen muscles around knee
- Braces with activity
  - Sports braces with hole at knee

## EXAMPLES OF SPORTS BRACES



## ARTHRALGIAS VS ARTHRITIS

#### Arthralgias

- Pain
- No inflammation

#### Arthritis

- Swelling or effusion
- Limitation on motion
- Pain on motion
- Tenderness on palpation
- Heat in a joint
- Occurs in one or more joints

## ARTHRALGIAS VS ARTHRITIS

#### Pain of Arthralgias

- Usually occurs as the day progresses
- Worsens with activity

#### Pain of arthritis

- Morning stiffness
- Worsens with prolonged sitting
- Improves with activity

#### JUVENILE IDIOPATHIC ARTHRITIS

- One of the most common rheumatic diseases of childhood
- Significant cause of childhood chronic illness
- Distinct entity from Adult Rheumatoid Arthritis

## ETIOLOGY AND PATHOGENESIS

- Genetic predisposition
  - Interactions of multiple genes
- Multiple disease onset triggers
  - Altered immunity, infections, trauma, stress
- Activation of autoreactive T cells
- B cells hyperactivity
- Release of inflammatory mediators
  - Cytokines (TNF, IL-1, IL-6, IL-18), prostaglandins
  - Damage cartilage and vessel wall

## ETIOLOGY AND PATHOGENESIS

- Hormonal factors
- Infection

## NOMENCLATURE

- JuvenileRheumatoidArthritis
  - Systemic
  - Polyarticular
  - Pauciarticular

- Juvenile Idiopathic Arthritis
  - Oligoarticular
    - Persistent
    - Extended
  - Polyarticular RF -ve
  - Polyarticular RF +ve
  - Systemic
  - Psoriatic arthritis
  - Enthesitis related
  - Other/Nondifferentiated

## ACR CRITERIA FOR CLASSIFICATION OF JIA

- Age of onset < 16 years</li>
- Arthritis in one or more joints
  - Swelling or effusion
  - Limitation of range of motion, tenderness or pain on motion, heat in a joints
- Duration longer than 6 weeks
- Exclusion of other forms of arthritis
  - Infection
  - Other rheumatic diseases
  - Malignancies
- Onset type defined by type of disease in 1st 6 months

## EHLER'S-DANLOS

- Rare genetic disorder
  - Defect in collagen synthesis
  - Fibronectin abnormality
  - Multiple defects involved
    - Fibrous proteins: COL1A1, COL1A2, COL3A1, COL5A1,
       COL5A2, and TNXB
    - Enzymes: ADAMTS2, PLOD1
- Affects 1 in 5000
- Males and females equally affected

## EHLERS-DANLOS

- Hypermobility
- Hyperelasticity of skin
- Vascular fragility and aneurysms (aorta, medium sized muscular arteries and cerebral arteries)
- Large eyes
- Lobeless ears

- Bilateral club feet
- Dislocated hips
- Peridontal disease
- Abnormal platelet aggregation

## EHLER'S-DANLOS CONTINUED

#### 13 major types

- Hypermobility
- Classical
- Vascular
- Kyphoscoliosis
- Arthrochalasis
- Dermatosparaxis

### HYPERMOBILITY TYPE

- Affects 1:10,000
- Symptoms
  - Loose, unstable joints
  - Soft, velvety skin
  - Chronic degenerative joint disease
  - Advanced premature osteoarthritis
  - Mitral valve prolapse

## PAIN AMPLIFICATION SYNDROME