

Rebalance<sup>MD</sup>

# PAEDIATRIC ORTHOPAEDICS

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PAEDIATRIC ORTHOPAEDIC SURGEON

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a place of mind  
THE UNIVERSITY OF BRITISH COLUMBIA

## DISCLOSURES

- I HAVE NO INDUSTRY CONFLICTS TO DECLARE
- I AM AN ORTHOPAEDIC SURGEON TRAINED IN PAEDIATRICS, NOT A PAEDIATRICIAN TRAINED IN ORTHOPAEDICS
- I AM A PARENT
- ALL IMAGES IN THIS PRESENTATION ARE FROM THE ROYAL CHILDREN'S HOSPITAL MELBOURNE ORTHOPAEDIC FACT SHEETS

## LEARNING OBJECTIVES

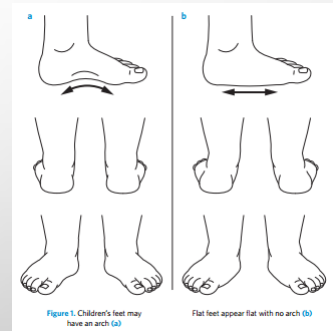
- DISCUSS NORMAL SO WE CAN IDENTIFY ABNORMAL
- COVER THE BASIC ORTHOPAEDIC CONCERNS IN YOUNG PATIENTS
  - DIAGNOSIS
  - BASIC TREATMENT
  - WHEN TO REFER
- LEAVE TIME FOR DISCUSSION

## DEFINING “NORMAL”

- ONE OF THE BIGGEST PARTS OF MY JOB IS SEPARATING THE “NORMAL” FROM THE PATHOLOGIC
  - ALMOST ALL CHILDREN MAKE THEIR WAY TO BEING A “NORMAL” ADULT
  - WHAT IS “ABNORMAL” IN ADULTS CAN BE “NORMAL” IN A CHILD
- PARENTAL CONCERN IS ONE OF THE MAIN REASONS KIDS SEE ME
- PAIN AND FUNCTIONAL LIMITATION ARE PROBABLY THE BEST MARKERS OF TRUE PATHOLOGY IN A CHILD

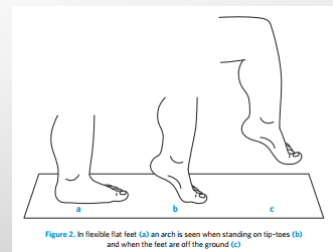
## FLAT FEET

- DEFINED AS LACKING THE LONGITUDINAL ARCH OF THE FOOT
- FLAT FEET ARE NORMAL IN ESSENTIALLY ALL INFANTS AND MANY YOUNG CHILDREN
  - IN INFANTS THE MEDIAL FAT PAD OBSCURES THE DEVELOPING ARCH
  - IN CHILDREN FLEXIBILITY CAN CREATE PHYSIOLOGIC FLEXIBLE FLAT FEET
  - MOST CHILDREN DEVELOP AN ARCH BY AROUND THE AGE OF 6
  - 1 IN 5 NEVER DEVELOP AN ARCH
    - VAST MAJORITY OF WHICH HAVE NO LONG TERM PROBLEMS



## FLAT FEET

- FLEXIBILITY OF THE FOOT IS THE MOST IMPORTANT FEATURE
  - PAINLESS FLEXIBLE FLAT FEET DO NOT REQUIRE TREATMENT (AT ANY AGE)
- ORTHOTICS AND EXERCISES DO NOT LEAD TO DEVELOPMENT OF AN ARCH
  - COMFORTABLE SHOES WHICH ACCOMMODATE THE FOOT SHAPE ARE MOST IMPORTANT
- ORTHOPAEDICS CAN HELP IF:
  - 1) PAINFUL FLAT FOOT
  - 2) FUNCTIONAL LIMITATION
  - 3) UNILATERAL FLAT FOOT
  - 4) RIGID/STIFF FLAT FOOT
    - LIKELY A TARSAL COALITION



# GAIT

- THERE IS NO NORMAL
- WE CALL THEM TODDLERS FOR A REASON
  - WIDER STANCE
  - RAPID CADENCE
  - SHORT STEPS
- IT TAKES UNTIL AROUND AGE 3 FOR KIDS TO DISPLAY MATURE WALKING PATTERNS
  - BY AGE 7, GAIT SHOULD MIMIC ADULT GAIT
- DURING YOUR CHILD'S FIRST FEW YEARS WALKING, THEY MAY HAVE OBVIOUS GAIT ABNORMALITIES
  - THE MOST COMMON TYPES OF GAIT ABNORMALITIES ARE INTOEING, OUTTOEING, LIMPING AND TOE WALKING
- MANY GAIT DISTURBANCES ARE COMMON AND CORRECT THEMSELVES ON THEIR OWN
- GAIT DISTURBANCES RARELY REQUIRE MEDICAL ASSISTANCE
- STILL BE THINKING ABOUT NEUROMUSCULAR DISORDERS AND DISLOCATED HIPs (AMOUNG OTHERS)



# GAIT CYCLE

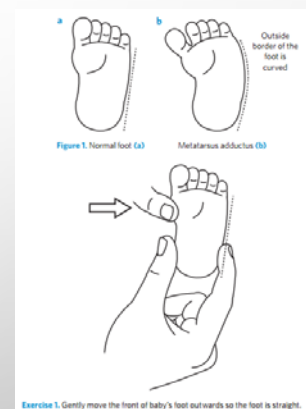
- STANCE PHASE (FOOT IS IN CONTACT WITH THE GROUND)
  - INITIAL DOUBLE-LIMB SUPPORT > SINGLE-LIMB STANCE > DOUBLE-LIMB SUPPORT
  - HEEL-STRIKE > FLAT FOOT > TOE OFF
- SWING PHASE (FOOT IS IN THE AIR)
- AGE CHANGES GAIT
  - WALKING VELOCITY, STEP LENGTH, AND THE DURATION OF THE SINGLE-LIMB STANCE INCREASE WITH AGE
  - NUMBER OF STEPS TAKEN PER MINUTE DECREASES
  - A MATURE GAIT PATTERN IS WELL ESTABLISHED BY THREE YEARS OF AGE, AND THE GAIT OF A SEVEN-YEAR-OLD CHILD CLOSELY APPROXIMATES THAT OF AN ADULT

# INTOEING

- DEFINED AS AN INTERNAL FOOT PROGRESSION ANGLE ON NORMAL GAIT
- OFTEN PRESENT BECAUSE OF CONCERNS REGARDING COSMESIS OF THE GAIT OR CLUMSINESS WITH RUNNING OR SPORTS
- EXTREMELY COMMON AND ALMOST ALWAYS OUTGROWN BEFORE AGE 10
- USUALLY COMES FROM ONE OF THREE CAUSES
  - 1) FOOT (METATARSUS ADDUCTUS)
  - 2) TIBIA (INTERNAL TIBIAL TORSION)
  - 3) HIP (FEMORAL ANTEVERSION)

# INTOEING

- METATARSUS ADDUCTUS
  - NORMAL LATERAL BORDER OF FOOT IS STRAIGHT
  - USUALLY PRESENT FROM BIRTH
  - OFTEN RESOLVES BY 2-3 YEARS OF AGE
  - OCCASIONALLY REQUIRES PT/STRETCHING
  - RARELY REQUIRES CASTING



# INTOEING

- INTERNAL TIBIAL TORSION
  - VERY COMMON IN INFANCY
  - USUALLY CORRECTS BY AROUND AGE 8
  - BEST CHECKED BY MEASUREMENT OF “THIGH-FOOT ANGLE”
    - PATIENT PRONE LOOKING DOWN ON FOOT IN RELATION TO THIGH
  - NO BRACING, FOOTWEAR OR ORTHOTICS MAKE ANY DIFFERENCE

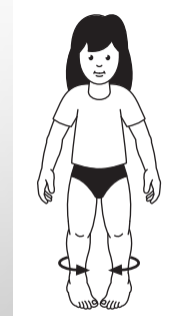


Figure 2. Internal tibial torsion — when the lower leg turns inwards between the knee and ankle

# INTOEING

- FEMORAL ANTEVERSION
  - NORMAL IN YOUNG CHILDREN
  - USUALLY RESOLVES BY AROUND 10 YEARS OF AGE
  - “W-SITTERS”
    - CHILDREN SIT WITH KNEES FORWARD AND BOTTOMS BETWEEN THEIR FEET
    - NO EVIDENCE THIS IS HARMFUL, BUT OFTEN SUGGEST TO AVOID IT
  - NO BRACING, FOOTWEAR OR ORTHOTICS MAKE ANY DIFFERENCE

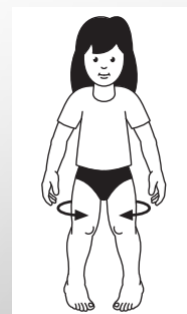


Figure 3. Internal femoral torsion — when the thigh bone turns inwards between the hip and the knee

## INTOEING

- ORTHOPAEDICS CAN HELP IF:
  - 1) UNILATERAL INTOEING
  - 2) SEVERE INTOEING
  - 3) FAILURE TO IMPROVE ON EXPECTED TIME COURSE
  - 4) SCHOOL AGE CHILD WITH TRIPPING AND DECLINED PARTICIPATION AS A RESULT
  - 5) STIFF METATARSUS ADDUCTUS

## TOE WALKING

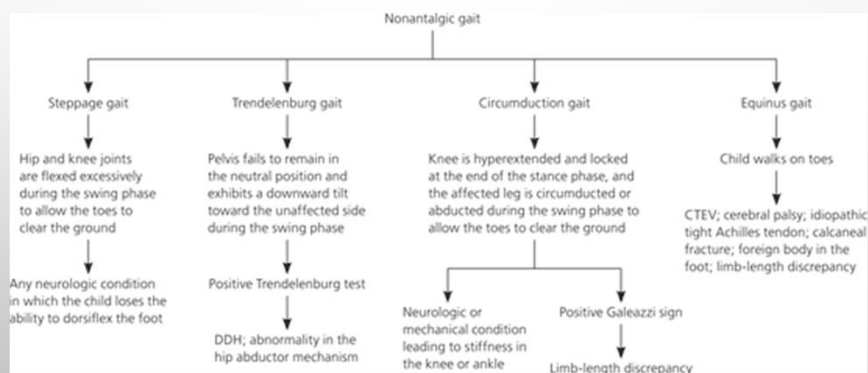
- MORE COMMON IN AUSTISM SPECTRUM
- DON'T WORRY AS MUCH THOSE WHO HAVE DONE IT FOREVER, DO WORRY ABOUT NEW ONSET TOE WALKERS
- STRETCHING, REMINDING AND HIGH-TOPS CAN HELP BREAK THE HABIT
- NEEDS EVALUATION IF:
  - PERSISTS BEYOND AGE 3
  - NEUROLOGIC FINDINGS
  - UNILATERAL
  - PAIN



## THE LIMPING CHILD

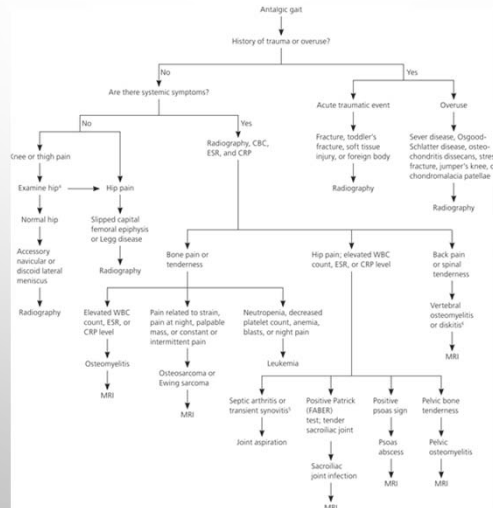
- HAVE A HIGH LEVEL OF SUSPICION, BUT APPRECIATE MOST DIAGNOSIS ARE BENIGN AND SELF-LIMITING
- THIS CAN BE A SIGN OF MSK OR NON-MSK RELATED DISEASE
  - GOOD PHYSICAL EXAM OF THE WHOLE CHILD IS REQUIRED
- HISTORY, PHYSICAL EXAM, LABS, IMAGING
  - JUST LIKE EVERY OTHER PROBLEM

## THE LIMPING CHILD



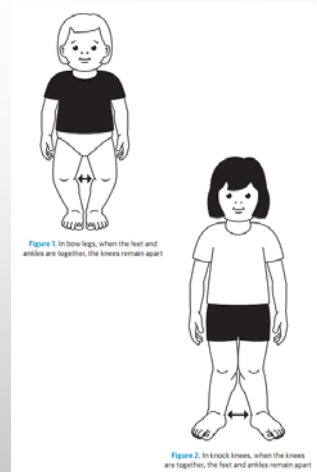


## THE LIMPING CHILD



## BOW LEGS AND KNOCK KNEES

- **GENU VARUM (BOW LEGS)**
  - NORMAL FOR INFANTS AND TODDLERS
  - USUALLY CORRECTS BY AROUND AGE 3
  - MORE OBVIOUS IN STANDING/WALKING CHILD
- **GENU VALGUM (KNOCK KNEES)**
  - NORMAL BETWEEN AGES 3-6 YEARS
  - USUALLY CORRECTS BY AROUND AGE 8
  - MUST ASSESS WITH PATELLAE FACING FORWARD



## BOW LEGS AND KNOCK KNEES

- ORTHOPAEDICS CAN HELP IF:
  - 1) DEFORMITY IS SEVERE
  - 2) DEFORMITY TYPE IS OUTSIDE THE NORMAL GROWTH TRAJECTORY AND AGE RANGE
  - 3) UNILATERAL DEFORMITY
  - 4) ASSOCIATED WITH PAIN OR LIMP
  - 5) SHORT STATURE

## GROWTH

- GROWTH IS NOT A BENIGN PROCESS
- I DESCRIBE TO PARENTS AS “THE SKELETON GROWS AND EVERYTHING ELSE JUST HAS TO STRETCH TO FIT”
  - PRE-EXISTING TIGHTNESS GETS WORSE
    - CEREBRAL PALSY
    - LOW BACK PAIN IN ADOLESCENTS
  - NEW TIGHTNESS DEVELOPS
    - ENTHESOPATHIES
    - SCOLIOSIS
- ALIGNMENT CHANGES DRASTICALLY THROUGH GROWTH

## GROWING PAINS

- DIAGNOSIS OF EXCLUSION
- OCCUR IN 15-30% OF CHILDREN
- PAIN IS CHARACTERISTIC
  - GENERALLY ACHE IN LARGE MUSCLE GROUPS (QUADS, CALF, HAMS)
  - TENDS TO BE EVENING OR NIGHT PAIN (CAN WAKE KIDS FROM SLEEP)
  - TENDS TO BE WORSE AFTER AN ACTIVE DAY
- TREATMENT IS SYMPTOMATIC RELIEF
  - MASSAGE
  - HEAT
  - TYLENOL/IBUPROFEN PRN
- STRETCHING IS PREVENTATIVE
- CONCERNS = ANY SIGN OF ANY OTHER DIAGNOSIS
  - SWELLING, HEAT, ATYPICAL PAIN

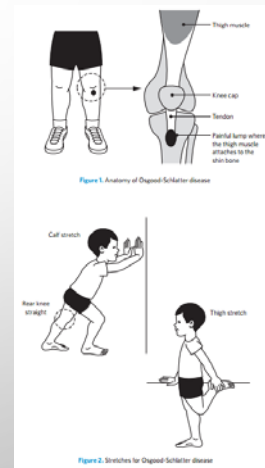
## ENTHESOPATHIES

- INFLAMMATION WHERE A TENDON INSERTS ON BONE
- MANY COMMON DISORDERS ARE ENTHESOPATHIES
  - IN CHILDREN THEY TEND TO BE APOPHYSITIS
    - TENDON INSERTING ADJACENT TO OR ON A GROWTH PLATE
  - THEY ALL HAVE NAMES, BUT NO ONE CAN REMEMBER MOST OF THEM
- ANOTHER GROWTH RELATED PHENOMENON



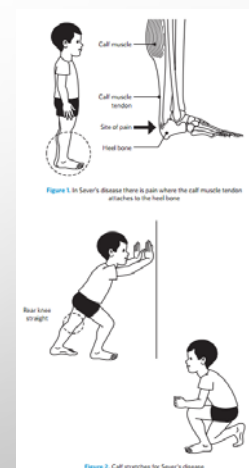
# OSGOOD-SCHLATTER DISEASE

- TIBIAL TUBERCLE APOPHYSITIS
- MORE COMMON IN BOYS
  - PATELOFEMORAL SYNDROME MORE COMMON IN GIRLS
- TENDS TO BE IN ACTIVE, RUNNING/JUMPING ATHLETES
- PAIN TENDS TO BE PRESENT DURING OR IMMEDIATELY AFTER ACTIVITY
- TENDER DIRECTLY OVER THE LUMP OF TIBIAL TUBERCLE
  - OFTEN SWELLING/ENLARGEMENT OF TUBERCLE
- SYMPTOM MANAGEMENT
  - REST, ICE, ANALGESIA
  - STRETCHING!!
  - TAPING/BRACING
- PAIN GOES AWAY BUT LUMP OFTEN STAYS



# SEVER'S DISEASE

- CALCANEAL APOPHYSITIS
- RESPONSIBLE FOR ALMOST ALL HEEL PAIN IN CHILDREN
- SAME RISK FACTORS AND PAIN PROFILE AS OSGOOD-SCHLATTER
- CONTINUING SPORT IN THESE CONDITIONS IS NOT HARMFUL BUT MAY EXACERBATE THE PAIN
- SYMPTOM MANAGEMENT
  - REST, ICE, ANALGESIA
  - STRETCHING!!
  - GEL HEEL CUPS IN ALL SHOES
- USUALLY GOES AWAY WITHIN 6-12 MONTHS OF ONSET WITH STRETCHING



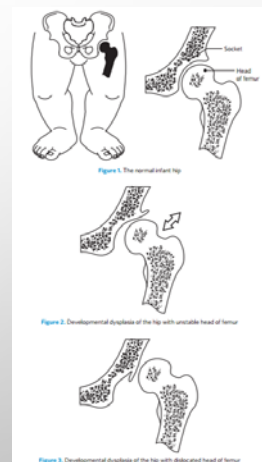
## CURLY TOES

- YES - THAT IS ACTUALLY ITS MEDICAL NAME
- THE OFFENDING TOES ARE THE ONES CURLED UNDER THE NEIGHBOUR
  - OVERRIDING SECOND TOE IS AN ENTITY
- MANY CORRECT ON THEIR OWN
  - I SAY WAIT UNTIL AT LEAST 3 YEARS OLD IN MOST CASES
- SURGICAL INDICATIONS
  - PAIN
  - BLISTERS OR SKIN CHANGES
  - NAIL PROBLEMS



## DEVELOPMENTAL DYSPLASIA OF HIP

- THE HIP IS A BALL-IN-SOCKET JOINT
- NORMAL INFANT HIP IS NOT MATURE AT BIRTH, BUT DEVELOPS INTO A STABLE JOINT WITH TIME
  - FAILURE OF THIS PROCESS IS DDH
- DDH IS A SPECTRUM OF DISEASE
  - ACETABULAR DYSPLASIA (SHALLOW SOCKET)
  - UNSTABLE HIP (BARLOW POSITIVE)
  - DISLOCATED HIP (ORTOLANI POSITIVE)



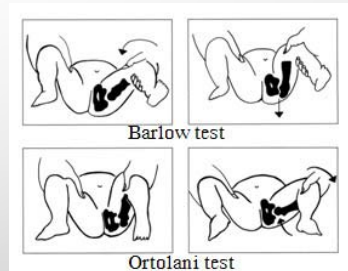
## DEVELOPMENTAL DYSPLASIA OF HIP

- OCCURS IN 1/1000 LIVE BIRTHS
- RISK FACTORS
  - FIRST BORN
  - FEMALE
  - FAMILY HISTORY
  - FRANK BREECH
  - CULTURAL (FINNISH, FIRST NATIONS)
  - TWINS (FRATERNITY)
  - HIGH BIRTH WEIGHT (FAT)



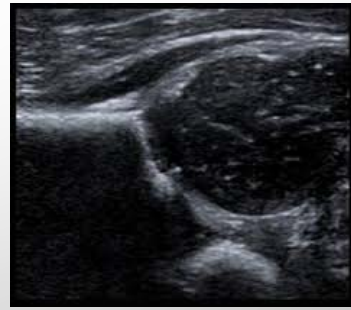
## DEVELOPMENTAL DYSPLASIA OF HIP

- DDH IS OFTEN PICKED UP AT BIRTH BUT CAN DEVELOP IN THE WEEKS OR MONTHS AFTER BIRTH
  - ROUTINE EXAM ON BABY CHECKS IS IMPORTANT
- DDH IS NOT PAINFUL – PAIN IS NOT A SIGN OF DDH
- ORTOLANI/BARLOW SIGNS USEFUL EARLY, BUT MORE DIFFICULT AS CHILD AGES
- LACK OF ABDUCTION IS MOST USEFUL SIGN
- ASYMMETRIC THIGH CREASES ONLY USEFUL IN NEWBORN PERIOD
- CLICKS ARE ALMOST ALWAYS BENIGN



## DEVELOPMENTAL DYSPLASIA OF HIP

- ULTRASOUND SCREENING
  - 2 OR MORE RISK FACTORS
    - FIRST BORN, FEMALE, BREECH, FAMILY HISTORY
  - ANY ABNORMAL EXAM
  - IF YOU ARE UNSURE FOR ANY REASON



## DEVELOPMENTAL DYSPLASIA OF HIP

- EARLY DETECTION AND TREATMENT IS CRITICAL
- IT IS A RELATIVELY EASY PROBLEM UNDER THE AGE OF 6 MONTHS AND GETS EXPONENTIALLY MORE DIFFICULT TO TREAT THEREAFTER
- LATE DIAGNOSIS OFTEN RESULTS IN LARGE SURGICAL TREATMENT WITH INFERIOR OUTCOMES



## DEVELOPMENTAL DYSPLASIA OF HIP

- HEALTHY HIP POSITIONING

- "THE FROG POSITION"
- KNEES UP AND OUT
- SAFE SWADDLING
- CARRIERS

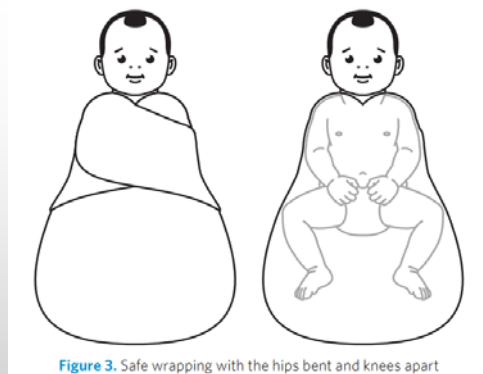


Figure 3. Safe wrapping with the hips bent and knees apart

## CLUBFOOT

- TALIPES EQUINOVARUS

- FOOT TURNED IN AND DOWN

- POSITIONAL VS. CONGENITAL

- FLEXIBLE VS RIGID

- AFFECTS 1/1000 LIVE BIRTHS

- WITH ONE AFFECTED CHILD, RISK IS 1/30 OF SUBSEQUENT CHILD BEING AFFECTED





# CLUBFOOT

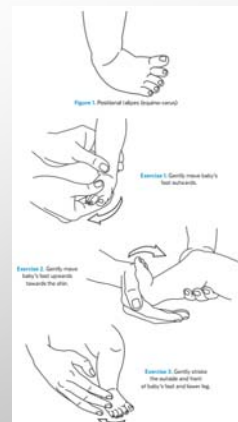
- TREATED WITH SERIAL CASTING
  - PONSETI TECHNIQUE
    - WEEKLY CAST CHANGES
    - LONG LEG CASTS
  - OFTEN ACCOMPANIED WITH ACHILLES TENOTOMY AT END OF TREATMENT
  - DENNIS-BROWNE BOOTS AND BARS FOLLOW CASTING
    - FULL TIME FOR 3 MONTHS
    - NIGHT WEAR UNTIL AGE 3-4 YEARS



Figure 3. Foot abduction brace

# CLUBFOOT

- POSITIONAL TALIPES
  - ANY FOOT THAT CAN BE PUT IN NORMAL POSITION IS NOT A TRUE CLUBFOOT
  - HIGH CORRELATION TO DDH
  - TREATED WITH STRETCHING
  - CASTING RARELY REQUIRED





THANK YOU



# *Perinatal Orthopaedics*

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Medical Director, Rebalance<sup>MD</sup>

## Outline

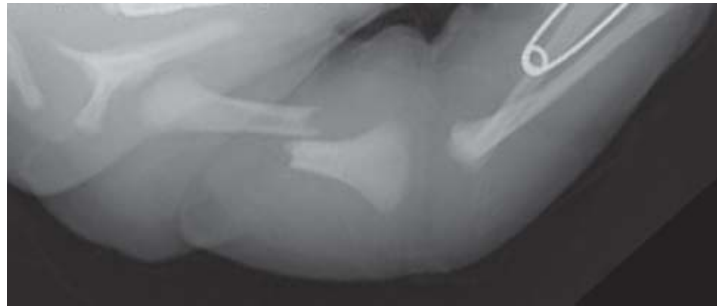
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- **Neonatal fractures**
- **Positional foot deformities**
- **Hip pathology**
- **Discussion**

# Neonatal Fractures

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- **6-8/1000 Live births**
- **Perinatal trauma**
- **Brittle bone diseases**



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# Neonatal Fractures

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## • Causes

- Shoulder dystocia
- Instrument deliveries
- Versions and extractions

## • Risk Factors

- Macrosomia (>4500g)
- Maternal obesity
- Abnormal presentation
- Prima gravida
- Cephalopelvic disproportion, small maternal stature, maternal pelvic anomalies
- Prolonged or rapid labor
- Oligohydramnios



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# Neonatal Fractures

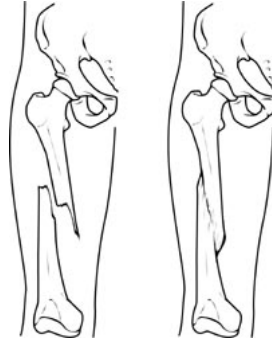
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- **Common injuries**

- Clavicle Fracture (most common)
- Humerus fracture
- Femoral fracture
- Brachial plexus injury

- **Findings**

- Loss of spontaneous movement of limb
- Swelling
- Pain on passive movement



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# Neonatal Fractures

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- **Management**

- Upper limb
  - Immobilize by pinning arm to body
  - Vietnam splint
- Lower limb
  - Pavlik harness



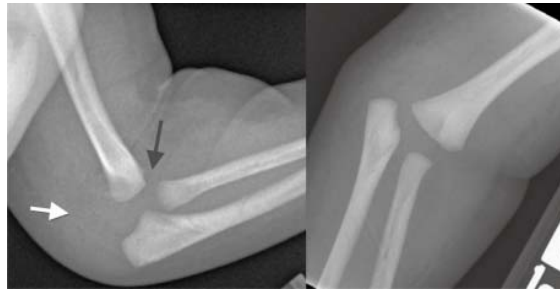
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# Neonatal Fractures

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- **Special cases**

- Epiphyseal dissociation
  - No fracture on XR
  - Something just isn't right to you



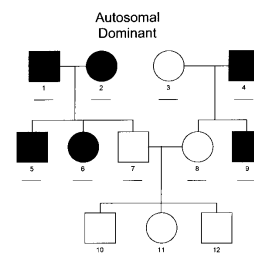
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# Brittle Bone Diseases

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- **Suspicious Findings**

- Translucent skin
- Bowing deformities
- Blue sclera
  - Can be normal – persistence is not
- Family history



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## Brittle Bone Diseases

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- **Metabolic bone disease of prematurity**
- **Osteogenesis imperfecta**
- **Rickets**
- **Bruck Syndrome**
- **Cole-Carpenter Syndrome**
- **Familial Osteoporosis**
- **Fibrous Dysplasia/Mc-Cune Albright Syndrome**
- **Hypophosphatasia**
- **Congenital insensitivity to pain**

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## Brittle Bone Diseases

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# Brittle Bone Diseases

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- **Metabolic bone disease of prematurity**

- < 28 weeks
- Late establishment of enteral feeds
- Chronic lung disease
- Jaundice (conjugated hyperbilirubinemia)
- Furosemide diuresis (> 2 weeks)
  - Urinary loss of Ca -> Bone remodelling
- Self-limiting disease (2 years)

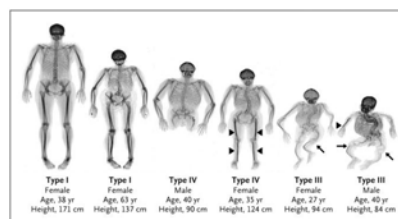
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# Brittle Bone Diseases

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- **Osteogenesis imperfecta**

- 1 in 15,000
- Type I collagen deficiency
- 4 types
  - I = mild
  - II = “lethal”
  - III = severe
  - IV = moderate
- Family history



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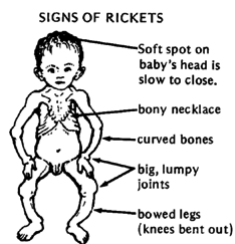


# Brittle Bone Diseases

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- **Rickets**

- Multiple types
  - Hypophosphatemic
  - Vitamin D deficiency
- Rising incidence in NA
- Breast fed without supplementation



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# Positional Foot Deformities

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- **Clubfoot**

- Congenital Talipes Equinovarus
- Positional clubfoot

- **Calcaneovalgus**

- Congenital vertical/oblique talus

- **Metatarsus adductus**



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# Congenital Clubfoot

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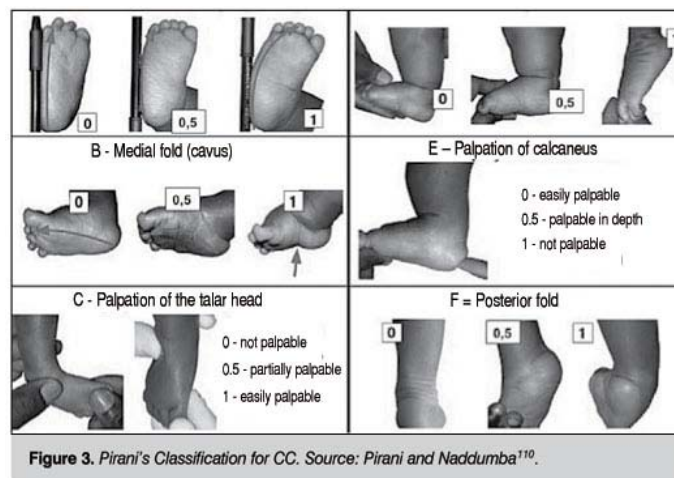
- **1/1000 live births**
  - 50% bilateral
- **Very Specific Deformity**
  - CAVE
    - Cavus (plantar flexed 1<sup>st</sup> ray)
    - Adductus (Metatarsal)
    - Varus (Hindfoot)
    - Equinus
- **Spectrum of severity**
  - Pirani score



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# Congenital Clubfoot

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# Congenital Clubfoot

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- **Treatment is by Ponseti serial casting**

- Start within first 2 weeks of life
- Serial casting to correct deformity
- Weekly cast changes (usually 5-8 casts)
- Achilles tenotomy

- **Boots and bars**

- 3 months full time
- Sleep time wear until at least 3<sup>rd</sup> birthday

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# Congenital Clubfoot

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## Clubfoot treatment over 4 – 6 weeks



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## Congenital Clubfoot

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## Positional Clubfoot

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- **Flexible deformity**
  - Foot held in same position as congenital clubfoot
- **Treatment is physiotherapy**
  - Occasionally do 1-2 casts to speed it up
  - No boots and bars or maintenance



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## Atypical Clubfoot

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- **Also known as syndromic clubfoot**

- Arthrogryposis
- Tibial Hemimelia
- Mobius Syndrome
- Larsen Syndrome
- Pierre Robin Sequence



- **Resistant to usual casting techniques**

- **Usually severe appearing**

- May be associated with other MSK deformities

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## Calcaneovalgus Foot

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- **Vast majority are normal**

- Intrauterine positioning
- Most resolve without treatment
- If no improvement by 3 months, need further assessment

- **Congenital vertical talus**

- Rocker bottom foot
- Often associated with neuromuscular abnormality



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# Metatarsus Adductus

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- **Most common foot deformity of infancy**

- Incidence as high as 12%
- Flexible require no treatment
- Resolve spontaneously
- Passive stretching recommended
- Casting may be appropriate if not resolved by 6-12 months



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# Hip Dysplasia

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- **Name changed from Congenital Dislocation of the Hip to Developmental Dysplasia of the Hip**

- Spectrum of disease
- Acetabular dysplasia
- Instability
- Dislocation

- **It is a failure of the normal development of the hip**

- Acetabulum and femoral head come from a single block of cartilage
- Acetabulum responds to pressure of femoral head on it to form a socket

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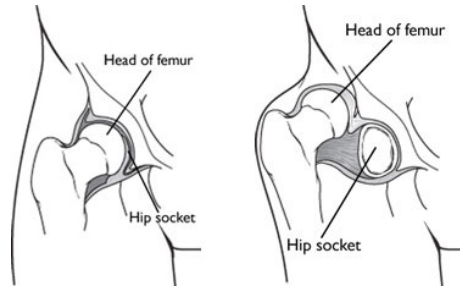
# Hip Dysplasia

Rebalance<sup>MD</sup>

- **Incidence 1-2%**

- **Cause**

- We have no idea...
- Genetic?
  - Family history
  - Ethnic variations
- “Packaging Syndrome”
  - First born
  - Oligohydramnios
  - Associated abnormalities (metatarsus adductus, torticollis...)
  - Left hip (forced adducted against spine in left occiput anterior)



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# Hip Dysplasia

Rebalance<sup>MD</sup>

- **Risk Factors**

- First Born
- Females
- Frank breech
- First Nations/Finnish
- Family history
- Foot deformity
- Facing one direction (torticollis)



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# Hip Dysplasia

Rebalance<sup>MD</sup>

- **Physical exam**

- Leg length difference
  - Galeazzi Sign



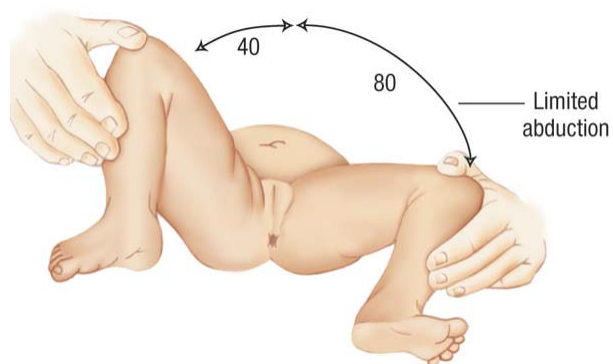
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# Hip Dysplasia

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- **Physical exam**

- Leg length difference
  - Galeazzi Sign
- Limitation in abduction



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# Hip Dysplasia

Rebalance<sup>MD</sup>

- **Physical exam**

- Leg length difference
  - Galeazzi Sign
- Limitation in abduction
- Asymmetric thigh folds
  - Only helpful in the NEWBORN period



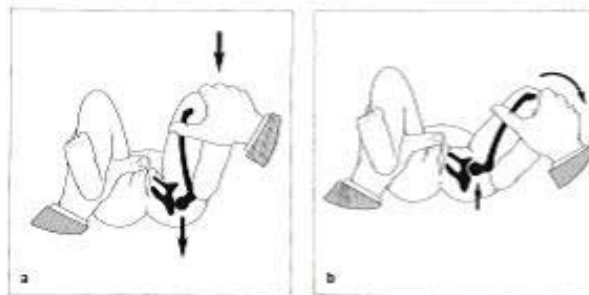
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# Hip Dysplasia

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- **Physical exam**

- Ortolani
  - Reducing a dislocated hip
  - Hip must be QUT to be QRTOLANI
- Barlow
  - Dislocating an unstable hip



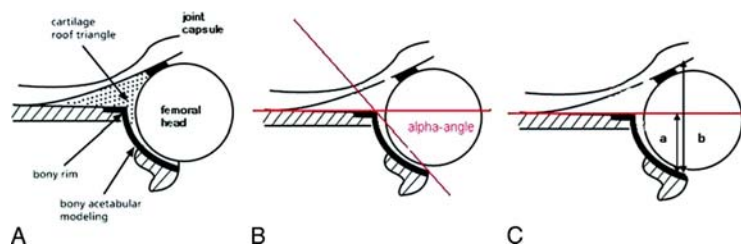
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# Hip Dysplasia

Rebalance<sup>MD</sup>

- **Ultrasound**

- We screen patients with risk factors
- We are the only screening program in Canada
- Literature does not support cost effectiveness of screening programs in DDH
- Early intervention is BETTER treatment

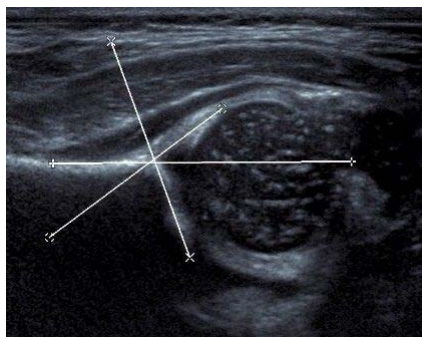


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# Hip Dysplasia

Rebalance<sup>MD</sup>

- **Ultrasound**



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# Hip Dysplasia

Rebalance<sup>MD</sup>

## • Treatment

- The whole krux of treatment is get the ball in to the socket as soon as possible to encourage the normal development of the hip
- Essentially replicating the Ortolani manoeuvre
  - Flexion (100°)
  - Abduction



Dennis Browne Bar



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# Hip Dysplasia

Rebalance<sup>MD</sup>

## • Treatment

- The whole krux of treatment is get the ball in to the socket as soon as possible to encourage the normal development of the hip
- Essentially replicating the Ortolani manoeuvre
  - Flexion (100°)
  - Abduction



Pavlik Harness



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# Hip Dysplasia

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- **Treatment of reduced but dysplastic hip**

- Hip development just needs to be encouraged
- Hold in abduction and flexion until normal parameter
- Usually 6 weeks full time wear + 6 weeks night time wear
- Older the patient starts, longer bracing goes on



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# Hip Dysplasia

Rebalance<sup>MD</sup>

- **Treatment of dislocated hip**

- Need to get reduced (Ortolani)
- Brace with ultrasounds every week to check progress
  - Surprising number actually reduce!
- Failure at 3 weeks = abandon bracing
- Surgical care



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# Hip Dysplasia

Rebalance<sup>MD</sup>

- **Treatment of dislocated hip (Surgical)**

- Attempt closed reduction and spica casting in OR
- Can release adductor tendons to improve ``zone of safety``
- Cast change at 6 weeks
  - 2 casts = total 12 weeks in cast
  - +/- bracing post casting
- Failed closed reduction = open reduction
  - Clear blockages to reduction
  - Capsuloraphy
- Older children (~18 months) require corrective osteotomies in conjunction



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# Hip Dysplasia

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## Congenital Dislocation of Knee

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## Arthrogryposis

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# Radial Club Hand

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# Thank You

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