



Paediatric Orthopedics: When does Physiology become Pathology?

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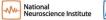


















SCOPE

FLATFEET

When to Worry?

LOWER LIMB MALALIGNMENT

INTOEING

ACHES AND PAINS

ABNORMAL GAIT/LIMPING CHILD



FLATFEET





















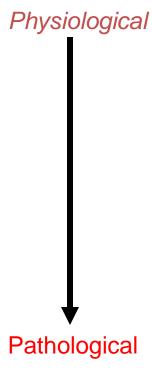




FLATFEET

Two diagnostic groups

- Dynamic
 - Due to generalised ligamentous laxity
 - Due to tight TA
- Fixed Flatfeet
 - Neuromuscular e.g. CP
 - Idiopathic
 - Tarsal coalition with/without peroneal spasm





FLATFEET Dynamic







FLATFEET Management

Dynamic with no tight TA

Watch and Masterly Inactivity

Dynamic with tight TA

Stretching and Medial Arch Support

Fixed

Treat underlying problem - usually needs surgery.



Good Shoeing

- Heel in line with back of leg
 - Firm heel counter
- Arch lifted off and not bearing weight
 - Built in soft arch supports
- No pressure on weight bearing area
 - Good padding of the sole with good flexibility



Arch Supports

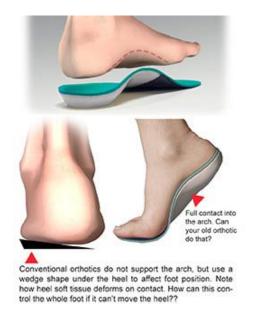
- Most effectively used when the heel is aligned in a good pair of shoes
- DOES NOT create an arch- only supports it and arch will collapse again once removed
- No need for custom made ones as child is still growing.



Arch Supports

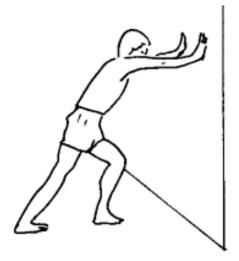


Stepping off the orthotic





Heel Cord Stretching



A. Sit with your legs straight in front of you. Wrap a towel around the ball of your foot and hold the loose ends in each hand. Pull the ends of the towel toward you until you feel a gentle stretch in the back of your calf.



B. Stand on an incline box with the higher edge toward the wall. With your heel on the floor and knee straight, lean toward the wall until you feel a gentle stretch in your calf.



C. Stand about an arm's length from a wall with one foot in front of the other. Bend the front knee and put your hands on the wall.

Lean forward until you feel a gentle stretch in the calf of your back leg.

Next, bring your back leg closer to the wall and bend both knees. Lean forward until you feel a gentle stretch in the heel cord of your back leg.



FLATFEET

When to Worry?











FLATFEET

- Dynamic
 - Due to generalised ligamentous laxity
 - Due to tight TA
- Fixed Flatfeet
 - Due to pathological ligament laxity.
 - Due to pathological tight muscles /contracture
 - Talar anatomy
 - oblique talus
 - Vertical talus
 - Tarsal coalition with/without peroneal spasm

TARSAL COALITION Calcaneo Navicular 9 to 12 years

oblique xray



Talo - calcaneum 12 to 15 years CT



50% bilateral

Treat if < 30% involved- treat with soft tissue interposition

- fat
- ext digitorum brevis

Spasmodic peroneal flatfoot.



LOWER LIMB MALALIGNMENT





























Normal alignment by age



BOWLEGGEDNESS

Tibial Intorsion and Tibia Vara



Corrects by age 3



Tibial Intorsion



When to worry?



5 YEAR OLD GIRL





KNOCK KNEES

Genu Valgum

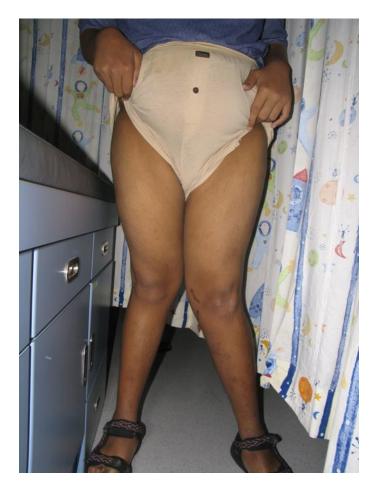
- Most commonly physiological
- Corrects to adult alignment by school going age
- Monitored with intermalleolar distance





When to worry?









INTOEING











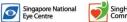
















INTOEING Physiological

Birth to walking

Metatarsus Adductus

Toddler to 3 yrs

Tibial Intorsion

School going to 10 yrs

Femoral Anteversion



INTOEING Metatarsus Adductus

Management

- Serial casting if diagnosed early
- Reverse Last shoes in older children
- Stretching will suffice if mild

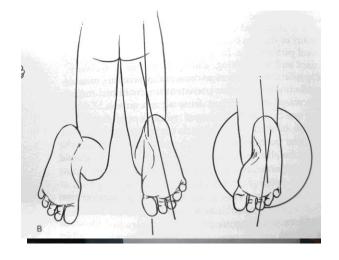




INTOEING Tibial Intorsion

Measured as

- Thigh foot Angle
- Trans Malleolar Axis





INTOEING Tibial Intorsion

Management

- Watch if not associated any other deformity
- Physiological and self limiting.



History of W-sitting





- History of W-sitting
- Increased Internal Rotation





- History of W-sitting
- Increased Internal Rotation

Reduced External Rotation





Management

- Cross legged sitting
- Reassurance will out turn by 10 yrs of age
- Consider behavior modification





When to worry?



ACHES AND PAINS









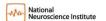


















ACHES AND PAINS

Growing Pains

- Cause unknown
- From age 3 or so.
- Pain mainly at night. Episodes for years.
- Clinic examination is normal.
- Massage and tender loving care usually will suffice.
- Self limiting gets better after school going



ACHES AND PAINS

Growing Pains When to worry?

- Pain with limping
- Pain extends into the day and affect play.
- Beyond school going
- Does not get better with simple remedies



ABNORMAL GAIT/LIMPING











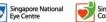
















ABNORMAL GAIT/LIMPING

- Gait is a function of brain development
- Brain development is mostly done when child achieve ability to converse in full sentences – around 3.
- Until then gait is still not stable or "adult pattern"



LIMPING CHILD

When to worry?



LIMPING CHILD

When to worry?

Before age of 3

- Biggest concern is an undiagnosed dislocated hip (DDH)
 - Significant limb length discrepancy
 - Asymmetrical thigh crease
 - Hip clicks
 - The hip "does not feel" the same on both sides







Limping 2½ yr old boy



LIMPING CHILD

When to worry?

3 to school going

School going through adolescents

Adolescents

To exclude
Trauma,
Infection and
Tumour
for All groups



LIMPING CHILD When to worry?

3 to school going

School going through adolescents

Adolescents

- Transient synovitis
 - Inability to bear weight
 - Associated with excessive running too much
 - May be related to upper respiratory infection
 - Gets better with rest



LIMPING CHILD When to worry?

3 to school going

School going through adolescents

Adolescents

Perthes

- Loss of blood supply to the hip
- Unknown cause
- Can cause early degenerative disease















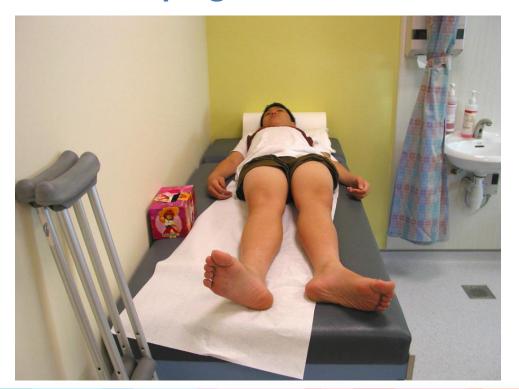


13 year old Limping with Left knee pain





13 year old limping adolescent









LIMPING CHILD When to worry?

3 to school going

School going through adolescents

Adolescents

- Slipped Upper Femoral Epiphysis
 - Thought to be due to excessive weight on a "weak" hip
 - Diagnosis delayed because they can present with knee pain
 - Always need surgery to stop progression



CONCLUSION

Most common conditions are thankfully physiological in nature.

It is however important to recognise a condition than is beyond physiological as most time than not early treatment will resolve the condition.





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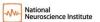






















A Primary Physicians' guide to Common Congenital and Developmental Conditions in Paediatrics Orthopaedics

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SCOPE

DDH

CLUBFEET

TORTICOLLIS

SCOLIOSIS



Developmental Dysplasia of Hips

- Incidence 1:1000 live births
- Risk factors
 - 1st female child
 - Family history
 - Breech
 - Condition predisposing to crowding of uterus
 - Multiple pregnancies
 - Primipararity
 - Fibroids or other uterine growths
 - Low AFI
 - Big baby eg in GDM



Management

Diagnosis

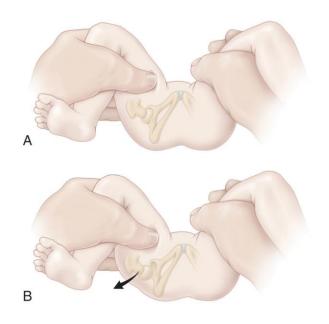
- Clinical
- Ultrasound
- Xrays



Management

Diagnosis

- Clinical
- Ultrasound
- Xrays



Barlow maneuver

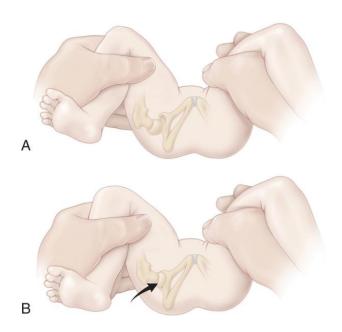
-Test of "dislocatability"



Management

Diagnosis

- Clinical
- Ultrasound
- Xrays



Ortolani maneuver



Management

Diagnosis

- Clinical
- Ultrasound
- Xrays

Beyond 3 months

- Reduced Abduction
- Asymmetrical crease
- LLD (Galeazzi positive)
- Treledenburg Gait



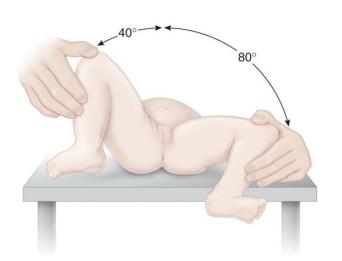
Reduced Abduction







Reduced hip abduction

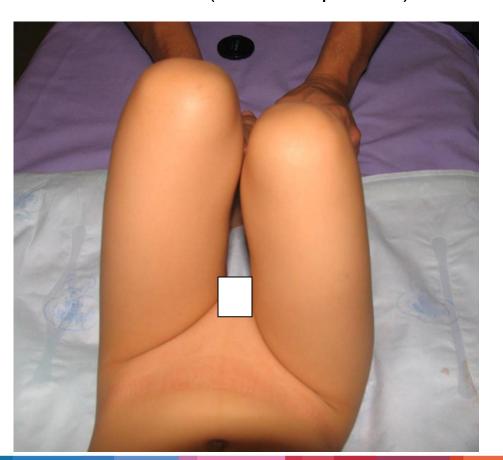




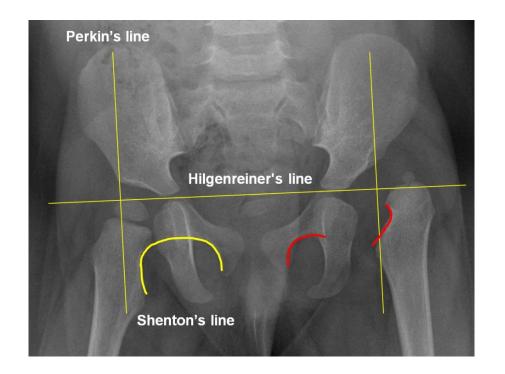


LLD (Galeazzi positive)













GOAL:

CONCENTRICALLY REDUCED AND STABLE HIPS WITH GOOD ACETABULAR COVER

To Avoid Early Degenerative Disease of the Hips















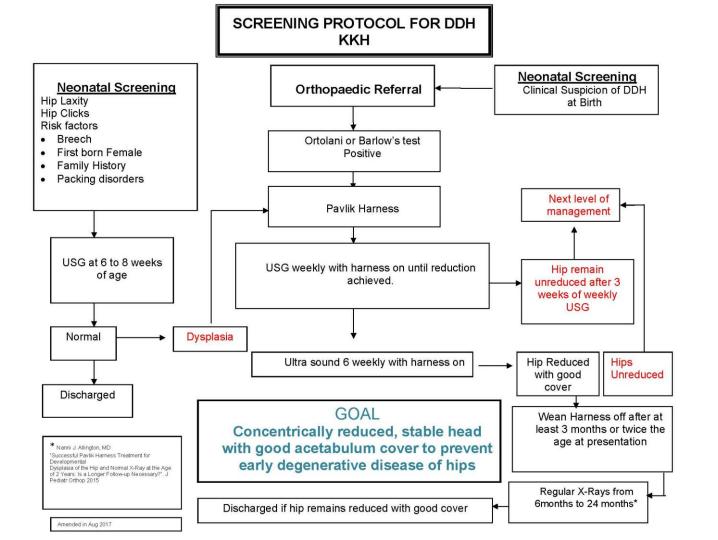












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Up to 6 Months

Clinically reducible hips

- Pavlik Harness
- Check U/S weekly until hip reduced
- Then 6 weekly
- Xrays at 6,and 18 months
- Discharged with
 - Spherical
 - Concentrically reduced
 - Well covered hips







Pavlik Harness

Anterior straps – to keep hips flexed at 100°

Posterior straps – to limit adduction of the hips (4 finger breadths)







Pavlik harness worn full time for 3 months then wean





DDHBirth to 6 Months



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1st Xray at 4 months

Follow up at 8 months



2 year old girl Neglected DDH

























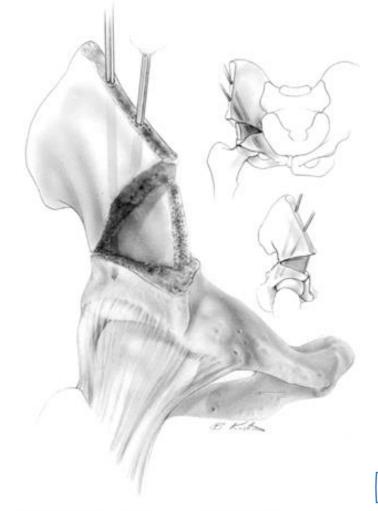


Beyond 18 months

- Remodelling potential of acetabulum thought to decrease after this age.
- Not enough to just reduce the hip
- Need to also provide acetabular cover



DDHSalter Osteotomy





Intra operative



























4 month post op





























3 years old (1 year post op)





4 years old (2 years post surgery)







Age		Treatment
Newborn to 6 months	Reducible hips Ultra sound diagnosed	Pavlik harness
	Irreducible hips Teratologic hips	Adductor release and Closed Reduction Adductor AND Iliospoas release by Medial Approach
6 months to 18 months		Adductor and Iliospoas release. Open reduction. Capsuloraphy.
> 18 months		Add Salter Possibly Femoral shortening after 2 years



Introduction

Incidence 1:1000

Subgroups

- Flexible or positional
 - If able to Dorsiflex foot then Positional
- Structural
 - Idiopathic
 - Neuromuscular / Paralytic (L4)
 - Syndromic



Introduction

Clinical features

- Hindfoot EquinoVarus
- Forefoot Supination
- Metatarsus Adductus
- Cavus

CAVE





Management

Goal of Treatment

- Painless foot
- Plantigrade foot
- Functional foot

Principles of Treament

- Correct Early
- Correct fully
- Maintain correction



- A particular way of serial casting and manipulation
- A particular sequence of correction
- Percutaneous Tenotomy part of serial casting and manipulation
- A particular way of maintaining the correction



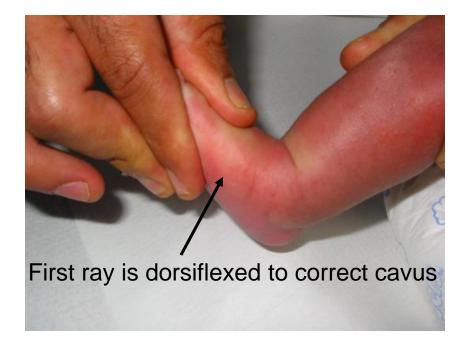
Cavus

Metatarsus Adductus

Hindfoot varus

Hindfoot equinus





Counter pressure On the medial Distal tibia Talus reduced

To correct MTA

No attempt at manipulating hindfoot to avoid rocker bottom foot





No attempt at manipulating hindfoot to avoid rocker bottom foot

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Serial Casting and Manipulation



Toe to Groin casting with Plaster of Paris

Hindfoot not forced into dorsiflexion

Forefoot in supination and abduction.



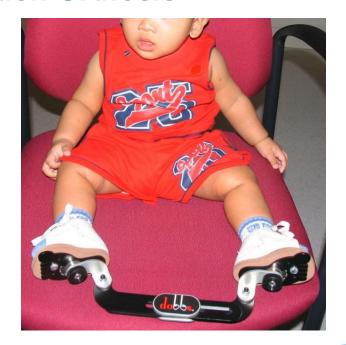
Percutaneous tenotomy after forefoot correction





Foot Abduction Orthosis

- 23hrs/ day for 3 months
- Then wean till child being to cruise
- Then night and nap till 4 years old.





Introduction

Clinical description, sometimes called **wry neck**Classification

- Muscular/idiopathic tight sternocleidomastoid (SCM)
- Secondary
 - Squint
 - Hearing problems
 - Flat head (plagiocephaly)
 - Bony fusion of cervical spine (Klippel Feil)
 - Tumours and infections in the neck
 - Trauma



1 month old boy

Pseudotumour of L SCMI POSNA





Clinical features

Childs tilts or looks to the sides opposite to the tight SCM

Facial asymmetry on the side of the torticollis





Clinical features

 Limited lateral rotation on the same side of torticollis

 Limited lateral flexion on the opp side of torticollis







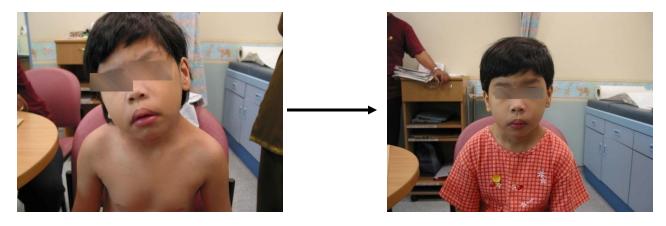
Management

- Exclude secondary causes
- If under 1 year can be treated effectively with stretching





TORTICOLLIS Surgery



- Surgery usually needed for those above 1 year old.
- Release of the SCM is done with or without a reconstruction of the sternal head



SCOLIOSIS



CLASSIFICATION

- Idiopathic
- Congenital
- Neuromuscular
- Syndromic e.g Marfan's syndrome



CLASSIFICATION

Idiopathic

- Adolescent
 - after 11 years of age to the end of skeletal growth
- Juvenile
 - at age 5 to 6 years
- Infantile
 - in the first year of life



CLASSIFICATION

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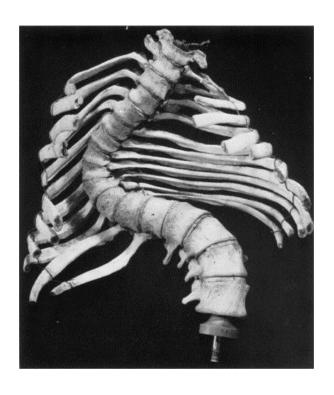
ETIOLOGY

- Neurologic Dysfunction
- Connective Tissue Abnormalities
- Genetic Factors

The exact cause of idiopathic scoliosis remains unknown, despite considerable investigation



PATHOPHYSIOLOGY



- Trapezoidal verterbrae
- Rotation of spine producing the hump
- Reduced lung capacity
- Neurologic deficits are rare



Epidemiology

- Prevalence
 - 6-7 years: 0.05% (girls), 0.02% (boys)
 - **11-12 years: 1.37% (girls),** 0.21% (boys)
 - **13-14 years: 2.22% (girls),** 0.66% (boys)
- Recommends
 - screening of girls between 11-14 years

Idiopathic Scoliosis in Singapore Schoolchildren

A Prevalence Study 15 Years Into the Screening Program

Hee-Kit Wong, FRCS, James H. P. Hui, FRCS, Uma Rajan, FA, MS, and Hwee-Pin Chia. MMed



Spinal screening

 The Forward Bending Test is the method used to screen for abnormal spinal curve.



An angle of 6° and greater requires an Xray and review by spine specialist.











PRESENTING COMPLAINT

- Cosmesis
- Backpain

Respiratory symptoms

Neurologic deficits



PRESENTING COMPLAINT

- Cosmesis
- Backpain uncommon
 - to investigate further if present
- Respiratory symptoms uncommon
 - Occurs when curve magnitude reaches 100°
- Neurologic deficits rare
 - More common in left sided curves e.g. syrinx



Physical examination

General inspection

- Syndromic features
- cafe au lait spots
- subcutaneous nodules
- axillary freckles, which are seen in neurofibromatosis
- hairy patches
- skin dimples: an underlying sign of spinal dysraphism
- Feet (Cavovarus)

Back examination

- Posture from front, side and back
 - Sagittal profile-kyphosis
 (AIS usually hypokyphosis
- Shoulder and pelvis level
- Spinal balance C7 plumbline, listing
- Adam Forward Bending Test
 - Scoliometer
 - Hump, rib asymmetry
- LLD, block test

Neurological examination

- LL (motor, sensory and reflexes)
- Abdominal reflexesasymmetrical (syringomyelia/ACM)
- Back sensation --> suspended sensory loss
- Hamstrings (poplitefemoral
- angle)
- Gait: short leg gait from LLD



C7 plumbline

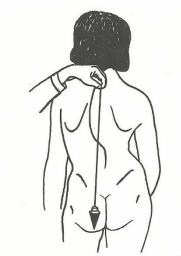


Figure 13.7. Diagram of a left thoracic scoliosis with a plumb dropped from the C7 spinous process. Weight shift can be noted by the plumb line striking the left buttock.



PHYSICAL EXAMINATION

Adams forward bending Test

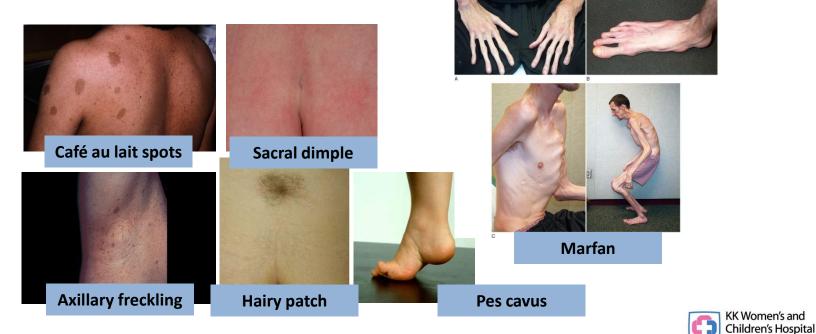




Angle of trunk rotation $(ATR) > 6^{\circ}$

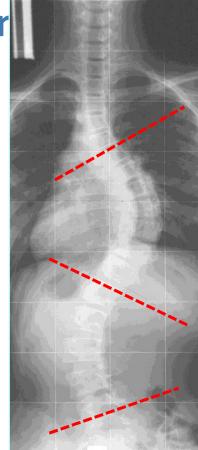


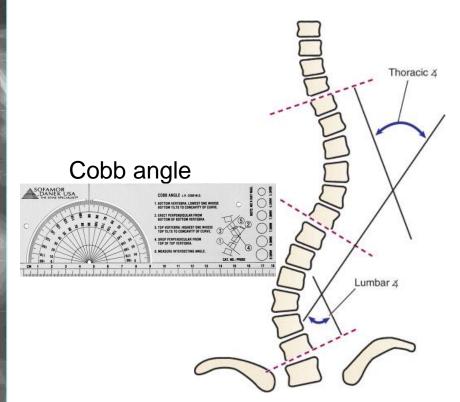
Signs of underlying disorders



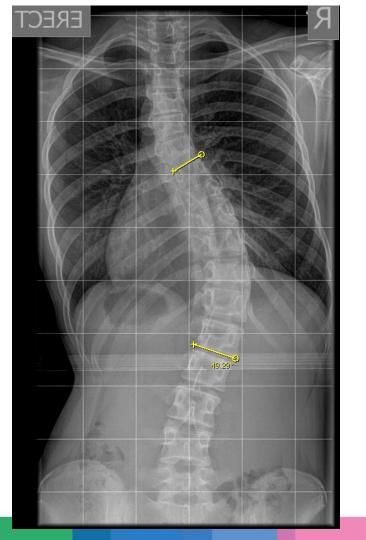
SingHealth

Plair











Digital measurement using PACS



TREATMENT GOAL

Stop Progression of Curve to prevent effects of natural history







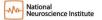


















TREATMENT

Observation Brace Surgery



Treatment Guide

	Risser's Sign		
Curve Magnitude	Grade 0 /Premenarchal	Grade 1 or 2	Grade 3, 4, or 5
< 25°	Observation	Observation	Observation
30°-40°	Brace therapy - begin when curve is >25°	Brace therapy	Observation
>45°	Surgery	Surgery	Surgery - when curve is >50°



Observation

No treatment is needed for curves < 25°, regardless of the patient's maturity.



Brace

Indication

- immature children(Risser grade 0, 1, or 2)
- curves in the range of 30° to 45° or who have documented progression exceeding 5°
- Risser grade 0 when their curves reach 25°

Contraindication

- large curves (>45°) in a growing adolescent
- patients who find wearing an orthosis to be emotionally intolerable



Brace Types

Milwaukee Brace Boston Brace



Brace Types

Milwaukee Brace

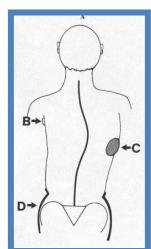
Boston Brace

a pelvic girdle, a suprastructure, and

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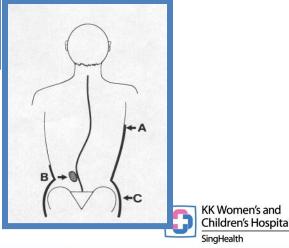
Brace Types

Milwaukee Brace

Boston Brace

- individually constructed by an orthotist
- Most commonly usedTLSO today
- apex at T7 or below





Surgery

Goals

- reduce the magnitude of the deformity
- to obtain fusion to prevent future curve progression
- to do so safely

Should result in a well-balanced spine in which the patient's head, shoulders, and trunk are centred over the pelvis



Surgery

Indications

- > 50° at skeletal maturity
- Thoracolumbar and lumbar curves >40° to 45°
- Smaller curve with poor cosmesis



Treatment Guide

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Conclusion

Main focus for the primary physician in the management of Common Congenital and Developmental Conditions in Paediatrics Orthopaedics are:

- Accurate diagnosis
- To determine or even assist in screening to change natural history for the better.





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