

# Idiopathic Scoliosis

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# Disclosures

- None relevant to this talk

# Learning Objectives

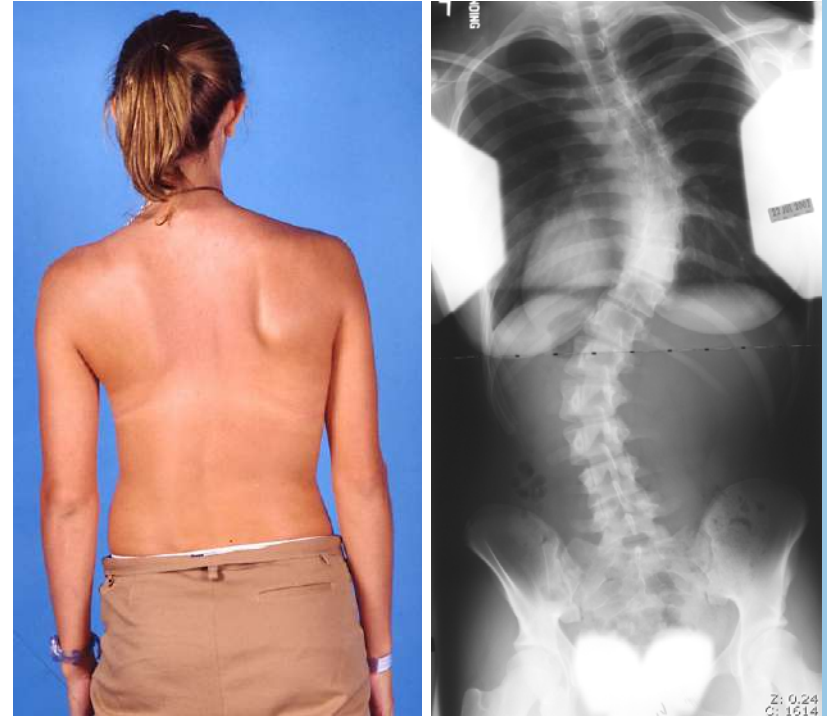
- Recognize different types of spinal deformity in children
- Understand common presentation & physical exam findings in children with spinal deformity
- Understand radiographic assessment of Scoliosis
- Recognize “atypical” presentations
- Understand the concepts behind treatment of childhood spinal deformity
- Understand the importance of early recognition of post-operative complications



# Pediatric Scoliosis

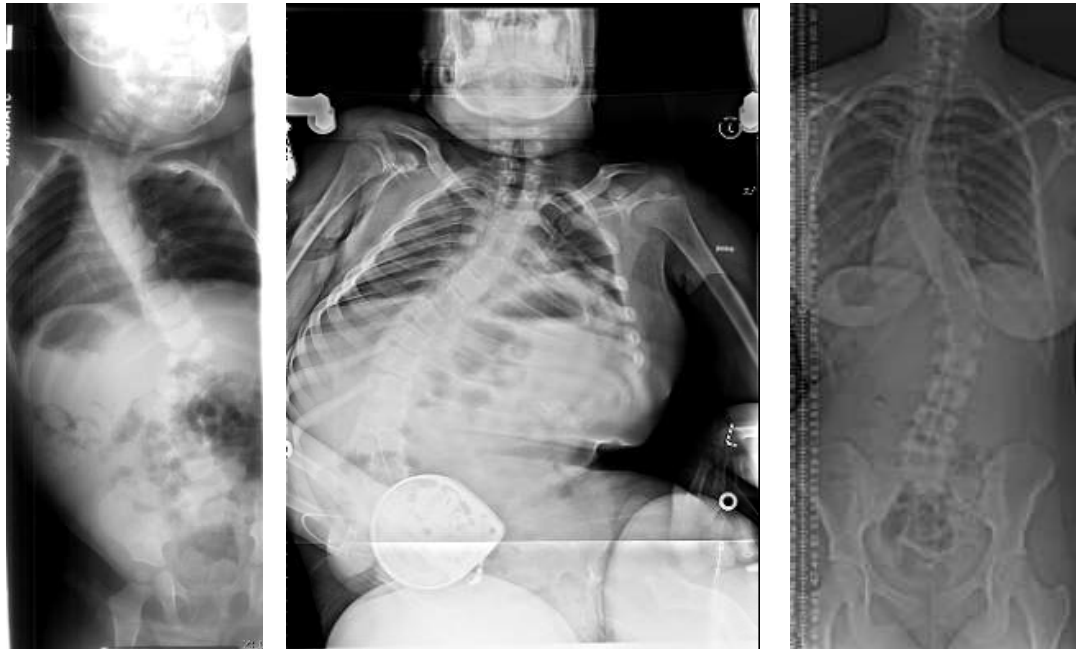
## Definition:

- Structural spinal deformity characterized by decompensation of the normal vertebral alignment during rapid skeletal growth
- Deformity is 3 dimensional
  - Coronal, sagittal, as well as abnormal vertebral rotation



# Types of Scoliosis

- **Congenital** - due to a congenital abnormality of the vertebrae or fused ribs
- **Neuromuscular** - caused by problems such as poor muscle control or muscle weakness, or paralysis due to diseases
  - Cerebral Palsy
  - Myelomeningocele
  - Muscular Dystrophy
- **Idiopathic scoliosis** - is scoliosis of unknown cause. Genetic in origin.



# Idiopathic Scoliosis

- By far the most common type >90%
- No definite cause
- Believed to be related to asymmetric growth of vertebral bodies
- Types
  - Early-Onset
    - Infantile Scoliosis
    - Juvenile Scoliosis
  - Late-Onset
    - Adolescent Scoliosis



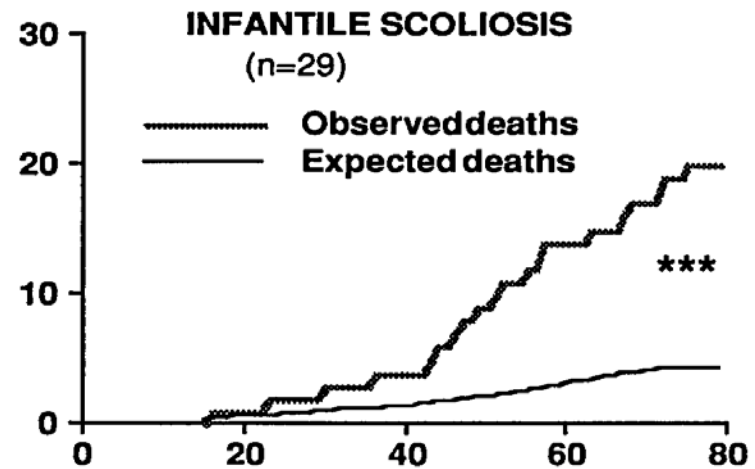
# Infantile Idiopathic Scoliosis

- Develops before 3 years of age
- Often occurs w/ other congenital abnormalities
- 80-90% resolve w/o tx
- 90% w/ left thoracic curve



# Infantile Idiopathic Scoliosis

- The good news: often resolves!  
Up to 92% of curves reportedly resolve
- The bad news: if it doesn't resolve  
it may kill you





# Bracing in EOS

- Advantages – removable, lighter, widely accessible
- Disadvantages – removable, rib wall deformity
- Contraindications (relative)
  - children with decreased pulmonary function



# Casting

- Used in progressive curves when unable to control with a brace, either due to noncompliance or increasing stiffness
- Can perform serial casting to decrease deformity for bracing



# Casting - Brief History

## Risser

- Introduced casting in early 1900s for AIS
- Principles elongation and derotation

## Morel and Cotrel (1964)

Built on Risser's principles but added

## Mehta (1975-2000)

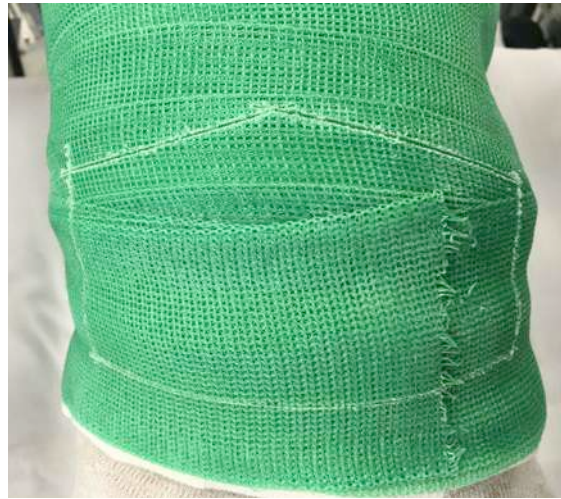
Modified technique of Morel and Cotrel

- Emphasized early intervention
- Demonstrated ability to cure EOS especially

# Casting Technique

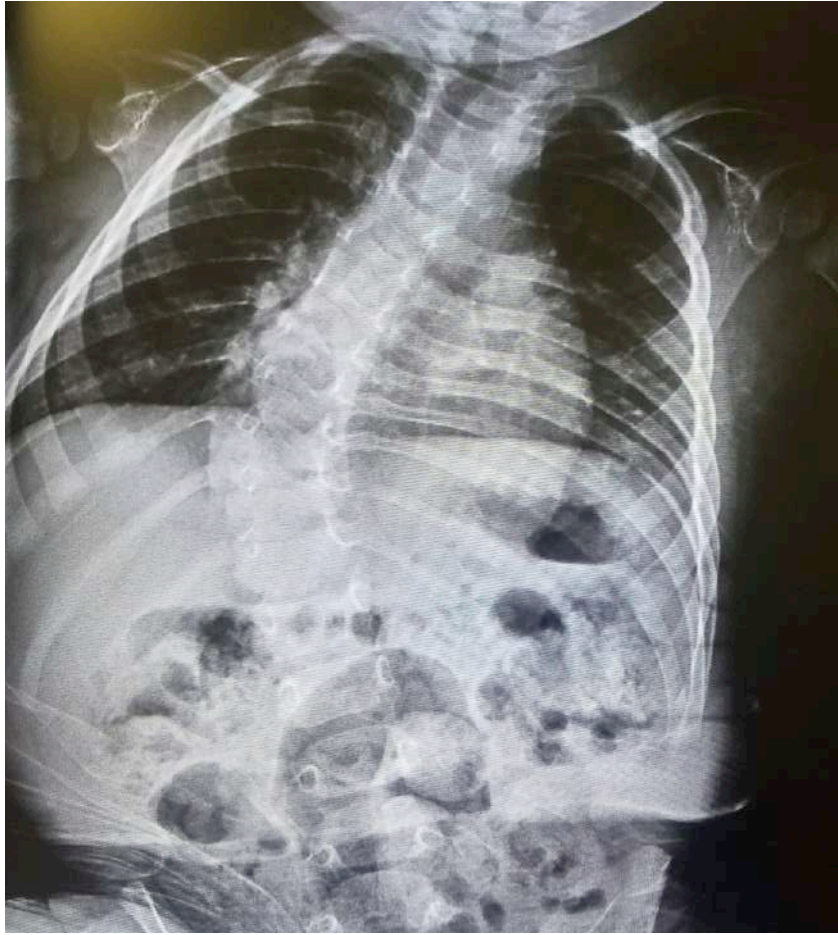


# Casting Technique



# Casting as definitive treatment

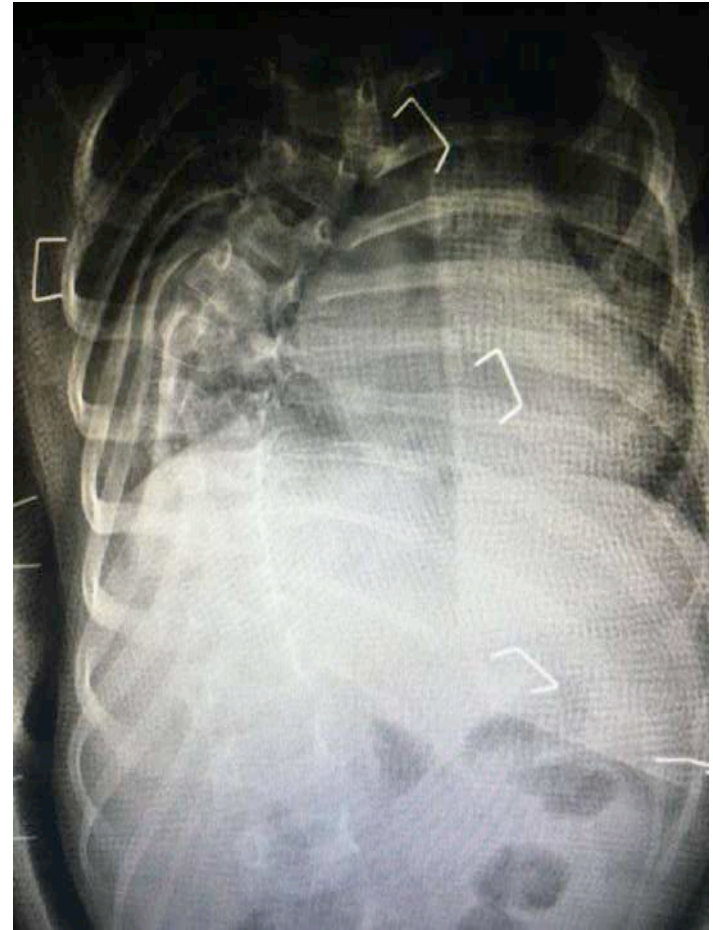
Age 12 months



# Cast to buy time



**10 months later**



# Complications of Casting

- Chest wall deformity
- Eating intolerance
- Decreased pulmonary function (temporary)
- Cast sores
- Brachial plexus injury
- Cranial nerve palsy
- Subclavian thrombosis





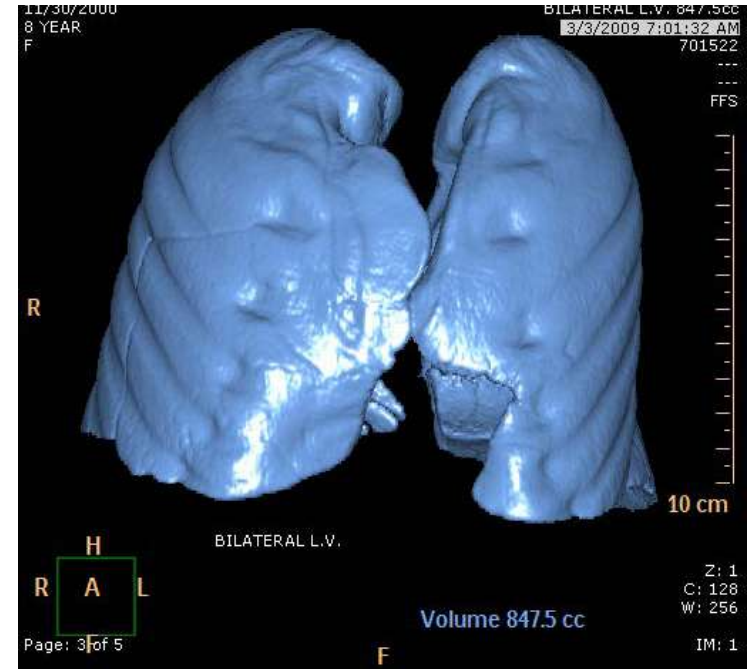
# What about when casting fails? Traction

- Indicated in stiff curves with inability to brace or cast due to larger curves or proximal thoracic curves
- Goal is to increase traction as tolerated until:
  - Tiptoes if standing
  - Barely on buttocks if in wheelchair



# Benefits of traction

- Coronal deformity (35%) \*
- Trunk shift (65%)\*
- Sagittal plane (30%)\*
- Pulmonary function?
- Mobilization
- Increased surgical correction
- Decreased neurologic complications intraoperatively













# Multiple Surgical Options



- Stay tuned – we are still figuring this out



# Juvenile Idiopathic Scoliosis

- 3-10 years of age
- Curves are often progressive
  - Potential for
    - severe truncal deformity
    - cardiopulmonary complications
    - Significant impact on lung maturation
  - 90% will need surgery due to magnitude of deformity and progression potential



# Adolescent Idiopathic Scoliosis

- 11-18 years of age
- Approximately 90% of childhood spinal deformity
  - Prevalence 1-3%
  - Female-male ratio
    - 6:1 non-operative
    - 7:1 operative
- > 90% w/ right thoracic curve
- Majority asymptomatic
  - Back pain is common
- Patient/Family perception of body asymmetry



# Risk Factors

- Family History
  - 30% incidence in daughters of women with AIS
  - 73-92% monozygotic twin concordance; 36-63% dizygotic
- Peak adolescent growth spurt



# Physical Examination

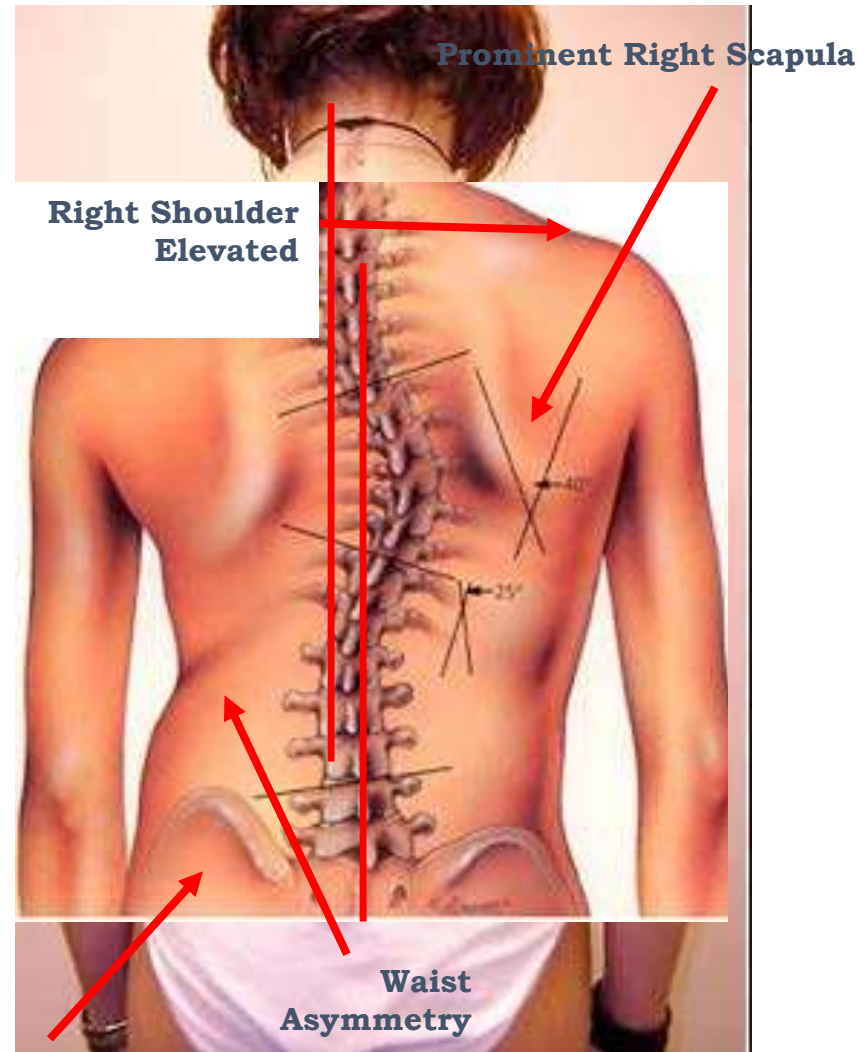
- Skin
  - Look for birthmarks, dimpling, hairy patches
- Strength
- Gait
  - Heel/Toe Walk
    - Can be used to elicit subtle motor weakness
- Neurologic Exam
  - Strength, sensation, deep tendon reflexes
    - Upper Extremities
    - Lower Extremities
    - Abdominal reflexes
    - Gag reflex
    - Supine straight leg raise
- Sensation



# Physical Examination

## Back Evaluation

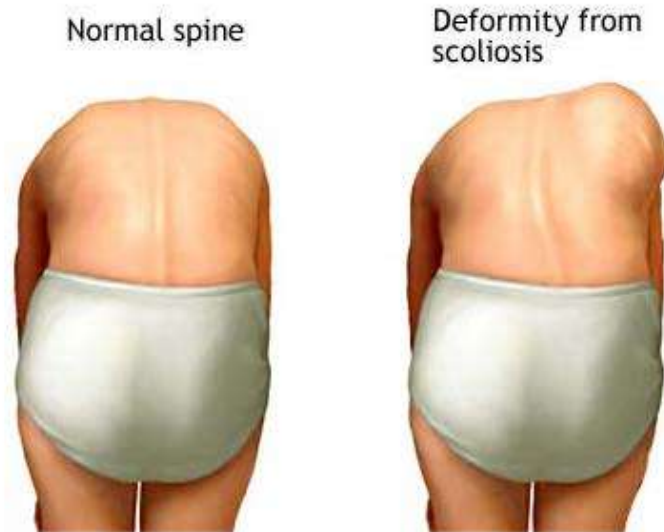
- Anterior, posterior, lateral view
- Truncal asymmetry
  - Thorax, ribs, breasts
- Shoulder height
- Waist asymmetry
- Leg-length inequality



# Physical Examination

## Adam's Forward Bend Test

- Forward bend at waist
  - Identify paraspinal prominences
  - Result of abnormal vertebral rotation and coronal
- **Scoliometer**
    - Used to quantify paraspinal prominences
    - Positive result is  $> 7$  degrees



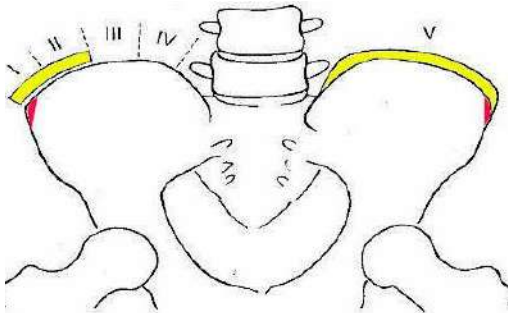
# Radiographs

- PA and Lat of the entire spine
- EOS



# Radiographs

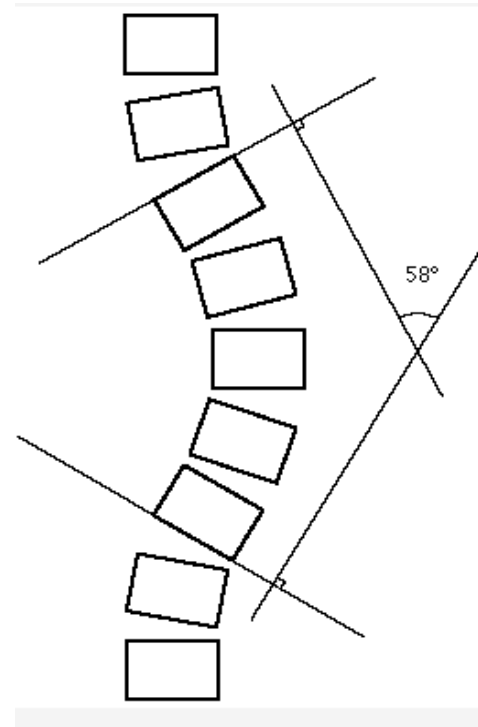
- Risser Sign:
  - Iliac apophysis develops from lateral to medial on AP view of pelvis
  - Risk of progression in Risser 1 or less as high as 70%
  - Risser of 3 has risk of progression ~ 10%
- Triradiate Cartilage
  - Closure coincides w/ the end of peak adolescent growth spurt





# Radiographic Examination

- **Cobb Angle:**
  - Should be measured for all curvatures present
  - > 10 degrees establishes a diagnosis of scoliosis



# Radiographic Examination

- **Kyphosis:**

- Assessment of sagittal thoracic contour
- Patients w/ AIS are usually hypokyphotic, but convex rib prominence may give appearance of increased kyphosis



# Radiographic Examination

- **MRI:**

- Used to evaluate patients with “atypical” presentation
- Should include brain (posterior fossa) and entire spine (C,T,LS)
- Include Gadolinium contrast
- Neural axis lesions
  - Syringomyelia
  - Chiari malformation
  - Tethered cord
  - Tumors



# Atypical Presentation

- Signs/symptoms that may suggest non-idiopathic deformity
  - Rapid progression
  - Large curve at dx
  - Left sided T curve
  - Pain that limits activity
  - ANY neurologic symptom/finding
  - Early onset



# Natural History?

- Progression with growth
- Risk of curve progression is related to patient's maturity (Risser sign, menarchal status) and to the size of the curve
- Larger curves with more growth more likely to progress

Lonstein (1984) *Progression >5° in curves:*

	<u>Less than 20°</u>	<u>20°-29°</u>
<i>Risser 0 or 1</i>	22%	68%
<i>Risser 2-4</i>	2%	23%

Risk of Scoliosis Progression			
Degree of Curve (Cobb Angle)	Age 10-12	Age 13-15	Age over 16
<20°	25%	10%	0%
20°-30°	60%	40%	10%
30° -60°	90%	70%	30%
>60°	100%	90%	70%

The above data has been obtained from the Scoliosis Research Society.

# Treatment

- Based on initial deformity and risk of curve progression
  - Overall assessment of growth potential
- Goal is to prevent progression until skeletal maturity
  - Risk of curve progression decreases

<b><u>Degree of Curve</u></b>	<b><u>Treatment</u></b>
10° to 25°	Close observation – PA spine Q4-5months
25° to 45°	TLSO bracing
>45°-50°	Posterior Spinal Fusion

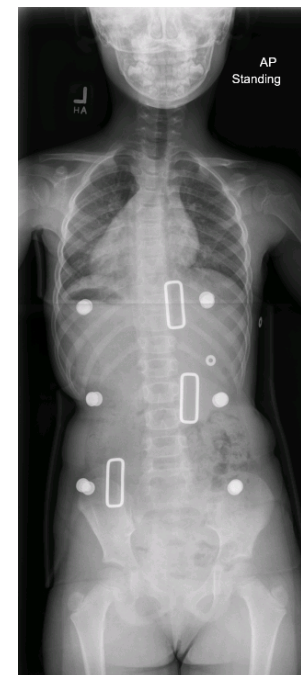
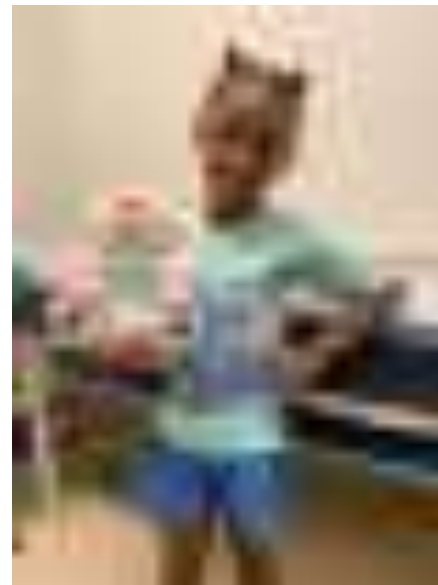
# Therapy

- **Observational Monitoring**
  - Follow-up standing PA and lateral scoliosis x-rays at 4-12 month intervals depending on growth potential
- **Core strengthening & conditioning exercises**
  - Recommended for all patients, especially those with pain
    - Physical Therapy techniques
    - Yoga/Pilates



# Bracing

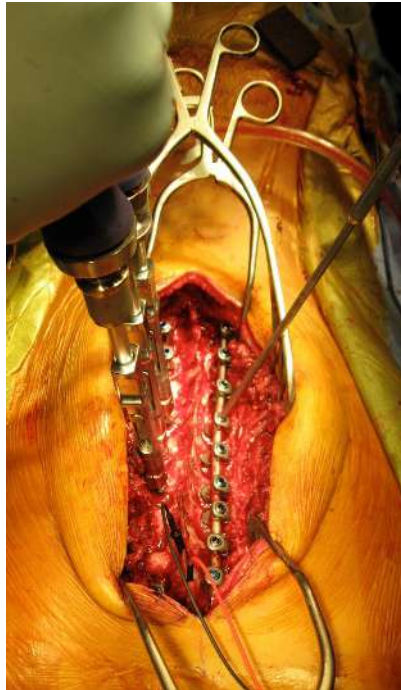
- Braist study (NEJM 2013 Weinstein et al.)
  - Study stopped short because of success
  - Brace wear positively associated with decreasing rate of progression to surgical grade curve
  - More brace wear associated with greater success with greatest success seen in those wearing the brace >12.9 hours daily
- Cobb angle between 25 & 45 degrees who are at increased risk of progression
  - Premenarchal
  - Risser < 2





# Surgical Treatment

- Cobb angle  $> 45$  degrees w/ risk of progression
- Spinal arthrodesis w/ instrumentation



# Goals of Surgical Treatment

- Stop curve progression
- Achieve maximal deformity correction
- Improve appearance w/ trunk balance
- Reduce short- and long-term complications



# Surgical Treatment Technique

- Choice of surgical approach and technique is dependent on deformity, flexibility, and surgeon preference
  - Most treated by posterior approach
  - Some by anterior approach
  - Occasionally combined approach



# Post-operative Care

- Typical hospital stay of 3 days
- No post-operative brace
- Return to school in 3-6 weeks
- Activity restrictions for at least 6 months



# Complication Risks with Surgical Treatment

	National	NCH
• Blood loss (transfusion rate)	~ 50%	0 since 2014
• Infection	.9-3-%	<.01%
• Neurologic injury	1-2 per 1000	NR*
• SMA Syndrome	1-4%	NR*

\*NR- non-reported



# Take Home Points

- Who
  - Spinal deformity in children is relatively common
  - Can develop at any age
  - Predominately effects girls
- Diagnosis
  - Clinical exam
  - Full length PA and Lat scoliosis films
  - Recognize “atypical” presentations
  - In all age groups larger curves are associated with greater risk of progression
  - Significant effects on thoracic and pulmonary growth and ultimately mortality in patients <8yrs old especially in IIS
- Treatment
  - Varies based on age and severity of curve

# Thank You

From the Nemours Children's Hospital Orthopaedic Team

