Children's Hospital Colorado
Orthopedics Institute

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The role of serial casting in early onset scoliosis

Jeannie Huh, MD
David M. Baulesh, BA
Timothy Judkins, MD
Sumeet Garg, MD
Nancy H. Miller, MD
Mark A. Erickson, MD

Affiliated with
School of Medicine
UNIVERSITY OF COLORADO
ANSCHUTZ MEDICAL CAMPUS

Disclosures

- Neither I nor any member of my family has a financial relationship or interest with any proprietary entity producing health care goods or services

Study Objectives

- To review our institution's experience with serial casting in the management of children with EOS.
- To determine if there are any differences in response to serial casting based on EOS curve etiology.
- To determine the effect of serial casting on thoracic height growth in the management of EOS.

Methods

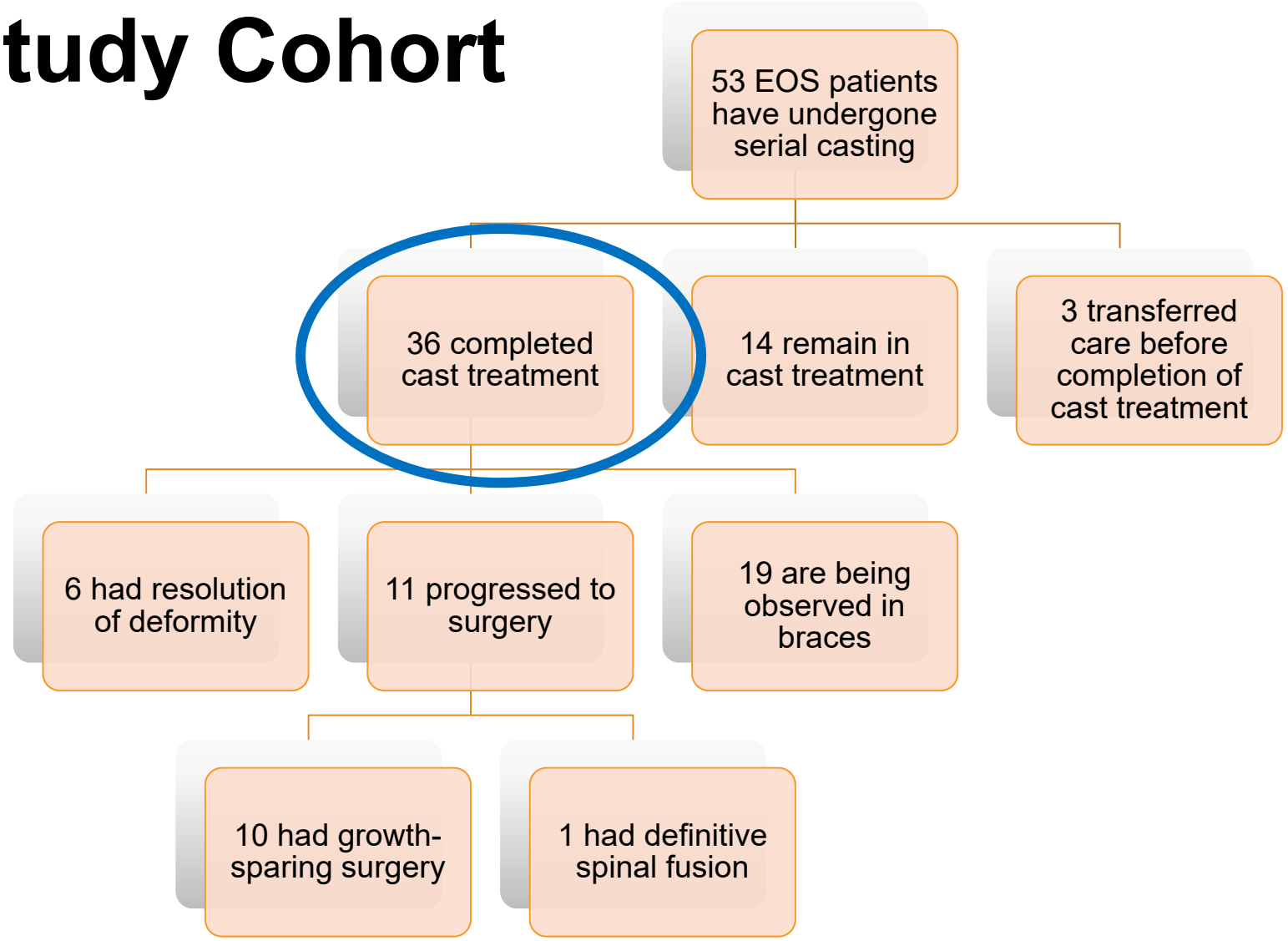
- Retrospective
- Serial casting database
- Single pediatric hospital
- Jan 2005 – Aug 2010
- Inclusion Criteria:
  - Diagnosis of EOS (<6yo)
  - Underwent at least 2 consecutive cast applications

Methods

- AMIL spine casting table
- Over-the-shoulder casts with anterior and posterior windows
- Casts changed every 8-12 weeks

- Classified into 2 groups
  - Idiopathic
  - Non-idiopathic

Study Cohort



Results

- No differences between idiopathic and nonidiopathic groups
  - Length of time in cast (range 3 – 47 months, average 13 months)
  - Follow-up (range 5 – 54 months, average 31 months)
  - # of casts (range 2-20, average 6)
  - Age at start of casting (range 0.2-6 yrs, average 2.6 yrs)

Complications

- Complication Rate: 19% (7/36)
  - Pulmonary (5)
    - 3 pneumonia
    - 1 bronchiolitis
    - 1 aspiration
  - Skin (2)
    - Superficial skin breakdown

- None required invasive interventions

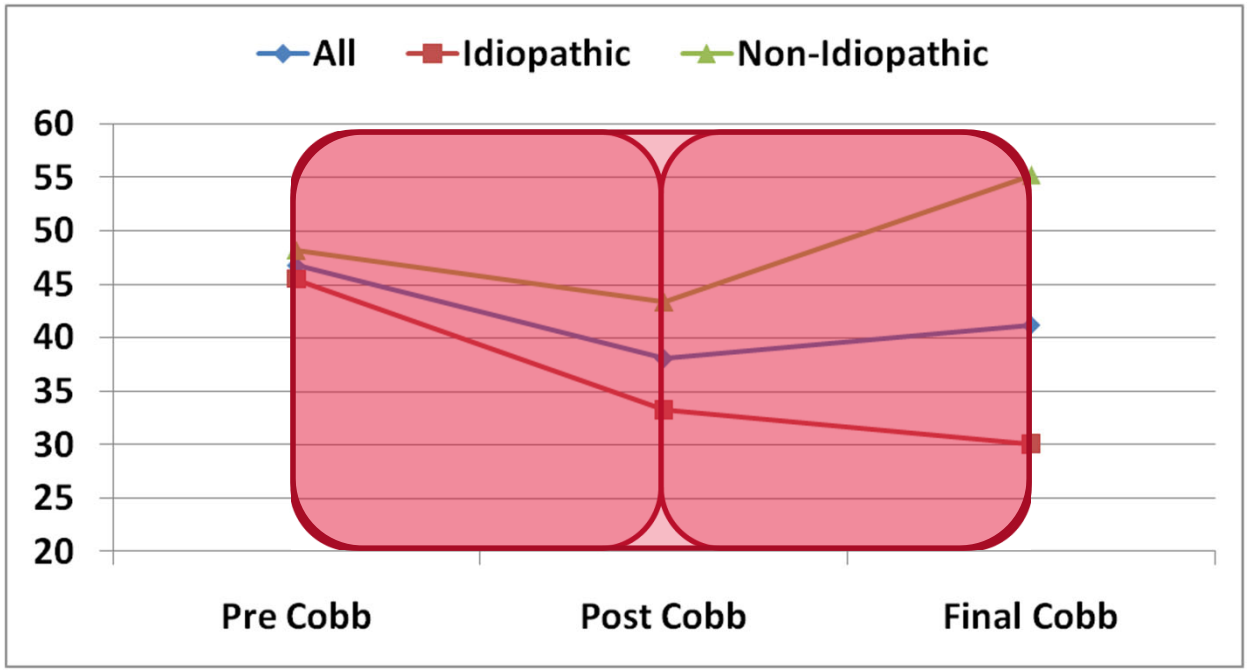
Results

- Surgery: 30.6% (11/36)
  - Avg age: 5.6 yo (2.7-8.0 yo)
  - Avg time delay from initial casting: **2.1 yrs** (0.4-4.2 yrs)

- Deformity Resolution: 16.7% (6/36)

	Resolution		
	n	**Percentage**	**P-Value**
Idiopathic	5	26.32%	p=0.1821*
Non-Idiopathic	1	5.88%	

*Fisher's Exact Test



	Pre-Post Cobb	Pre-Final Cobb	Post-Final Cobb
All*	**8.7 (p=0.0014)**	5.7 (p=0.1248)	-3.1 (p=0.1792)
Idiopathic†	**12.2 (p=0.0032)**	**15.4 (p=0.0052)**	3.2 (p=0.2532)
Non-Idiopathic†	4.8 (p=0.1018)	-6.5 (p=0.1392)	**-11.0 (p=0.0071)**

*Student's t-test
†Wilcoxon Sign Ranked

Thoracic Height Velocity

	T-Height Velocity: Subject Mean (cm/year)	T-Height Velocity: Population Mean	P-value*
All	1.56	1.4 cm/year	p=0.6501
Idiopathic	1.59	1.4 cm/year	p=0.5619
Non-Idiopathic	1.52	1.4 cm/year	p=0.8926

*Wilcoxon Sign Ranked

Discussion

- Study Limitations:
  - Retrospective study design
  - No control observed or braced group
  - Small sample size
  - Follow-up is not to skeletal maturity
  - Limited objective parameters to assess treatment success
  - No functional or quality of life measures

Discussion

- First study evaluate efficacy of serial casting in EOS using thoracic height velocity

- Correction, surgical delay and complications comparable to recent studies of casting
  - Tsuji SRS 2008
  - Fletcher ICEOS 2010

Conclusions

- Serial casting leads to improved curve correction in idiopathic compared to non-idiopathic EOS

- Serial casting in EOS delayed surgery an average of 2.1 years in our cohort

Conclusions

- Thoracic height growth continues at normal velocity during serial casting for EOS.
  - Benefit of casting vs. natural history?

- Further studies on serial casting in the management of EOS are warranted to explore :
  - Outcomes using other objective measures of treatment success (e.g. quality of life, pulmonary function)
  - Results at skeletal maturity

Thank You!