

Intraspinal MRI Abnormalities in Early-Onset Scoliosis – Rates Across a Global Cohort

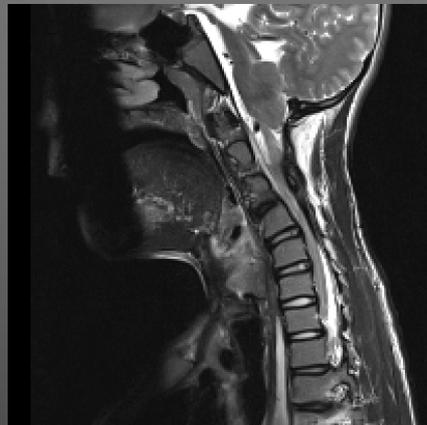
Anna McClung, Brendan A. Williams, Suken A. Shah, Laurel C. Blakemore, Jeff Pawelek, Paul D. Sponseller, Stefan Parent, John B. Emans, Peter F. Sturm, Burt Yaszay, Behrooz A. Akbarnia, and the Growing Spine Study Group

Background

- Spinal MRI is commonly included in the evaluation of EOS due to higher frequencies of intraspinal abnormalities reported in this population
- MRI findings across a diverse, multi-center EOS cohort have not been previously described

Objectives

- 1) To report on the rate and type of abnormalities identified by spinal MRI from a diverse EOS cohort within an international patient registry
- 2) To identify patient-related factors associated with a higher likelihood of MRI abnormality



Methods

Design: *Retrospective review of a prospective, multi-center database*

Inclusion criteria: *Idiopathic, Congenital, Neuromuscular or Syndromic EOS patients in whom MRI was obtained*

Exclusion criteria:

- *Incomplete or unverifiable data regarding pre-treatment imaging*
- *Structural deformities secondary to tumor or infection*

Independent variables:

- *Patient demographics: Age, race/ethnicity*
- *Etiology of EOS*
- *Major curve size (Degrees)*
- *Type of treatment (Operative or Non-operative)*

Dependent variable: *Presence (**MRI+**) or absence (**MRI-**) of MRI abnormality*

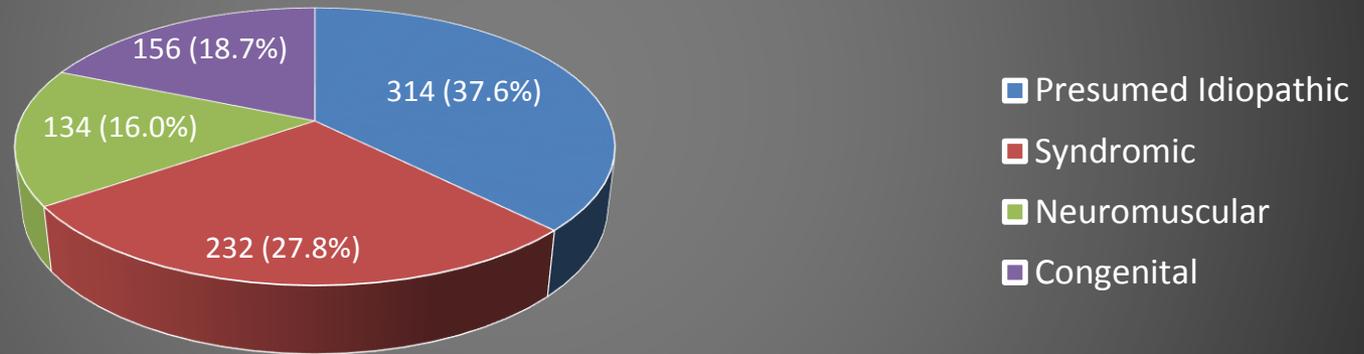
Statistical Analysis

- Demographic, clinical and radiographic characteristics summarized with descriptive statistics
- MRI findings were summarized with frequency distributions by abnormality type, patient and EOS Etiology
- Univariate analyses were performed using Pearson's chi-square (χ^2) for categorical variables and two-tailed student's t-test for continuous variables
- Multivariate logistic regression was performed to identify significant predictors of MRI abnormality

Cohort Demographics

- MRIs were obtained in 836 of 1,343 (62%) of registry subjects meeting inclusion criteria at mean age of 5.8 +/- 4.0 years old

Etiology Distribution of Patients Undergoing MRI

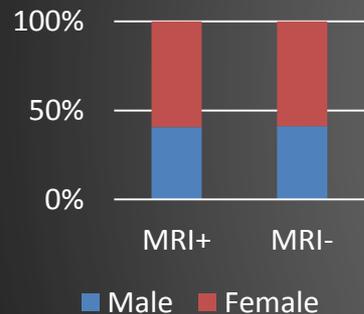


- 23.6% (197/836) of patients had positive MRI findings
- 247 unique MRI abnormalities were identified

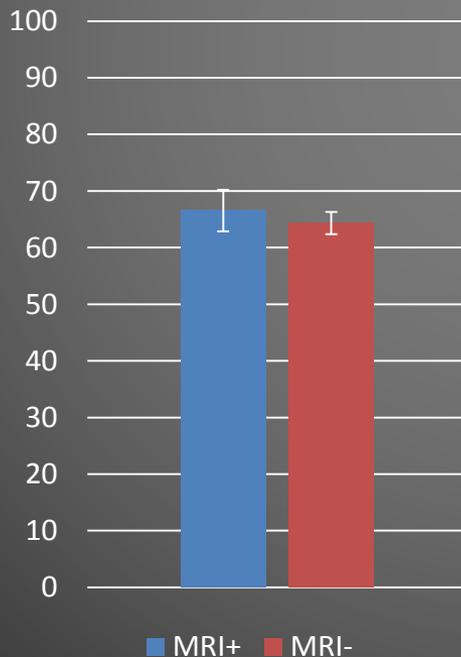
Univariate Analyses

- MRI+ showed no association ($p > 0.05$) with gender, treatment type, major curve size, age at MRI and age at treatment

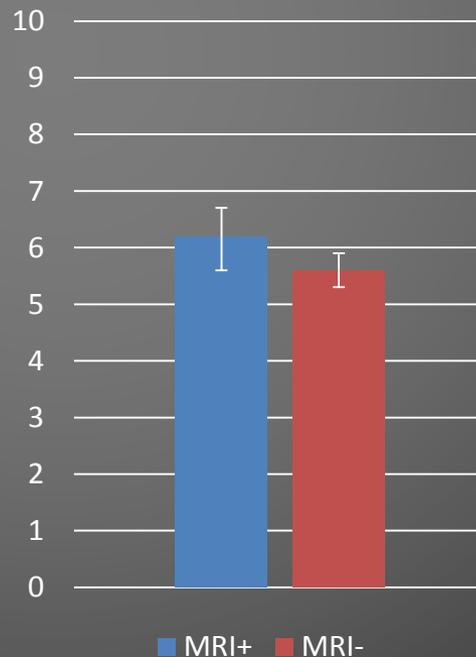
Gender



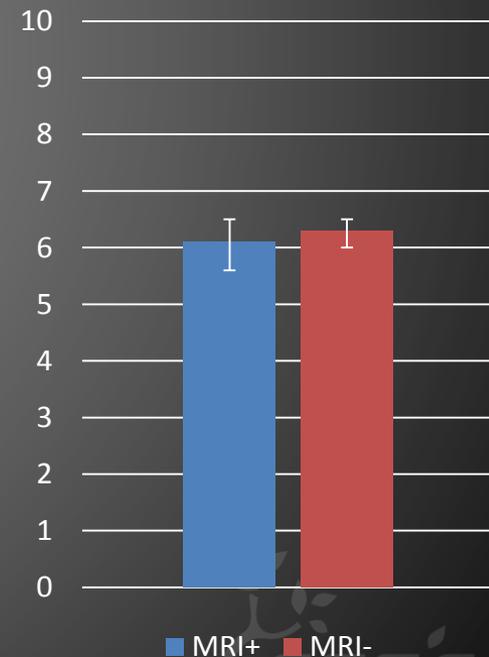
Major Curve (degrees)



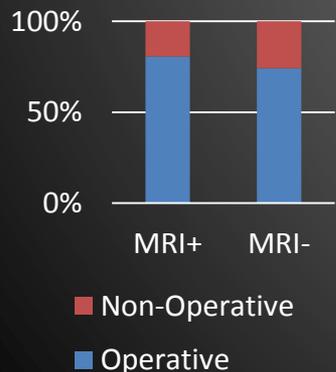
Age at MRI



Age at Treatment



Treatment Type



MRI Abnormalities - By Race/Ethnicity

Univariate Analysis

Race/Ethnicity	Abnormal (MRI+) n = 197 (23%)	Normal (MRI-) n = 639 (76%)	p-value
White/Caucasian	114 (57.9%)	410 (64.2%)	P = 0.002
African/African-American	23 (11.7%)	89 (13.9%)	
Hispanic	15 (7.6%)	57 (8.9%)	
Asian/Asian-American	18 (9.1%)	19 (3.0%)	
Other/Unspecified	27 (13.7%)	64 (10.0%)	

Multivariate Regression

Race/Ethnicity	Odd Ratio	95% CI	Adjusted OR*	95% CI
White/Caucasian	1*		1*	
African/African-American	0.9	(0.56, 1.54)	0.8	(0.48, 1.37)
Hispanic	1	(0.52, 1.73)	0.7	(0.37, 1.31)
Asian/Asian-American	3.4	(1.73, 6.71)	2.8	(1.39, 5.68)
Other/Unspecified	1.5	(0.93, 2.49)	1.3	(0.77, 2.15)

*Adjusted for etiology

MRI Abnormalities - By Etiology

Univariate Analysis

Etiology	Abnormal (MRI+) n = 197 (23%)	Normal (MRI-) n = 639 (76%)	P - value
Presumed Idiopathic	42 (21.3%)	272 (42.6%)	P < 0.001
Syndromic	48 (24.4%)	184 (28.8%)	
Neuromuscular	52 (26.4%)	82 (12.8%)	
Congenital	55 (27.9%)	101 (15.8%)	

Multivariate Regression

Etiology	Odd Ratio	95% CI	Adjusted OR*	95% CI
Syndromic	1*		1*	
Neuromuscular	2.4	(1.52, 3.89)	2.5	(1.55, 4.02)
Idiopathic	0.6	(0.38, 0.93)	0.6	(0.38, 0.96)
Congenital	2.1	(1.32, 3.30)	2.1	(1.31, 3.33)

*Adjusted for race/ethnicity

MRI Abnormality Distribution - By Patient

Patient MRI Findings	N (%)
Syrinx	43 (21.8%)
Tethered Cord with or without Fatty Filum	39 (19.8%)
Canal Abnormality	24 (12.2%)
Chiari Malformation	19 (9.6%)
Chiari Malformation and Syrinx	10 (5.1%)
Fatty Filum	9 (4.6%)
Syrinx And Tethered Cord	7 (3.6%)
Lipoma/Lipomeningocele	7 (3.6%)
Chiari Malformation and Spina Bifida	6 (3.0%)
Spina Bifida	6 (3.0%)
Other	5 (2.5%)
Dural Ectasia	4 (2.0%)
Syrinx and Fatty Filum	2 (1.0%)
Chiari Malformation, Syrinx and Tethered Cord	2 (1.0%)
Chiari Malformation and Tethered Cord	1 (0.5%)

Top 5 MRI Findings Within Each EOS Etiology

	Syndromic (n=48)	Neuromuscular (n=52)	Presumed Idiopathic (n = 41)	Congenital (n=55)
1.	Tethered cord - 11	Chiari Malformation - 9	Syrinx - 18	Tethered Cord +/- Fatty Filum - 19
2.	Syrinx - 9	Canal Abnormality +/- Other finding - 8	Chiari Malformation - 9	Syrinx - 10
3.	Canal Abnormality - 8	Chiari Malformation and Spina Bifida - 6	Chiari Malformation + Syrinx - 5	Canal Abnormality - 5
4.	Fatty Filum - 4	Syrinx - 6	Canal Abnormality - 4	Lipoma or Lipomeningocele - 4
5.	Dural Ectasia - 3	Tethered Cord - 6	Tethered Cord +/- Fatty Filum - 3	Fatty Filum - 4

Discussion

- In the largest and most diverse EOS cohort to date, a 24% rate of MRI abnormality was identified
- Multivariate logistic regression demonstrated:
 - Increased Odds for MRI Abnormality among *Asian/Asian-American* (2.8x vs. White/Caucasian), *Congenital* (2.1x vs. Syndromic) and *Neuromuscular* (2.5x vs. Syndromic) patients
 - Decreased Odds for MRI Abnormality among *Idiopathic* (0.6 vs. Syndromic) patients
- The most frequent abnormalities seen were Syrinx (22%) and Tethered cord (20%)
- The most common MRI findings in each etiologic subgroup are described

Limitations

- Registry studies rely upon the accuracy and consistency of data collected at participating centers
- No standardization of MRI review - Reported imaging findings based on each institution's local radiology report
- Other potential risk factors for MRI abnormality (e.g. physical exam findings) not available for inclusion

References

1. Rajasekaran S, Kamath V, Kiran R, Shetty AP. Intraspinous anomalies in scoliosis: An MRI analysis of 177 consecutive scoliosis patients. *Indian J Orthop*. 2010;44(1):57-63.
2. Pereira EAC, Oxenham M, Lam KS. Intraspinous anomalies in early-onset idiopathic scoliosis. *Bone and Joint Journal*. 2017;99-B(6):829-833.
3. Mohanty S, Kumar N. Patterns of presentation of congenital scoliosis. *J Orthop Surg* . 2000;8(2):33-37.
4. Gupta N, S R, G B, Shetty A. Vertebral and Intraspinous Anomalies in Indian Population with Congenital Scoliosis: A Study of 119 Consecutive Patients. *Asian Spine J*. 2016;10(2):276-281.
5. Zhang W, Sha S, Xu L, Liu Z, Qiu Y, Zhu Z. The prevalence of intraspinal anomalies in infantile and juvenile patients with “presumed idiopathic” scoliosis: a MRI-based analysis of 504 patients. *BMC Musculoskelet Disord*. 2016;17(1):189.
6. Inoue M, Minami S, Nakata Y, et al. Preoperative MRI analysis of patients with idiopathic scoliosis: a prospective study. *Spine* . 2005;30(1):108-114.
7. Koç T, Lam KS, Webb JK. Are intraspinal anomalies in early onset idiopathic scoliosis as common as once thought? A two centre United Kingdom study. *Eur Spine J*. 2013;22(6):1250-1254.
8. Nakahara D, Yonezawa I, Kobanawa K, et al. Magnetic resonance imaging evaluation of patients with idiopathic scoliosis: a prospective study of four hundred seventy-two outpatients. *Spine* . 2011;36(7):E482-E485.
9. Dobbs MB, Lenke LG, Szymanski D a., et al. Prevalence of neural axis abnormalities in patients with infantile idiopathic scoliosis. *J Bone Joint Surg Am*. 2002;84-A(12):2230-2234.
10. Evans SC, Edgar M a., Hall-Craggs MA, Powell MP, Taylor B a., Noordeen HH. MRI of “idiopathic” juvenile scoliosis. A prospective study. *J Bone Joint Surg Br*. 1996;78(2):314-317.
11. Pahys JM, Samdani AF, Betz RR. Intraspinous Anomalies in Infantile Idiopathic Scoliosis. *Spine* . 2009;34(12):E434-E438.
12. Williams BA, Matsumoto H, McCalla DJ, et al. Development and initial validation of the Classification of Early-Onset Scoliosis (C-EOS). *J Bone Joint Surg Am*. 2014;96(16):1359-1367.
13. Gupta P, Lenke LG, Bridwell KH. Incidence of Neural Axis Abnormalities in Infantile and Juvenile Patients With Spinal Deformity. *Spine* . 1998;23(2):206-210.
14. Lewonowski K, King JD, Nelson MD. Routine use of magnetic resonance imaging in idiopathic scoliosis patients less than eleven years of age. *Spine* . 1992;17(6 Suppl):S109-S116.
15. Fribourg D, Delgado E. Occult spinal cord abnormalities in children referred for orthopedic complaints. *Am J Orthop* . 2004;33(1):18-25.
16. Basu PS, Elsebaie H, Noordeen MHH. Congenital spinal deformity: a comprehensive assessment at presentation. *Spine* . 2002;27(20):2255-2259.
17. Suh SW, Sarwark JF, Vora A, Huang BK. Evaluating congenital spine deformities for intraspinal anomalies with magnetic resonance imaging. *J Pediatr Orthop*. 2001;21(4):525-531.
18. Prahinski JR, Polly DW Jr, McHale KA, Ellenbogen RG. Occult intraspinal anomalies in congenital scoliosis. *J Pediatr Orthop*. 2000;20(1):59-63.
19. Belmont PJ Jr, Kuklo TR, Taylor KF, Freedman BA, Prahinski JR, Kruse RW. Intraspinous anomalies associated with isolated congenital hemivertebra: the role of routine magnetic resonance imaging. *J Bone Joint Surg Am*. 2004;86-A(8):1704-1710.
20. Shen J, Wang Z, Liu J, Xue X, Qiu G. Abnormalities associated with congenital scoliosis: a retrospective study of 226 Chinese surgical cases. *Spine* . 2013;38(10):814-818.
21. Liu Y-T, Guo L-L, Tian Z, et al. A retrospective study of congenital scoliosis and associated cardiac and intraspinal abnormalities in a Chinese population. *Eur Spine J*. 2011;20(12):2111-2114.
22. Ghandhari H, Tari HV, Ameri E, Safari MB, Fouladi DF. Vertebral, rib, and intraspinal anomalies in congenital scoliosis: a study on 202 Caucasians. *Eur Spine J*. 2015;24(7):1510-1521.
23. Xue X, Shen J, Zhang J, et al. Rib deformities in congenital scoliosis. *Spine* . 2013;38(26):E1656-E1661.