



Risk Factors for Coronal Decompensation After Posterior Spinal Instrumentation and Fusion in Adolescent Idiopathic Scoliosis

Jaime A. Gomez, MD^a, Hiroko Matsumoto, PHDC^{a,*}, Nicholas D. Colacchio, MD^b,
David P. Roye, Jr, MD^{a,c}, Daniel J. Sucato, MD^d, B. Stephens Richards, MD^d,
John B. Emans, MD^e, Mark A. Erickson, MD^f, James O. Sanders, MD^g,
Lawrence G. Lenke, MD^h, Michael G. Vitale, MD, MPH^{a,c}

^aDivision of Pediatric Orthopedic Surgery, Department of Orthopaedic Surgery, Columbia University Medical Center, New York, NY 10032, USA

^bDepartment of Orthopedics, Tufts Medical Center, Boston, MA 02111, USA

^cDivision of Pediatric Orthopedic Surgery, New York-Presbyterian Morgan Stanley Children's Hospital, Columbia University Medical Center, New York, NY 10032, USA

^dDepartment of Pediatric Orthopedic Surgery, Texas Scottish Rite Hospital for Children, Dallas, TX 75219, USA

^eDepartment of Orthopedic Surgery, Children's Hospital Boston, Boston, MA 02115, USA

^fDepartment of Orthopedic Surgery, Children's Hospital Colorado, Aurora, CO 80045, USA

^gDepartment of Orthopedic Surgery, University of Rochester Medical Center, Rochester, NY 14642, USA

^hDepartment of Orthopedic Surgery, Washington University School of Medicine, St. Louis, MO 63110, USA

Received 14 November 2013; revised 9 May 2014; accepted 12 May 2014

Author disclosures: JAG (none); HM (nonfinancial support from Spinal Deformity Study Group, during the conduct of the study; grants from Scoliosis Research Society, grants from Pediatric Orthopedic Society of North America, grants from Children's Spine Foundation (Grant no.: CWSD005, CWSD0004, CWSD0022, CWSD0026, CWSD0049), grants from Cerebral Palsy International Research Foundation (CPIRF Grant no.: R-808-12), grants from AOSpine, outside the submitted work); NDC (Spinal Deformity Study Group, during the conduct of the study); DPR (Spinal Deformity Study Group, during the conduct of the study; nonfinancial support from Stryker, grants from Scoliosis Research Society, grants from Pediatric Orthopaedic Society of North America, grants from Cerebral Palsy International Research Foundation, other from International Society of Orthopaedic Surgery and Traumatology, nonfinancial support from Children's Spine Foundation (Grant no.: CWSD005, CWSD0004, CWSD0022, CWSD0026, CWSD0049), grants from Chest Wall and Spine Deformity Research Foundation, grants from AOSpine, grants from Orthopaedic Research and Education Foundation, grants and nonfinancial support from Medtronic, grants from OMeGA (Grant no.: 001167, nosh5532NO, 000786), grants from Biomet, nonfinancial support from Broadwater [Biomet, Synthes, Stryker, Medtronic, K2], outside the submitted work); DJS (none); BSR (Wolters Kluwer Health—Lippincott Williams & Wilkins, other from Pfizer, outside the submitted work; and board member/committee member, Scoliosis Research Society; Medical/Orthopaedic Publications Editorial Governing Board, *Journal of Pediatric Orthopaedics*); JBE (Medtronic spine, other from Synthes spine, outside the submitted work); MAE (none); JOS (Grants from Children's Spine Foundation (Grant no.: CWSD005, CWSD0004, CWSD0022,

CWSD0026, CWSD0049), grants from POSNA, grants from National Institute of Arthritis and Musculoskeletal and Skin Diseases (Grant no.: 2R01AR052113), outside the submitted work; stock owner of Abbott Labs, GE, Hospira, and Abbvie); LGL (Grants from Axial Biotech, grants from DePuy-Synthes, grants from AOSpine/SRS/Norton Healthcare, other from AOSpine, other from John and Marcella Fox Research Agreement, personal fees from Medtronic, personal fees from K2M, personal fees from DePuy-Synthes Spine, personal fees from Medtronic, personal fees from Quality Medical Publishing, outside the submitted work); MG (Spinal Deformity Study Group, during the conduct of the study; other from AAP Section on Orthopaedics, personal fees, nonfinancial support and other from CWSDSG, personal fees from Stryker, personal fees from Biomet, grants from AOSpine, grants from Children's Spine Foundation (Grant no.: CWSD005, CWSD0004, CWSD0022, CWSD0026, CWSD0049), grants from Orthopaedic Research and Education Foundation, grants from SRS, grants and other from POSNA, grants and nonfinancial support from Medtronic, grants from OMeGA, nonfinancial support from Broadwater [Biomet, Syntes, Stryker, Medtronic, K2], nonfinancial support from FoxPSDSG, outside the submitted work).

This study was performed through the Spinal Deformity Study Group (SDSG), which was funded by Medtronic Sofamor Danek.

*Corresponding author. Division of Pediatric Orthopedic Surgery, Department of Orthopedic Surgery, Columbia University Medical Center, New York, NY 10032, USA. Tel.: (212) 305-5028; fax: (212) 305-8271.

E-mail address: hm2174@columbia.edu (H. Matsumoto).

Abstract

Study Design: Retrospective review of multicenter data set with adolescent idiopathic scoliosis (AIS) patients with at least 2 years of follow-up after posterior spinal instrumentation and fusion (PSIF).

Objectives: The purpose of this study is to investigate risk factors for coronal decompensation 2 years after PSIF for AIS.

Summary of Background Data: Coronal decompensation is a potential complication of spinal instrumentation for AIS. This can result in problems requiring revision surgery.

Methods: Demographic, clinical, and radiographic measures were reviewed on 890 identified patients. Coronal decompensation was defined as a change farther away from midline from 6 weeks postoperatively to 2 years in any one of the following radiographic parameters: change in coronal balance > 2 cm; change in coronal position of the lowest instrumented vertebra (LIV) > 2 cm; change in thoracic trunk shift > 2 cm; or change in LIV tilt angle > 10°. Patients with decompensation were compared to those without. The relationship between the LIV and lowest end vertebra (LEV) was examined as an independent variable.

Results: Two years postoperation, 6.4% (57/890) of patients exhibited coronal decompensation. Multivariate regression revealed that decompensated patients were twice as likely to be male, have lower preoperative Risser score, and lower percentage major curve correction. The relationship between the LIV and LEV as well as quality of life surveys were not significantly different between decompensated and nondecompensated patients at 2 years.

Conclusions: Two years after PSIF, 6.4% of patients with AIS exhibit radiographic coronal decompensation. Although this study did not demonstrate a significant association between the relationship of LIV and LEV and decompensation 2 years postoperation, results of this study indicate that skeletal immaturity, male gender, and less correction of the major curve may be related to higher rates of coronal decompensation.

© 2014 Scoliosis Research Society.

Keywords: Adolescent idiopathic scoliosis; Coronal balance; Posterior spinal instrumentation and fusion; Decompensation

Introduction

Segmental instrumentation using pedicle screws has become the standard of care for surgical treatment of adolescent idiopathic scoliosis (AIS). Although posterior spinal instrumentation and fusion (PSIF) techniques allow dramatic correction of multidimensional spinal deformities, there can be unintended effects on spinal alignment in both the coronal and sagittal planes. Poor spinal alignment directly resulting from surgery, or decompensation that develops postoperatively over time, may lead to pain, further decompensation, or other problems that require revision surgery [1]. As such, the characteristics and risk factors for spinal malalignment and decompensation must be further elucidated in order to assist surgeons in their preoperative planning to achieve optimal spinal alignment in scoliosis surgical correction.

The propensity of the unfused spine to respond to and change after the correction of coronal scoliotic deformity has been consistently demonstrated by the observation that nonstructural lumbar deformities in the coronal plane spontaneously correct after selective instrumentation and correction of the thoracic deformity [2–7]. On the basis of this concept and the presence of postoperative decompensation, it is now clear that these changes in coronal alignment may develop over time after PSIF for AIS. As such, coronal decompensation is a recognized potential complication that may develop after surgical correction of scoliosis. Several hypotheses have been reported in attempts to understand the etiology of decompensation. Coronal decompensation has been associated with the Cotrel-Dubousset rod derotation maneuver and a hypercorrection of the main thoracic curve, with management ranging from

observation or bracing, to revision surgery [8–10]. Postoperative coronal decompensation in specific curve types, such as Lenke 3C, has also been correlated to derotation of lumbar apical vertebrae [11]. Despite the wide attention to correction of coronal deformity, there is inconsistency in the literature concerning the definition of coronal decompensation, and investigation of its associated risk factors is incomplete.

It is the purpose of this study to investigate risk factors for coronal decompensation 2 years after PSIF for AIS. The term *coronal decompensation* is used in this study to describe a worsening of coronal spinal alignment that develops over time after spinal fusion, rather than a static measure of coronal alignment at any specific point in time.

Materials and Methods

Study design

After institutional review board approval from the senior author's institution (MGV), a survey was conducted of the prospective multicenter database of the Spinal Deformity Study Group (SDSG) for AIS patients (10–18 years at diagnosis). There were 21 centers who contributed patients to this study. On obtaining their consent, patients in this cohort were enrolled into the database between 2002 and 2009 and had their 2-year follow-up visit between 2004 and 2011. The database identified 2,833 patients with AIS who underwent PSIF during the period with average of 3.3 years of follow-up. Out of 2,833 patients, 1,766 patients had minimum of 2-year follow-up. There were 890 (152 males, 738 females) patients with any Lenke type curve who had PSIF with a minimum 2 years of complete clinical and

Download English Version:

<https://daneshyari.com/en/article/4095408>

Download Persian Version:

<https://daneshyari.com/article/4095408>

[Daneshyari.com](https://daneshyari.com)