

Clinical Study

Incidence, impact, and risk factors of adverse events in thoracic and lumbar spine fractures: an ambispective cohort analysis of 390 patients

R. Andrew Glennie, MD, FRCSC^a, Tamir Ailon, MD, MPH, FRCSC^a,
Kyun Yang, MD^a, Juliet Batke, MSc^a, Charles G. Fisher, MD^a, Marcel F. Dvorak, MD^a,
Alexander R. Vaccaro, MD, PhD^{b,c}, Michael G. Fehlings, MD, PhD, FRCSC^d,
Paul Arnold, MD^e, James S. Harrop, MD^{b,c}, John T. Street, MD, PhD, FRCSI^{a,*}

^aCombined Neurosurgical and Orthopedic Spine Program, Blusson Spinal Cord Center, University of British Columbia, 818 West 10th Ave., Vancouver, British Columbia, Canada V5Z1M9

^bDepartment of Orthopedic Surgery, Thomas Jefferson University, 925 Chestnut Street, Philadelphia, PA 19107, USA

^cDepartment of Neurological Surgery, Thomas Jefferson University, 925 Chestnut Street, Philadelphia, PA 19107, USA

^dDepartment of Surgery and Spinal Program, University of Toronto and University Health Network, 585 University Ave. Toronto, Ontario M5G2C4, Canada

^eDepartment of Neurosurgery, University of Kansas Medical Center, 3901 Rainbow Blvd., Kansas City, KS 66160, USA

Received 18 April 2014; revised 15 October 2014; accepted 21 November 2014

Abstract

BACKGROUND CONTEXT: Adverse events (AEs) in thoracic and lumbar spine fractures are common, but little is known about the type of AEs that are specific to this population. Furthermore, very little is known about the incidence and clinical impact of these AEs on patients in the presence of traumatic spinal cord injury and whether they are treated operatively or nonoperatively.

PURPOSE: The purpose of this study was to determine primarily the incidence of AEs in patients with thoracic or lumbar spine fractures treated both operatively and nonoperatively and their impact on length of stay (LOS) and secondarily the difference in the incidence of AEs in both neurologically intact and compromised patients.

STUDY DESIGN/SETTING: This is an ambispective cohort study at a quaternary referral center.

FDA device/drug status: Not applicable.

Author disclosures: **RAG:** Nothing to disclose. **TA:** Nothing to disclose.

KY: Nothing to disclose. **JB:** Research Support (Investigator Salary, Staff/Materials): Medtronic (F, Paid directly to institution); Fellowship Support: Medtronic (D, Paid directly to institution). **CGF:** Royalties: Medtronic (F); Consulting: Medtronic (F), Nuvasive (F); Grants: OREF (D); Fellowship Support: Medtronic (F, Paid directly to institution), AO Spine (F, Paid directly to institution). **MFD:** Royalties: Medtronic (F); Consulting: Medtronic (E); Speaking and/or Teaching Arrangements: Medtronic (F); Trips/Travel AO spine International (F), Medtronic (F); Endowments: Chair in Spinal Cord Injury Research (E, Paid directly to institution); Grants: Rick Hansen Institute (H, Paid directly to institution); Fellowship support: Medtronic (F, Paid directly to institution), Depuy Synthesis (F, Paid directly to institution), AO Spine (F, Paid directly to institution). **ARV:** Royalties: DePuy (C), Medtronic (H), Stryker Spine (G), Biomet Spine (C), Globus (F), Nuvasive (B), Aesculap (B); Stock Ownership: Replication Medica (15,000 shares, B), Globus (123,398 shares, L), K-2 Medical, Paradigm Spine (97,500 units, F), Stout Medical (1% company, E), Spine Medica (25,000 stock options, value unknown), Computational Biodynamics (50% ownership, value unknown), Progressive Spinal Technologies (30% ownership, value unknown), Spinology (8,125 shares, value unknown), Small Bone Innovations (15,000 shares, value unknown), Cross Current (62,500 shares, D), Syndicom (2,750 shares, B), In Vivo (123,935 shares, value unknown), Flagship Surgical (D), Advanced Spinal Intellectual Properties (30% ownership, value unknown), Cytonics (25,000 shares, value unknown), Bonovo Orthopedics (100,000 shares, paid F), Electrocore (50,000 shares, value unknown), Gamma Spine (15% ownership, value unknown), Location-Based Intelligence (20% ownership, value

unknown), Flowpharma (nonqualified stock options 200,000, value unknown), R.S.I. (50% ownership, value unknown), Rothman Institute and Related Properties (practice, value unknown), Innovative Surgical Design (30% ownership, value unknown), Spinicity (53,000 shares, 3.4% ownership); Consulting: Gerson Lehrman Group (B), Guidepoint Global (B), Medacorp (B), Stout Medical (B), Innovative Surgical Design (B), Orthobullets (A); Board of Directors: AO Spine, Innovative Surgical Design, Association of Collaborative Spine Research, Spinicity; Grants: Stryker Spine, Cerapedics, Nuvasive (C). **MGF:** Nothing to disclose. **PA:** Stock Ownership: Z-plasty (A); Consulting: Medtronic, Lifespine (A), Integra life (B), Spinewave, Stryker Spine (C), Fziomed (B), MIEMS (B), AO Spine North America (B), Cerapedics-past relationship (B); Speaking and/or Teaching Arrangements: University of Missouri (A); Board of Directors: AO Spine North America-past relationship; Grants: AO Spine North America (E). **JSH:** Consulting: Depuy Spine (D, Paid directly to institution); Scientific Advisory Board/Other Office: Tejin (B); Scientific Advisory Board/Other Office: Bioventus (B). **JTS:** Speaking and/or Teaching Arrangements: Medtronic (B); Research Support (Investigator Salary, Staff/Materials): Medtronic (F, Paid directly to institution); Fellowship Support: Medtronic (D, Paid directly to institution).

The disclosure key can be found on the Table of Contents and at www.TheSpineJournalOnline.com.

No funding was received or used in this study.

* Corresponding author. Combined Neurosurgical and Orthopedic Spine Program, Blusson Spinal Cord Center, University of British Columbia, 818 West 10th Ave., Vancouver, British Columbia, Canada V5Z1M9. Tel.: (1) 604-875-5328; fax: (1) 604-875-8223.

E-mail address: john.street@vch.ca (J.T. Street)

PATIENT SAMPLE: Patients admitted at our institution with thoracic or lumbar fractures from January 2009 to December 2013 were identified. Patients with full Spine Adverse Events Severity System (SAVES) data were included.

OUTCOME MEASURES: Number and type of AEs collected from SAVES were assessed. Impact of AE on acute LOS was also determined.

METHODS: Data on intraoperative, preoperative, and postoperative AEs were prospectively collected using the SAVES data collection. Logistic regression was used to model the likelihood of experiencing at least one AE based on the patient characteristics. The impact of the total number of AEs experienced by a patient and that of each of the most common AEs on LOS was determined using Poisson regression.

RESULTS: Three hundred and ninety patients were included in the final analysis. Two hundred and seventy-six patients (70.8%) were treated operatively. One hundred and forty patients (36%) experienced neurologic deficit as a result of their initial injury. Adverse events occurred 56% of the time in the operatively treated patients and only 13% of the time in the nonoperative group. The presence of neurologic deficit increased the risk of AEs especially in high thoracic (T1–T6) trauma increasing the odds of having an AE by 12.1 ($p < .0001$). The most common AEs were urinary tract infections (19.7%), neuropathic pain (12.3%), pneumonias (11.8%), delirium (10.5%), and ileus (6.2%). Length of hospital stay increased significantly with pneumonia ($p < .0001$) and delirium ($p = .0001$).

CONCLUSIONS: The presence of neurologic injury and the need for operative fixation of thoracic or lumbar injuries lead to a greater risk of AEs. Only pneumonia and delirium consistently increase LOS. © 2015 Elsevier Inc. All rights reserved.

Keywords:

Thoracic; Lumbar; Adverse event; Operative; Traumatic spinal cord injury; Spine

Introduction

Thoracic and lumbar spine traumas encompass a wide variety of injury patterns from low-energy vertebral compression fractures to high-energy fracture dislocations. A great deal has been published on the management of thoracic and lumbar spine traumas and the associated wide spectrum of injuries. This is reflected in the variability of analyses and publications on the subject matter. The majority of the literature focuses on a number of critical areas: appropriately classifying different injury patterns, operative versus nonoperative treatment, timing of operative intervention, operative techniques for fixation of unstable fractures, radiographic outcomes, and health-related quality of life (QOL) [1–6]. There is a paucity of literature, however, specifically about adverse events (AEs) in thoracic and lumbar traumas.

Operative and nonoperative management of these injuries has been debated over numerous decades. Classically, these fractures were initially treated nonoperatively with bed rest and body casting [7]. Practices have changed over time with improvements in operative management, spinal instrumentation, and evidence demonstrating a significant morbidity associated with prolonged immobilization. Surgeons transitioned into treating a greater number of fractures operatively with the rationale that this would lead to better early pain relief, better fracture reduction, early ambulation, and an overall improvement in clinical outcome [8].

More recently, the evidence for these concepts is being challenged with large series demonstrating equivalent clinical

outcomes in fractures treated nonsurgically [9,10]. In those injury patterns that are considered “stable,” most will be treated nonsurgically and will focus on early ambulation with appropriate analgesia [11]. However, there is great difficulty in defining “stable”; thus, presently there is considerable effort invested in identifying those fractures that will fail nonoperative management and require surgical stabilization [2,12]. Clinical outcomes and QOL have been compared for operative and nonoperative groups. Very little is known, however, about any differences that may exist with regard to AE rates in patients treated surgically or nonsurgically [13–15].

Fortunately, most of these injuries will not result in neurologic injury. Larger trials have reported a 10% to 25% incidence of neurologic deficit [16,17]. The majority of these patients will be treated operatively because of ongoing compression of the neural elements, residual instability, and malalignment of the spinal column. Spinal cord injury (SCI) will often translate to a higher incidence of AEs with some authors quoting as high as 77% [18,19]. When only considering thoracic or lumbar traumatic SCI, however, there is less known about the incidence of AEs that are specific to this group of patients. For example, it is debated whether thoracic and lumbar SCI patients are susceptible to the same pulmonary complications that can be quite common with cervical SCI.

Identifying AEs is critical in this population and establishing predictive models to help potentially avoid these complications depending on which treatment patients receive. We present an analysis of thoracic and lumbar trauma patients undergoing treatment at a Level 1 trauma

Download English Version:

<https://daneshyari.com/en/article/4096348>

Download Persian Version:

<https://daneshyari.com/article/4096348>

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