Title

Opioid Use Prior to Adult Spine Deformity Correction Surgery is Associated With Worse Pre- and Postoperative Back Pain and Prolonged Opioid Demands.

Permalink

https://escholarship.org/uc/item/93d8t95t

Authors

Sardi, Juan
Smith, Justin
Gum, Jeffrey
et al.

Publication Date

2024-06-04

DOI

10.1177/21925682241261662

Peer reviewed

Opioid Use Prior to Adult Spine Deformity Correction Surgery is Associated With Worse Pre- and Postoperative Back Pain and Prolonged Opioid Demands

Juan P. Sardi, MD[1] , Justin S. Smith, MD, PhD[1] , Jeffrey L. Gum, MD[2] ,
Brett Rocos, MD[3] , Anastasios Charalampidis, MD, PhD[4,5] , Lawrence G. Lenke, MD[6],
Christopher I. Shaffrey, MD[3], Kenneth M. C. Cheung, MD[7,8] , Yong Qiu, MD[9],
Yukihiro Matsuyama, MD[10], Ferran Pellisé, MD[11] , David W. Polly JR, MD[12,13],
Jonathan N. Sembrano, MD[12,13], Benny T. Dahl, MD[13], Michael P. Kelly, MD[14],
Marinus de Kleuver, MD[15], Maarten Spruit, MD[16], Ahmet Alanay, MD[17],
Sigurd H. Berven, MD[18], Stephen J. Lewis, MD[19] , and AO Spine Knowledge Forum
Deformity

Abstract

Study Design: Prospective multicenter database post-hoc analysis.

Objectives: Opioids are frequently prescribed for painful spinal conditions to provide pain relief and to allow for functional improvement, both before and after spine surgery. Amidst a current opioid epidemic, it is important for providers to understand the impact of opioid use and its relationship with patient-reported outcomes. The purpose of this study was to evaluate

[1] Department of Neurosurgery, University of Virginia, Charlottesville, VA, USA
[2] Norton Leatherman Spine Center, Louisville, KY, USA
[3] Departments of Neurosurgery and Orthopedic Surgery, Duke University, Durham, NC, USA
[4] Department of Reconstructive Orthopaedics, Karolinska University Hospital, Stockholm, Sweden
[5] Department of Clinical Science, Intervention and Technology (CLINITEC), Karolinska Institutet, Stockholm, Sweden
[6] Department of Orthopedic Surgery, Columbia University, New York, NY, USA
[7] Department of Orthopaedic & Traumatology, The University of HK, Hong Kong
[8] The HKU-Shenzhen Hospital, Shenzhen China
[9] The Affiliated Drum Tower Hospital of Nanjing University Medical School, Nanjing, China
[10] Department of Orthopaedic Surgery, Hamamatsu University School of Medicine, Shizuoka, Japan
[11] Spine Surgery Unit, Vall d'Hebron Hospital, Barcelona, Spain
[12] University of Minnesota, Minneapolis, MN, USA
[13] Texas Children's Hospital, Houston, TX, USA
[14] Rady Children's Hospital, San Diego, CA, USA
[15] Department of Orthopedic Surgery, VU University Medical Center, Amsterdam Movement Sciences, Amsterdam, The Netherlands
[16] St. Maartenskliniek, Nijmegen, Netherlands
[17] Department of Orthopedics and Traumatology, Acibadem Mehmet Ali Aydınlar University School of Medicine. Istambul, Turkey
[18] University of California San Francisco Spinal Disorders Service, San Francisco, CA, USA
[19] Toronto Western Hospital, Toronto, ON, Canada

Corresponding Author:
Juan P. Sardi, MD, Department of Neurosurgery, University of Virginia Health Sciences Center, PO Box 800212, Charlottesville, VA 22908, USA.
Email: js9yu@uvahealth.org

pre-/postoperative opioid consumption surrounding ASD and assess patient-reported pain outcomes in older patients undergoing surgery for spinal deformity.

Methods: Patients ≥60 years of age from 12 international centers undergoing spinal fusion of at least 5 levels and a minimum 2-year follow-up were included. Patient-reported outcome scores were collected using the Numeric Rating Scale for back and leg pain (NRS-B; NRS-L) at baseline and at 2 years following surgery. Opioid use, defined based on a specific question on case report forms and question 11 from the SRS-22r questionnaire, was assessed at baseline and at 2-year follow-up.

Result: Of the 219 patients who met inclusion criteria, 179 (81.7%) had 2-year data on opioid use. The percentages of patients reporting opioid use at baseline (n = 75, 34.2%) and 2 years after surgery (n = 55, 30.7%) were similar (*P* = .23). However, at last follow-up 39% of baseline opioid users (Opi) were no longer taking opioids, while 14% of initial non-users (No-Opi) reported opioid use. Regional pre- and postoperative opioid use was 5.8% and 7.7% in the Asian population, 58.3% and 53.1% in the European, and 50.5% and 40.2% in North American patients, respectively. Baseline opioid users reported more preoperative back pain than the No-Opi group (7.0 vs 5.7, *P* = .001), while NRS-Leg pain scores were comparable (4.8 vs 4, *P* = .159). Similarly, at last follow-up, patients in the Opi group had greater NRS-B scores than Non-Opi patients (3.2 vs 2.3, *P* = .012), but no differences in NRS-Leg pain scores (2.2 vs 2.4, *P* = .632) were observed.

Conclusions: In this study, almost one-third of surgical ASD patients were consuming opioids both pre- and postoperatively world-wide. There were marked international variations, with patients from Asia having a much lower usage rate, suggesting a cultural influence. Despite both opioid users and nonusers benefitting from surgery, preoperative opioid use was strongly associated with significantly more back pain at baseline that persisted at 2-year follow up, as well as persistent postoperative opioid needs.

Introduction

Over the last decades, the use of medical opioids has increased by a factor of 10 and opioid-related deaths have nearly quadrupled.[1-3] Insurance data suggest that opioids are the most commonly prescribed drugs for back pain and more than half of routine opioid users report back pain.[4-6] The aging of the global population has led to an increased prevalence of adult spinal deformity (ASD), with reports as high as 68% among older adults.[7,8] Pain is the hallmark presenting symptom of ASD, it causes significant disability and has a negative impact on patient-reported quality of life. With pain manifesting in nearly 90% of these patients, most of them will eventually take opioids as part of either their initial non-operative treatment or after surgery, or both.[6,9] Moreover, substantial advances in surgical approaches, instrumentation, technology and critical care have helped broaden surgical indications, thus allowing for older and more frail patients to be considered for deformity correction surgery.[10] This growing number of elderly adults with painful spinal conditions and the parallel rise in surgical interventions have led to an over 600% increase in spine-related pain opioid expenditure over the past 20 years.[6,11-14]

It is known that in the acute postoperative setting opioids provide efficacious analgesia, which translates into faster rehabilitation and functional recovery.[15-17] However, optimal opioid utilization for subacute or chronic non-cancer pain is still debated, and non-scientific formulations have been shown to increase the risk of side effects and misuse.[3,18,19]

Furthermore, marketing campaigns such as "*Pain is the 5th vital sign*" have pushed a cultural perception demanding opioids, thus resulting in an on-going public health emergency.[20,21] This has allowed for increasing recognition of the risks of perioperative opioids including tolerance, respiratory depression, persistent opioid use and dependence. Therefore, strategies to mitigate opioid consumption, in conjunction with their deleterious side-effects and costs, are being sought.

There is particular interest in establishing ideal preoperative opioid usage since despite a growing cognizance of higher morbidity and worse surgical outcomes associated with their use, these medications are still widely prescribed.[22,23] Additionally, there are inherent cultural differences regarding the perception of opioid utilization regardless of indication. The purpose of this study was to evaluate pre- and postoperative opioid consumption surrounding ASD and assess the effects of pre-operative opioid use on patient-reported pain scores at 2 years following multi-level spinal deformity surgery in older patients.

Materials and Methods

Study Design and Inclusion Criteria

Patient records were obtained from a prospective, multicenter, international cohort of patients (Prospective Evaluation of Elderly Deformity Surgery -PEEDS database), an AO-Spine

funded study to assess operative treatment and outcomes of elderly patients undergoing spine deformity surgery.[24] The original prospective, randomized trial received institutional review/ethics board approval at all 12 participating sites across North America, Asia and Europe and was then registered at clinicaltrials.gov (Identifier: NCT02035280). Written informed consent was obtained upon enrollment for all patients in the original study; however, the present study was approved for a Waiver of Consent, granting exemption from informed consent requirement, due to the retrospective nature of data collection. All cases were patients aged ≥60 years undergoing primary spinal fusion surgery of ≥5 levels for a coronal, sagittal or combined deformity who were capable and willing to sign the consent (No objective values to define deformity were given). Patients meeting inclusion criteria had outcome forms completed at baseline, 10 weeks, 12 months and 24 months.

Data Collection and Statistical Analysis

We retrospectively extracted data that were prospectively collected to gather patient demographic data, surgical information, and clinical characteristics. Reported outcome scores included the Numeric Rating Scale for back and leg pain (NRS-B; NRS-L; range from 0 to 10, with 0 being no pain and 10 corresponding to the most severe pain) and opioid use was collected from standardized case report forms that queried medication use and question 11 from the SRS 22r questionnaire: "Which one of the following best describes your pain medication use for back pain?" Possible choices were: (1) none; (2) non-narcotics weekly or less; (3) non-narcotics daily; (4) narcotics weekly or less; or (5) narcotics daily.[25] Patients were categorized dichotomously as any opioid use (Opi) or no-opioid use (No-Opi) and surgical centers were divided into North America, Europe and Asia.

The primary objectives of this study were to assess the effects of pre-operative opioid use on patient-reported pain scores at baseline and at 2 years after spine deformity correction surgery, as well as to assess international differences regarding pre- and post-operative opioid consumption. We compared reported outcome scores of patients receiving preoperative opioids to those who were not receiving opioids both at baseline and at 2-year follow-up. We also assessed changes in pain scores after surgery among these same groups and calculated least square differences with 95% confidence interval. Finally, we compared opioid use at baseline and at 2 years among patients from North America, Europe, and Asia, and used the NRS leg and back pain as well as the SRS 22-r satisfaction domain scores to assess for any significant regional differences. To detect significant differences between the groups and to compare pre/post-op results, Wilcoxon rank-sum test was performed. Means with standard deviations and medians with IQRs were used to describe continuous and ordinal variables, and frequencies with percentages were used for categorical variables. Differences were considered statistically significant for $P < .05$.

All statistical analyses were performed using SAS (version 9.4, SAS Institute Inc., Cary, NC, USA).

Results

A total of 219 patients met the inclusion criteria. Most patients were women (80.4%) and Caucasian (56.6%). The mean age at the time of surgery was 67.5 years (range: 60-83) and the median number of spinal levels fused was 9 (IQR: 5, 12) (Table 1). Preoperatively, 75 (34.2%) patients were using opioids, while 144 (65.8%) were opioid naïve. At 2-year follow-up, 179 patients (81.7%) had data on opioid use, 55 (30.7%) were still using opioids and 124 (69.3%) had stopped using opioids (Figure 1). Overall, the proportions of patients reporting opioid use at baseline and at 2 years after surgery were similar. Odds ratio for continued opioid use at 2 years postoperative was 9.65 (95% confidence interval = 4.65-20.00, $P < .001$) for opioid vs non-opioid users. Furthermore, persistent postoperative opioid need was seen in approximately two-thirds of the baseline users that were still present at last follow-up (39/64 patients) (Figure 1).

At last follow up, of the 75 patients in the Opi group, 63 had available NRS back pain scores and 61 had available NRS leg pain scores; and out of the 144 patients in the No-Opi group, 114 had data for NRS back and 113 for NRS leg pain. Baseline NRS back pain was 7.0 (SD 2.0) in the Opi group compared to 5.7 (SD 2.8) in the No-Opi group ($P = .001$). Baseline NRS leg was 4.8 (SD 3.4) in the Opi group compared to 4.0 (SD 3.3) in the No-Opi patients ($P = .159$) (Table 2). At 2-year follow-up, NRS back pain was 3.2 (SD 2.5) in the Opi group and 2.3 (SD 2.6) in the No-Opi group ($P = .012$), while NRS leg pain was 2.2 (SD 2.7) in the Opi group and 2.4 (SD 2.7) in the No-Opi group ($P = .632$) (Table 3). Pain scores significantly improved after surgery in both No-Opi (Δ NRS back pain score: −3.3 (SD 3.3) and Δ NRS Leg pain score of −1.7 (SD 3.8)) and Opi (Δ NRS back pain score: −3.6 (SD 2.5) and Δ NRS Leg pain score of −2.4 (SD 3.8)) groups, however the magnitudes of improvement were not significantly different between the groups (Table 4).

Regarding regional differences, 5.8% and 7.7% of patients from the Asia group were using opioids pre- and postoperatively, respectively. In contrast, 58.3% of European and 50.5% of North American patients were taking opioids at baseline and 53.1% and 40.2% were taking them postoperatively, respectively. Patients taking opioids at baseline in the North American cohort had worse baseline NRS-Back (6.6 vs 5.0, $P = .003$), 2-year NRS-Back (3.3 vs 1.4, $P = .001$) and NRS-Leg (2.6 vs 1.0, $P = .007$) scores than No-Opi users. Similarly, the Opi group in the Asia cohort had worse baseline NRS-Leg scores (7.6 vs 4.2, $P = .023$), but otherwise there was no significant difference in baseline NRS-Back or 2-year NRS-Back or NRS-Leg scores. There was no difference in NRS-Back or NRS-Leg for European patients at baseline or at 2 years. regardless of opioid use. The satisfaction component of the SRS 22-r demonstrated no statistically significant

Table 1. Baseline Demographic Variables of Patients Undergoing Spine Deformity Correction Surgery.

Characteristic	Total (N = 219)	No Preoperative Opioids (N = 144)	Preoperative Opioids (N = 75)	P Value
Age, mean (SD)	67.5 (5.4)	67.9 (5.6)	66.8 (4.9)	.179
Body Mass index, mean (SD)	26.1 (5.4)	25.5 (5.1)	27.2 (5.9)	**.021**
Female sex, n (%)	176 (80.4)	113 (78.5)	63 (84.0)	.328
Work status, n (%)				
Employed	23 (10.5)	13 (9.0)	10 (13.3)	.188
Self-employed	11 (5.0)	7 (4.9)	4 (5.3)	
Unemployed	11 (5.0)	4 (2.8)	7 (9.3)	
Homemaker	34 (15.5)	25 (17.4)	9 (12.0)	
Retired	132 (60.3)	88 (61.1)	44 (58.7)	
Not aplicable	8 (3.7)	7 (4.9)	1 (1.3)	
Preoperative radiographic parameters				
Pelvic incidence, mean (SD)	51.4 (12.0)	51.2 (11.5)	51.9 (12.9)	.715
Pelvic tilt, mean (SD)	17.0 (8.8)	16.6 (8.7)	17.7 (9.1)	.453
Sacral slope, mean (SD)	34.4 (11.0)	34.6 (11.1)	34.2 (10.8)	.839
Lumbar lordosis, mean (SD)	48.4 (14.4)	48.9 (14.8)	47.5 (13.8)	.534
SVA, mean (SD)	32.0 (42.8)	25.7 (38.6)	43.4 (47.7)	**.012**
T1PA, mean (SD)	14.2 (8.8)	13.5 (8.7)	15.5 (8.9)	.181
PI-LI missmatch, mean (SD)	3.1 (13.5)	2.3 (13.8)	4.4 (12.8)	.32
Coronal balance (absolute value), mean (SD)	−2.6 (28.1)	−2.1 (27.6)	−3.6 (29.1)	.746
Coronal lumbar cobb angle (absolute value), mean (SD)	14.1 (13.4)	12.4 (11.4)	16.5 (15.7)	.103
Coronal thoracic cobb angle (absolute value), mean (SD)	13.0 (11.9)	11.8 (9.6)	15.1 (15.2)	.288
Surgical characteristics				
Number of levels fused, mean (SD)	10.8 (3.9)	10.5 (3.8)	11.3 (4.1)	.177
Surgical time, mean minutes (SD)	406.6 (112.7)	410.6 (111.9)	399.0 (114.7)	.474
Blood loss, mean mL (SD)	1684.2 (1185.7)	1652.3 (1272.5)	1745.6 (1003.3)	.553
Three column osteotomies, mean (SD)	29 (18.6)	18 (18.2)	11 (19.3)	.863

[a]p < 0.05.
Note. Statistical significant with p < 0.05.

differences between No-Opi and Opi groups in the European (4.4 vs 4.5, $P = .8$) and Asian (3.5 vs 3.7, $P = .645$) populations, however No-Opi users were more satisfied after surgery than Opi users in the North American population (4.5 vs 4.1, $P = .003$) (Table 5).

Discussion

Impact of Opioid Use on Clinical Outcomes

In this multicentric, multicontinental study of patients ≥60 years undergoing elective spine deformity correction surgery, we found that nearly one-third of patients were consuming opioids pre- and postoperatively world-wide. This is consistent with opioid prescription data in national registries, where 38% of adult patients are found to use opioids at some point during the 12 months prior to any surgical intervention.[26] Previous reports have shown that regular opioid use before surgery is one of the strongest predictors for poorly controlled postoperative pain and is the single most influential predictive factor for opioid refills.[27,28] When compared to

opioid naïve patients, even minimal preoperative opioid use has been reported to increase the likelihood of requiring prescription refills after surgery (OR 1.49, 95% CI 1.45 – 1.53; $P < .001$) and an even stronger association is seen if these medications are taken chronically (OR 60.79, 95% CI 27.81–132.92, $P < .001$).[26] Furthermore, given that nearly 20% of opioid users undergoing spine surgery may be opioid-dependent, high rates of chronic postoperative opioid use are to be expected after these interventions.[29-31]

Our results indicated that odds ratio for continued opioid use at 2 years postoperative was 9.65 (95% confidence interval = 4.65-20.00, $P < .001$) for opioid vs non-opioid users. These findings are supported by previous reports also demonstrating the strong association between opioid use prior to spine surgery and their continuation in the postoperative period.[3,22,23,29,31-34] However, we observed, that despite 33.3% (25 patients) of baseline opioid users being able to stop taking opioids by the last follow-up, the percentage of patients overall still depending on them 2 years after surgery was considerably high (30.7%). Since only 16 of the initial non-opioid users were taking these medications long after surgery,

nearly 70% of the sustained postoperative opioid use observed in our study largely resulted from the high incidence of preoperative use itself (34.2%) (Figure 1). Unlike most ASD studies that usually include revision surgeries, all our patients underwent primary interventions, which makes the incidence of preoperative opioid use strikingly high.

Prolonged postoperative opioid use might suggest surgery failed to provide the expected results in terms of symptomatic relief. However, we observed that pain scores significantly improved after surgery in both opioid users (Δ NRS back pain

score: −3.6 and Δ NRS Leg pain score of −2.4) and non-users (Δ NRS back pain score: −3.3 and Δ NRS Leg pain score of −1.7) (Table 4). Pain and disability are characteristically what lead patients with ASD to seek surgical treatment and there is substantial evidence supporting significant improvement in pain, HRQoL outcomes, patient function and satisfaction after surgery.[7,12,13,24,35] Nevertheless, the majority of these patients have struggled with chronic pain long before surgery and have likely exhausted non-operative interventions, including conventional analgesics and opioids. These chronic painful conditions and their subsequent opioid exposure have been shown to alter pain perception pathways with unintended consequences like allodynia, opioid-induced hyperalgesia (OIH), opioid tolerance and withdrawal-associated hyperalgesia (WAH).[36] Therefore, despite successful surgical interventions, it is possible that these patients may experience higher levels of postoperative pain, increased opioid demands and pose a considerable challenge when trying to wean them off opioids, when postoperative pain should theoretically be controlled.

This distorted nociceptive sensitization may potentially explain why patients in the opioid group, who were presumably receiving "stronger" opioid-based preoperative pain regimes, had significantly more back pain at baseline compared to patients who were not using opioids (NRS Back pain 7.0 [SD 2.0] vs 5.7 [SD 2.8], P = .001). However, while leg pain scores were similar in those with and without preoperative opioid use (NRS Leg pain 4.8 [SD 3.4] vs 4.0 [SD 3.3], P = .159), ASD can present with both back and radicular pain, but there is usually a predominant and most debilitating symptom that directs the invasiveness of the procedure.[37-39] Since our patients consisted of coronal and sagittal deformities requiring multilevel instrumentation, their main concern was axial mechanical back pain, thus reflected in the substantial difference between back- and leg pain intensity scores across all patients. Except for the SVA, all spinopelvic parameters
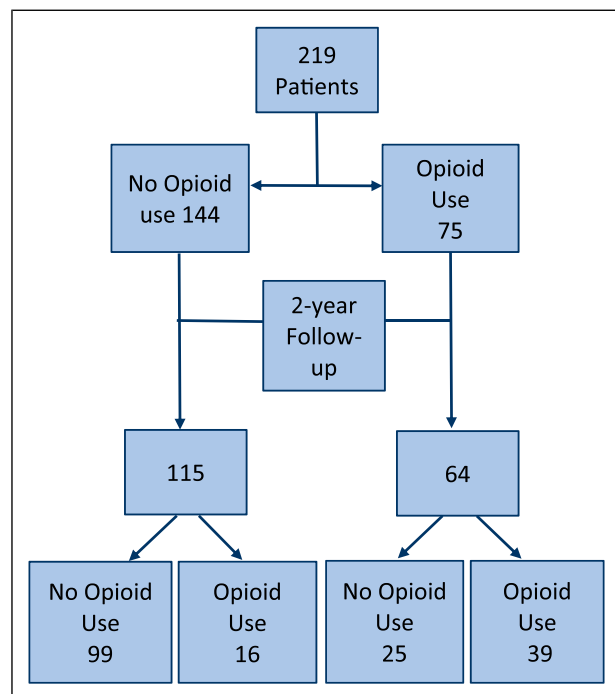

Figure 1. Distribution of patients reporting opioid intake at baseline and at 2-year follow-up.

Table 2. Pain NRS Data at Baseline by Opioid Use Derived From SRS-22r Questionnaire and Prescribed Drugs Statement.

	Baseline Opioid Use Based on SRS-22r Questionnaire and Prescribed Drugs Statement			
	No Opioid Use[a]	Opioid Use	Total	
Variable	N = 144	N = 75	N = 219	P Value
NRS leg pain at preoperative				.159[b]
n	143	72	215	
Mean (SD)	4.0 (3.3)	4.8 (3.4)	4.3 (3.3)	
Median (Q1; Q3)	3.0 (1.0; 7.0)	5.0 (1.0; 7.5)	4.0 (1.0; 7.0)	
NRS back pain at preoperative				.001[b]
n	144	74	218	
Mean (SD)	5.7 (2.8)	7.0 (2.0)	6.1 (2.7)	
Median (Q1; Q3)	6.0 (3.0; 8.0)	8.0 (6.0; 8.0)	7.0 (5.0; 8.0)	

[a]Includes 3 patients for whom question 11 of the SRS-22r was not answered at baseline although they declared no prescribed drugs.
[b]Wilcoxon rank sum test.

Table 3. Pain NRS Data at 2-Year Follow-Up by Opioid Use Derived From SRS-22r Questionnaire and Prescribed Drugs Statement.

	Baseline Opioid Use Based on SRS-22r Questionnaire and Prescribed Drugs Statement			
	No Opioid Use[a]	Opioid Use	Total	
Variable	N = 144	N = 75	N = 219	P Value
NRS leg pain at 2 Years				.632[b]
n	113	61	174	
Mean (SD)	2.4 (2.7)	2.2 (2.7)	2.3 (2.7)	
Median (Q1; Q3)	1.0 (.0; 4.0)	1.0 (.0; 4.0)	1.0 (.0; 4.0)	
NRS back pain at 2 Years				.012[b]
n	114	63	177	
Mean (SD)	2.3 (2.6)	3.2 (2.5)	2.6 (2.6)	
Median (Q1; Q3)	1.5 (.0; 4.0)	3.0 (1.0; 5.0)	2.0 (.0; 5.0)	

[a]Includes 3 patients for whom question 11 of the SRS-22r was not answered at baseline although they declared no prescribed drugs.
[b]Wilcoxon rank sum test.

Table 4. Change in Pain NRS Scores at 2-Year Follow-Up by Opioid Use Derived From SRS-22r Questionnaire and Prescribed Drugs Statement.

	Baseline Opioid Use Based on SRS-22r Questionnaire and Prescribed Drugs Statement							
	No Opioid Use			Opioid Use			Total	
	N = 144			N = 75			N = 219	
Variable	Preoperative	2 years	Change in Pain (Δ NRS)	Preoperative	2 years	Change in Pain (Δ NRS)	LS-Mean Difference in Narcotic Use (95% CI)	P Value
NRS leg pain							−.7 (−1.9; .5)	.266[a]
Mean (SD)	4.0 (3.3)	2.4 (2.7)	−1.7 (3.8)	4.8 (3.4)	2.2 (2.7)	−2.4 (3.8)		
Median (Q1; Q3)	3.0 (1.0; 7.0)	1.0 (.0; 4.0)	−2.0 (−4.0; .5)	5.0 (1.0; 7.5)	1.0 (.0; 4.0)	−2.0 (−5.0; .0)		
NRS back pain							−.3 (−1.3; .6)	.483[a]
Mean (SD)	5.7 (2.8)	2.3 (2.6)	−3.3 (3.3)	7.0 (2.0)	3.2 (2.5)	−3.6 (2.5)		
Median (Q1; Q3)	6.0 (3.0; 8.0)	1.5 (.0; 4.0)	−4.0 (−6.0; −1.0)	8.0 (6.0; 8.0)	3.0 (1.0; 5.0)	−3.0 (−5.0; −2.0)		

[a]Wilcoxon rank sum test.

were similar among both groups, indicating comparable magnitudes of sagittal and coronal deformities between users and non-users (Table 1). Similar results were reported by Line et al[40] where a propensity score matched analysis of 262 operatively treated ASD patients demonstrated greater baseline NRS back- (7.7 vs 6.8) and leg pain (2.5 vs 2.3) in patients using opioids before surgery as compared to non-opioid users. In their study, both groups of patients had comparable deformity magnitudes, comorbidity burden, history of mental illness and surgical invasiveness.

Interestingly, despite substantial symptomatic relief indicating both groups benefitted from surgery, at last follow-up there was still a statistically significant difference in NRS back pain scores between the Opioid- and No-Opioid groups (3.2 [SD 2.5] vs 2.3 [SD 2.6], $P = .012$). These findings are consistent with those by Hills et al[34] where preoperative opioid therapy was associated with significantly higher odds of not achieving a clinically meaningful improvement at 1 year in extremity pain (aOR, 1.55; 95% CI, 1.21-1.99; $P = .001$) and axial pain (aOR, 1.73; 95% CI, 1.37-2.18; $P < .001$). However, unlike our study where opioid use encompassed any opioid consumption, they only included chronic users (having an active prescription >50% of the days in each month for 3 consecutive months prior to surgery) of which 73% were on

Table 5. Opioid Use With Patient Satisfaction and Pain by Region Derived From SRS-22r Questionnaire and Prescribed Drugs Statement.

	Baseline Opioid Use			2 Years Opioid Use		
	No Opioids	Opioids	P Value	No Opioids	Opioids	P Value
Asia N (%)	81 (94.2%)	5 (5.8%)		60 (92.3%)	5 (7.7%)	
SRS-22r Satisfaction score, mean (SD)	3.1 (.7)	3.3 (1.5)	.956[a]	3.7 (.9)	3.5 (1.3)	.8[a]
NRS leg pain score, mean (SD)	4.2 (3.2)	7.6 (2.1)	.023[a]	2.8 (2.6)	4.2 (3.6)	.334[a]
NRS back pain score, mean (SD)	5.8 (2.9)	7.4 (1.9)	.271[a]	2.8 (2.6)	3.4 (3.2)	.674[a]
Europe N (%)	15 (41.7%)	21 (58.3%)		15 (46.9%)	17 (53.1%)	
SRS-22r Satisfaction score, mean (SD)	3.0 (1.7)	3.7 (1.0)	.566[a]	4.5 (.6)	4.4 (.6)	.645[a]
NRS leg pain score, mean (SD)	4.7 (3.6)	4.7 (3.8)	1.000[a]	2.4 (3.2)	3.3 (2.4)	.23[a]
NRS back pain score, mean (SD)	7.2 (2.1)	7.8 (1.3)	.545[a]	2.7 (2.8)	4.1 (2.1)	.191[a]
North America N (%)	48 (49.5%)	49 (50.5%)		49 (59.8%)	33 (40.2%)	
SRS-22r Satisfaction score, mean (SD)	3.2 (1.0)	2.8 (.9)	.077[a]	4.5 (.8)	4.1 (.8)	.003[a]
NRS leg pain score, mean (SD)	3.6 (3.3)	4.5 (3.2)	.177[a]	1.0 (1.9)	2.6 (3.0)	.007[a]
NRS back pain score, mean (SD)	5.0 (2.8)	6.6 (2.2)	.003[a]	1.4 (2.1)	3.3 (2.6)	.001[a]

[a]Wilcoxon rank sum test.

high opioid doses (daily morphine milligram equivalents >30). This might suggest that chronicity and dosage have an impact on outcomes and should also be considered prior to surgery. There are also numerous publications involving smaller spine surgeries, including decompressions, cervical and lumbar short-segment fusions and even minimally invasive techniques, in which opioids also were shown to have a negative effect on outcomes.[3,21,30,33,41,42] Our results add to the growing body of literature suggesting a negative impact of preoperative opioid use on postoperative outcomes that seems to exists regardless of the invasiveness of the procedure.

Furthermore, preoperative opioid use has been shown to impact other outcome measures besides pain scores. In a systematic review by Yerneni et al[31] that included 45 studies, opioid use was a negative predictor of return to work status, hospital stay, healthcare costs, wound complications and risk of revision surgery. Other studies have also shown a negative impact over ICU length of stay, overall postoperative complications, the 12-item Short Form Health Survey, Oswestry Disability Index and Neck Disability index.[23,28,43] There are no definitive explanations as to why preoperative opioid use is associated with worse clinical outcomes, and though it is probably multifactorial, the intrinsic hyperalgesia induced by opioids likely plays a critical role. In light of our results, one could make the argument that preoperative opioid use might be one of the most prevalent modifiable risk factors and that discontinuing or reducing opioid medications prior to ASD correction surgery could lead to better clinical outcomes. Potentially, detoxification and interdisciplinary collaborations could allow for a safe and tolerable transition to non-opioid alternatives and improved preoperative management. However, further research is warranted to recognize if reversing opioid use could lead to better outcomes after spine surgery and if so, establishing proper pathways to achieve this goal.

Our results also highlight the impact of sustained postoperative opioid use. Similar to our findings in which 14% of opioid naïve patients reported still using opioids at last follow-up, a study by Deyo et al[44] that included nearly 2500 patients undergoing spine surgery found that 12.8% of preoperative non-opioid users were also utilizing long-term opioids for pain control. Effective perioperative pain management can significantly improve functional recovery and rehabilitation after spine surgery, hence pain control is a fundamental aspect of the postoperative care.[15-17] However, though opioids are effective analgesics in the acute setting, subacute and chronic use remain controversial and have not been shown to be superior to non-opioid formulations in terms of pain relief, safety and functional outcomes.[15-17,28,45] Bearing in mind indiscriminate use of opioids and long-term consumption increases the risks of side effects, misuse, dependence, abuse and health-related costs, numerous clinical practice guidelines are now recommending against their use and suggest that alternate strategies for pain control should be considered.[3,18,19,46] Multimodal pain management should target reduced opioid use in the perioperative period, ideally reserving opioids only as rescue analgesia; and in cases of opioid-dependent or opioid-tolerant patients, medications like gabapentin, pregabalin and intraoperative ketamine infusions show promising opioid-sparing effects with significant reduced postoperative opioid consumption after spine surgery.[47-51]

Regional Differences in Perioperative Opioid Use

To our knowledge, this is the first report comparing regional differences in opioid use before- and after spine surgery, between North American, European, and Asian populations. While out of the 3 patient groups Asians reported significantly less pre- and postoperative opioid use, the percentage of

opioid users seemed to slightly increase after surgery (5.8% vs 7.7%). However, the total number of opioid users in the Asian patients remained unchanged (5) and this increase resulted from 21 nonusers being lost during follow-up. In contrast, postoperative opioid use in the North American and European patients decreased after surgery from 50.5% to 40.2% and 58.3% to 53.1% respectively, which suggests that surgery provided enough symptomatic improvement that allowed for opioid discontinuation. Much like the results of our data, and consistent with previous publications, subgroup analysis revealed that the vast majority of postoperative opioid users result from persistent preoperative use.

There are few reports in the literature addressing opioid use in spine surgery among European patients, but the high proportion of opioid users observed in our study contrasts significantly with previous published data. An observational study of by Holmberg et al[52] that included over 30 000 patients with degenerative lumbar spine disorders from 2 Norwegian registries found that only 8.7% of patients were using opioids in the year previous to their surgery, however similar to our results nearly two-thirds of these patients continued to use opioids 2 years postoperatively. In 2 other multicentric studies published by the European Spine Study Group, preoperative opioid use was reported to be 18% and 24.2%.[53,54] We believe these reports may underrepresent the true prevalence of opioid use in European patients undergoing spine surgery mainly because of how opioid consumption was defined. In the Norwegian paper opioid use was defined as 180 Defined Daily Doses (DDDs) or >4500 oral morphine equivalents (OMEQ) for 365 days, with prescriptions dispensed in 3 of 4 quarters of the year; while Bourghli et al[54] defined opioid use as daily narcotic consumption and patients using opioids on a weekly basis or less were included within the No-painkiller/Minor user group. Even minimal opioid consumption impacts postoperative opioid use and "minor" use should not be overlooked when a patients is deemed candidate for any spine surgical intervention.

Interestingly, despite having the highest percentage of baseline users, there was no difference in NRS-B or NRS-L scores for European patients at baseline or at 2 years following surgery, regardless of opioid use. Asian patients taking opioids at baseline did have worse mean NRS-L scores (7.6 vs 4.2, $P = .023$) but otherwise there was no difference in baseline NRS-B or 2-year NRS-B or NRS-L scores. These regional results differ from our pooled data and from previous publications in which baseline opioid use is associated with worse baseline and postoperative pain scores. We believe this might be secondary to variations in intake chronicity, dosage and opioid potency among populations. In a study by Lee et al[43] multivariate analyses demonstrated that increased preoperative opioid use was a significant predictor of decreased SF-12 PCS and MCS and increased ODI scores postoperatively; in fact, every increase in 10-mg morphine equivalent amount taken preoperatively predicted a .3 decrease in SF-12 and a .6 increase in ODI

postoperatively ($P < .05$). This is consistent with findings by Holmberg et al[52] where European patients receiving homogenously high opioid doses reported larger mean ODI scores at baseline than nonusers (51.0 points vs 41.9 points, $P < 0.001$) and larger mean NRS scores for both back pain and leg pain in the year following surgery. Local regulations and health system demands likely cause significant differences in opioid prescriptions between regions which were not discerned by the questionnaires used in our study.

Though prescription and intake dosages in our study were not specified, opioid formulation trends in Europe appear to be more conservative than in North America. Abul et al[53] reported that 96% of opioid users from 5 cites across Europe were found to be receiving weak opioids prior to spine deformity correction surgery, while in a North American based study, Hills et al[34] identified high-preoperative opioid dosage (>30 MME/d) in 73% of patients. These differences might explain why opioid use in North American patients was associated with worse baseline NRS-B (6.6 vs 5.0, $P = .003$) and 2-yr NRS-B (3.3 vs 1.4, $P = .001$) and NRS-L (2.6 vs 1.0, $P = .007$). Also, this might explain why there was no statistically significant difference in satisfaction between nonusers and opioid users in the European (4.4 vs 4.5, $P = .8$) and Asian (3.5 vs 3.7, $P = .645$) populations, while North American nonusers were more satisfied after surgery than opioid users (4.5 vs 4.1, $P = .003$). The amount, potency and chronicity of opioid consumption that leads to chronic postoperative use and significant changes in clinical outcomes remain unknown and further studies will be needed to assess this.

Limitations

Since our study was a post-hoc analysis, the definition of "spine deformity" relied on the inclusion criteria used by the authors of the original paper. Unfortunately, Nielsen et al[24] did not specify the objective coronal nor sagittal values used to define deformity, but rather included patients who underwent multilevel lumbar fusion >5 levels. Furthermore, one of the major limitations we encountered was the lack of specific data regarding opioid dosing before and after surgery. This limited the ability to evaluate dose reduction after surgery, considered an indirect marker of success; and from properly assessing the impact of chronicity and the dose-dependent effect over clinical outcomes and postoperative opioid dependence. Since this was a multicentric transcontinental study, regional differences between pre- and postoperative pain management likely resulted in heterogeneous formulations that were reflected in our results among the different populations. Studies with structured protocols are needed to adequately interpret these variations. Also, the retrospective nature of the design does not allow us to draw conclusions about causality, despite the evident association between preoperative opioid use and dependence as well as over clinical outcomes. Finally, since our study relied on patient questionnaires there was a potential for recall bias that in future research can be addressed using

prescription data which will also add granularity to the amounts and time of consumption.

Conclusions

The present study indicates that almost one-third of surgical ASD patients were consuming opioids both pre- and post-operatively world-wide; there is a drastic international variation, with Asia having a much lower usage rate, suggesting a cultural influence. Despite both users and nonusers benefitting from surgery and experiencing significant improvement in pain scores, our results support the growing evidence that preoperative opioid use is strongly associated with persistent postoperative opioid consumption. Furthermore, preoperative opioid users present with significantly more back pain at baseline that persists to at least 2 years following surgery compared to non-opioid users, highlighting the concerning difficulty of discontinuing opioid analgesics in patients who have been using them prior to their intervention. Preoperative opioid consumption may be a potentially modifiable risk factor to promote improved clinical outcomes for ASD surgery. Further studies are needed to determine if individualized preventative approaches and opioid-sparing perioperative management may lead to reduced postoperative opioid utilization and improved clinical outcomes.

Authors Contribution

All authors listed contributed sufficiently to the project to be included as authors, and all those who are qualified to be authors are listed in the author byline.

Declaration of Conflicting Interests

The author(s) declared no potential conflicts of interest with respect to the research, authorship, and/or publication of this article.

ORCID iDs

Juan P. Sardi ⓘ https://orcid.org/0000-0002-0696-9213
Justin S. Smith ⓘ https://orcid.org/0000-0003-0467-5534
Jeffrey L. Gum ⓘ https://orcid.org/0000-0003-0471-9437
Brett Rocos ⓘ https://orcid.org/0000-0002-0808-5585
Anastasios Charalampidis ⓘ https://orcid.org/0000-0001-5228-738X
Kenneth M. C. Cheung ⓘ https://orcid.org/0000-0001-8304-0419
Ferran Pellisé ⓘ https://orcid.org/0000-0002-0644-7757
Stephen J. Lewis ⓘ https://orcid.org/0000-0002-9173-8443

References

1. Okie S. A flood of opioids, a rising tide of deaths. *N Engl J Med*. 2010;363(21):1981-1985. doi:10.1056/NEJMp1011512
2. Paulozzi LJ, Ryan GW. Opioid analgesics and rates of fatal drug poisoning in the United States. *Am J Prev Med*. 2006;31(6): 506-511. doi:10.1016/j.amepre.2006.08.017
3. Kalakoti P, Hendrickson NR, Bedard NA, Pugely AJ. Opioid utilization following lumbar arthrodesis: trends and factors associated with long-term use. *Spine (Phila Pa 1976)*. 2018; 43(17):1208-1216. doi:10.1097/brs.0000000000002734
4. Ivanova JI, Birnbaum HG, Schiller M, Kantor E, Johnstone BM, Swindle RW. Real-world practice patterns, health-care utilization, and costs in patients with low back pain: the long road to guideline-concordant care. *Spine J*. 2011;11(7):622-632. doi:10.1016/j.spinee.2011.03.017
5. Hudson TJ, Edlund MJ, Steffick DE, Tripathi SP, Sullivan MD. Epidemiology of regular prescribed opioid use: results from a national, population-based survey. *J Pain Symptom Manage*. 2008;36(3):280-288. doi:10.1016/j.jpainsymman.2007.10.003
6. Deyo RA, Korff MV, Duhrkoop D. Opioids for low back pain. *BMJ Br Med J (Clin Res Ed)*. 2015;350:g6380. doi:10.1136/bmj.g6380
7. Smith JS, Shaffrey CI, Glassman SD, et al. Risk-benefit assessment of surgery for adult scoliosis: an analysis based on patient age. *Spine (Phila Pa 1976)*. 2011;36(10):817-824. doi:10.1097/BRS.0b013e3181e21783
8. Schwab F, Dubey A, Gamez L, et al. Adult scoliosis: prevalence, SF-36, and nutritional parameters in an elderly volunteer population. *Spine (Phila Pa 1976)*. 2005;30(9):1082-1085. doi:10.1097/01.brs.0000160842.43482.cd
9. Lo YT, Lim-Watson M, Seo Y, et al. Long-term opioid prescriptions after spine surgery: a meta-analysis of prevalence and risk factors. *World Neurosurg*. 2020;141:e894-e920. doi:10.1016/j.wneu.2020.06.081
10. Yadla S, Maltenfort MG, Ratliff JK, Harrop JS. Adult scoliosis surgery outcomes: a systematic review. *Neurosurg Focus*. 2010; 28(3):E3. doi:10.3171/2009.12.Focus09254
11. Kostuik JP, Israel J, Hall JE. Scoliosis surgery in adults. *Clin Orthop Relat Res*. 1973;(93):225-234. doi:10.1097/00003086-197306000-00022
12. Birknes JK, White AP, Albert TJ, Shaffrey CI, Harrop JS. Adult degenerative scoliosis: a review. *Neurosurgery*. 2008;63(3 Suppl):94-103. doi:10.1227/01.NEU.0000325485.49323.B2
13. Youssef JA, Orndorff DO, Patty CA, et al. Current status of adult spinal deformity. *Global Spine J*. 2013;3(1):51. doi:10.1055/s-0032-1326950
14. Martin BI, Deyo RA, Mirza SK, et al. Expenditures and health status among adults with back and neck problems. *JAMA*. 2008; 299(6):656-664. doi:10.1001/jama.299.6.656

15. Bajwa SJ, Haldar R. Pain management following spinal surgeries: an appraisal of the available options. *J Craniovertebr Junction Spine*. 2015;6(3):105-110. doi:10.4103/0974-8237.161589

16. Borgeat A, Blumenthal S. Postoperative pain management following scoliosis surgery. *Curr Opin Anaesthesiol*. 2008;21(3):313-316. doi:10.1097/ACO.0b013e3282f82baa

17. Lamperti M, Tufegdzic B, Avitsian R. Management of complex spine surgery. *Curr Opin Anaesthesiol*. 2017;30(5):551-556. doi:10.1097/aco.0000000000000494

18. Chang AK, Bijur PE, Esses D, Barnaby DP, Baer J. Effect of a single dose of oral opioid and nonopioid analgesics on acute extremity pain in the emergency department: a randomized clinical trial. *JAMA*. 2017;318(17):1661-1667. doi:10.1001/jama.2017.16190

19. da Costa BR, Nüesch E, Kasteler R, et al. Oral or transdermal opioids for osteoarthritis of the knee or hip. *Cochrane Database Syst Rev*. 2014. 2014(9):Cd003115. doi:10.1002/14651858.CD003115.pub4

20. US Office of National Drug Control Policy. The President's commission on combating drug addiction and the Opioid crisis. https://www.whitehouse.gov/sites/whitehouse.gov/files/images/Final_Report_Draft_11-1-2017.pdf. Accessed 23 October 2023.

21. Wang MC, Lozen AM, Laud PW, Nattinger AB, Krebs EE. Factors associated with chronic opioid use after cervical spine surgery for degenerative conditions. *J Neurosurg Spine*. 2020;32(1):1-8. doi:10.3171/2019.7.spine19563

22. Jain N, Phillips FM, Weaver T, Khan SN. Preoperative chronic opioid therapy: a risk factor for complications, readmission, continued opioid use and increased costs after one- and two-level posterior lumbar fusion. *Spine (Phila Pa 1976)*. 2018;43(19):1331-1338. doi:10.1097/brs.0000000000002609

23. Raad M, Jain A, Neuman BJ, et al. Association of patient-reported narcotic use with short- and long-term outcomes after adult spinal deformity surgery: multicenter study of 425 patients with 2-year follow-up. *Spine (Phila Pa 1976)*. 2018;43(19):1340-1346. doi:10.1097/brs.0000000000002631

24. Nielsen CJ, Lewis SJ, Oitment C, et al. Stratifying outcome based on the Oswestry Disability Index for operative treatment of adult spinal deformity on patients 60 years of age or older: a multicenter, multi-continental study on Prospective Evaluation of Elderly Deformity Surgery (PEEDS). *Spine J*. 2021;21(11):1775-1783. doi:10.1016/j.spinee.2021.07.007

25. Inclan P, CreveCoeur TS, Bess S, et al. SRS-22r question 11 is a valid opioid screen and stratifies opioid consumption. *Spine Deform*. 2022;10(4):913-917. doi:10.1007/s43390-022-00473-0

26. Vu JV, Cron DC, Lee JS, et al. Classifying preoperative opioid use for surgical care. *Ann Surg*. 2020;271(6):1080-1086. doi:10.1097/sla.0000000000003109

27. Kampe S, Wendland M, Welter S, et al. Independent predictors for higher postoperative pain intensity during recovery after open thoracic surgery: a retrospective analysis in 621 patients. *Pain Med*. 2018;19(8):1667-1673. doi:10.1093/pm/pnx238

28. Yang M, Riva-Cambrin J, Cunningham J. Development and validation of a clinical prediction score for poor postoperative pain control following elective spine surgery. *J Neurosurg Spine*. 2020;34:3.

29. Dunn LK, Yerra S, Fang S, et al. Incidence and risk factors for chronic postoperative opioid use after major spine surgery: a cross-sectional study with longitudinal outcome. *Anesth Analg*. 2018;127(1):247-254. doi:10.1213/ane.0000000000003338

30. Walid MS, Hyer L, Ajjan M, Barth AC, Robinson JS Jr. Prevalence of opioid dependence in spine surgery patients and correlation with length of stay. *J Opioid Manag*. 2007;3(3):127-128, 130-2. doi:10.5055/jom.2007.0050

31. Yerneni K, Nichols N, Abecassis ZA, Karras CL, Tan LA. Preoperative opioid use and clinical outcomes in spine surgery: a systematic review. *Neurosurgery*. 2020;86(6):E490-E507. doi:10.1093/neuros/nyaa050

32. Schoenfeld AJ, Belmont PJ Jr., Blucher JA, et al. Sustained preoperative opioid use is a predictor of continued use following spine surgery. *J Bone Joint Surg Am*. 2018;100(11):914-921. doi:10.2106/jbjs.17.00862

33. Lawrence JT, London N, Bohlman HH, Chin KR. Preoperative narcotic use as a predictor of clinical outcome: results following anterior cervical arthrodesis. *Spine (Phila Pa 1976)*. 2008;33(19):2074-2078. doi:10.1097/BRS.0b013e3181809f07

34. Hills JM, Pennings JS, Archer KR, et al. Preoperative opioids and 1-year patient-reported outcomes after spine surgery. *Spine (Phila Pa 1976)*. 2019;44(12):887-895. doi:10.1097/brs.0000000000002964

35. Berven SH, Kamper SJ, Germscheid NM, et al. An international consensus on the appropriate evaluation and treatment for adults with spinal deformity. *Eur Spine J*. 2018;27(3):585-596. doi:10.1007/s00586-017-5241-1

36. Tompkins DA, Campbell CM. Opioid-induced hyperalgesia: clinically relevant or extraneous research phenomenon? *Curr Pain Headache Rep*. 2011;15(2):129-136. doi:10.1007/s11916-010-0171-1

37. Heary RF, Kumar S, Bono CM. Decision making in adult deformity. *Neurosurgery*. 2008;63(3 Suppl):69-77. doi:10.1227/01.NEU.0000320426.59061.79

38. Pritchett JW, Bortel DT. Degenerative symptomatic lumbar scoliosis. *Spine*. 1993;18(6):700-703. doi:10.1097/00007632-199305000-00004

39. Liu W, Chen XS, Jia LS, Song DW. The clinical features and surgical treatment of degenerative lumbar scoliosis: a review of 112 patients. *Orthop Surg*. 2009;1(3):176-183. doi:10.1111/j.1757-7861.2009.00030.x

40. Line B, Bess S, Gum JL, et al. Opioid use prior to surgery is associated with worse preoperative and postoperative patient reported quality of life and decreased surgical cost effectiveness for symptomatic adult spine deformity; A matched cohort analysis. *N Am Spine Soc J*. 2022;9:100096. doi:10.1016/j.xnsj.2021.100096

41. Villavicencio AT, Nelson EL, Kantha V, Burneikiene S. Prediction based on preoperative opioid use of clinical outcomes

after transforaminal lumbar interbody fusions. *J Neurosurg Spine*. 2017;26(2):144-149. doi:10.3171/2016.7.SPINE16284

42. Kha ST, Scheman J, Davin S, Benzel EC. The impact of preoperative chronic opioid therapy in patients undergoing decompression laminectomy of the lumbar spine. *Spine (Phila Pa 1976)*. 2020;45(7):438-443. doi:10.1097/brs.0000000000003297

43. Lee D, Armaghani S, Archer KR, et al. Preoperative opioid use as a predictor of adverse postoperative self-reported outcomes in patients undergoing spine surgery. *J Bone Joint Surg Am*. 2014;96(11):e89. doi:10.2106/jbjs.M.00865

44. Deyo RA, Hallvik SE, Hildebran C, et al. Use of prescription opioids before and after an operation for chronic pain (lumbar fusion surgery). *Pain*. 2018;159(6):1147-1154. doi:10.1097/j.pain.0000000000001202

45. Busse JW, Craigie S, Juurlink DN, et al. Guideline for opioid therapy and chronic noncancer pain. *CMAJ*. 2017;189(18):E659-e666. doi:10.1503/cmaj.170363

46. Bernstein IA, Malik Q, Carville S, Ward S. Low back pain and sciatica: summary of NICE guidance. *BMJ*. 2017;356:i6748. doi:10.1136/bmj.i6748

47. Loftus RW, Yeager MP, Clark JA, et al. Intraoperative ketamine reduces perioperative opiate consumption in opiate-dependent patients with chronic back pain undergoing back surgery. *Anesthesiology*. 2010;113(3):639-646. doi:10.1097/ALN.0b013e3181e90914

48. Waelkens P, Alsabbagh E, Sauter A, Joshi GP, Beloeil H. Pain management after complex spine surgery: a systematic review and procedure-specific postoperative pain management recommendations. *Eur J Anaesthesiol*. 2021;38(9):985-994. doi:10.1097/eja.0000000000001448

49. Pacreu S, Fernández Candil J, Moltó L, Carazo J, Fernández Galinski S. The perioperative combination of methadone and ketamine reduces post-operative opioid usage compared with methadone alone. *Acta Anaesthesiol Scand*. 2012;56(10):1250-1256. doi:10.1111/j.1399-6576.2012.02743.x

50. Urban MK, Ya Deau JT, Wukovits B, Lipnitsky JY. Ketamine as an adjunct to postoperative pain management in opioid tolerant patients after spinal fusions: a prospective randomized trial. *HSS J*. 2008;4(1):62-65. doi:10.1007/s11420-007-9069-9

51. Yu L, Ran B, Li M, Shi Z. Gabapentin and pregabalin in the management of postoperative pain after lumbar spinal surgery: a systematic review and meta-analysis. *Spine (Phila Pa 1976)*. 2013;38(22):1947-1952. doi:10.1097/BRS.0b013e3182a69b90

52. Holmberg ST, Skurtveit S, Gulati S, et al. Persistent use of prescription opioids before and after lumbar spine surgery: observational study with prospectively collected data from two Norwegian national registries. *Spine*. 2023;48(14):969-977. doi:10.1097/brs.0000000000004710

53. Abul K, Yilgor C, Yucekul A, et al. Long-term opioid medication profile of European adult spinal deformity patients: minimum five years follow-up study. *Spine J*. 2023;23(2):209-218. doi:10.1016/j.spinee.2022.10.017

54. Bourghli A, Boissiere L, Larrieu D, et al. Opioids and analgesics use after adult spinal deformity surgery correlates with sagittal alignment and preoperative analgesic pattern. *Eur Spine J*. 2020;29(1):73-84. doi:10.1007/s00586-019-06141-8