



7-1-2016

Adverse Drug Effects and Preoperative Medication Factors Related to Perioperative Low-Dose Ketamine Infusions.

Eric S. Schwenk

Thomas Jefferson University, eric.schwenk@jefferson.edu

Stephen F. Goldberg

Thomas Jefferson University, Stephen.Goldberg@jefferson.edu

Ronak D. Patel

Thomas Jefferson University

Jon Zhou

University of California, Davis

Douglas R. Adams

*Thomas Jefferson University, Douglas.Adams@jefferson.edu**See next page for additional authors*

[Let us know how access to this document benefits you](#)

Follow this and additional works at: <https://jdc.jefferson.edu/anfp> Part of the [Anesthesiology Commons](#)

Recommended Citation

Schwenk, Eric S.; Goldberg, Stephen F.; Patel, Ronak D.; Zhou, Jon; Adams, Douglas R.; Baratta, Jaime L.; Viscusi, Eugene R.; and Epstein, Richard H., "Adverse Drug Effects and Preoperative Medication Factors Related to Perioperative Low-Dose Ketamine Infusions." (2016). *Department of Anesthesiology Faculty Papers*. Paper 33.

<https://jdc.jefferson.edu/anfp/33>

This Article is brought to you for free and open access by the Jefferson Digital Commons. The Jefferson Digital Commons is a service of Thomas Jefferson University's Center for Teaching and Learning (CTL). The Commons is a showcase for Jefferson books and journals, peer-reviewed scholarly publications, unique historical collections from the University archives, and teaching tools. The Jefferson Digital Commons allows researchers and interested readers anywhere in the world to learn about and keep up to date with Jefferson scholarship. This article has been accepted for inclusion in Department of Anesthesiology Faculty Papers by an authorized administrator of the Jefferson Digital Commons. For more information, please contact: JeffersonDigitalCommons@jefferson.edu.

Authors

Eric S. Schwenk, Stephen F. Goldberg, Ronak D. Patel, Jon Zhou, Douglas R. Adams, Jaime L. Baratta, Eugene R. Viscusi, and Richard H. Epstein

Adverse Drug Effects and Preoperative Medication Factors Related to Perioperative Low-Dose Ketamine Infusions

Corresponding Author:

Eric S. Schwenk, MD
Sidney Kimmel Medical College, Thomas Jefferson University
Department of Anesthesiology
Suite 8130, Gibbon Building
111 South 11th Street
Philadelphia, PA 19107
Phone: 215-955-6161
Fax: 215-955-0677
Email: Eric.Schwenk@jefferson.edu

Co-Authors:

Stephen F. Goldberg, MD
Sidney Kimmel Medical College, Thomas Jefferson University

Ronak Patel, MD
Thomas Jefferson University Hospital

Jon Zhou, MD
University of California, Davis

Douglas R. Adams, BS
Sidney Kimmel Medical College, Thomas Jefferson University

Jaime L. Baratta, MD
Sidney Kimmel Medical College, Thomas Jefferson University

Eugene R. Viscusi, MD
Sidney Kimmel Medical College, Thomas Jefferson University

Richard H. Epstein, MD
Sidney Kimmel Medical College, Thomas Jefferson University

Institutional Affiliation of Manuscript:

Sidney Kimmel Medical College, Thomas Jefferson University

Source of Funding:

Departmental funding

Portions of this manuscript were presented at the 2013 annual meeting of the American Society of Anesthesiologists and at the 2014 spring meeting of the American Society of Regional Anesthesia and Pain Medicine.

Conflicts of Interest:

Eric Schwenk, Stephen Goldberg, Ronak Patel, Jon Zhou, Douglas Adams, Jaime Baratta, and Richard Epstein have no conflicts of interest to declare.

Eugene Viscusi has served as a consultant for AcclRx, Medicines Company, Mallinkrodt, Trevena, Cara Pharmaceuticals, and Astra Zeneca. He has received grant money in the past from AcclRx and Pacira. He has been a paid lecturer for AcclRx, Merck, Salix, and Mallinkrodt.

1 **Abstract**

2 High-dose opioid administration is associated with significant adverse events.
3 Evidence suggests that low-dose ketamine infusions improve perioperative analgesia over
4 conventional opioid management, but usage is highly variable. Ketamine's adverse drug
5 effects (ADEs) are well known, but their prevalence during low-dose infusions in a
6 clinical setting and how often they lead to infusion discontinuation are unknown. The
7 purposes of this study were threefold: 1) to identify patient factors associated with
8 initiation of ketamine infusions during spine surgery; 2) to identify specific spine
9 procedures in which ketamine has been used most frequently; and 3) to identify ADEs
10 associated with postoperative ketamine infusions and which ADEs most frequently led to
11 discontinuation. Spine surgery was chosen because of its association with moderate to
12 severe pain and a relatively high use of ketamine infusions in this population at our
13 hospital.

1 **Introduction**

2 Patients presenting for surgery due to conditions associated with chronic pain
3 frequently are being treated with opioids, often at alarmingly high doses. One study of
4 Medicaid enrollees found that 63.5% of patients with non-cancer chronic pain had taken
5 an opioid in the prior 12 months, an increase of 18.9% from five years prior.¹ Yet despite
6 escalating doses of opioids, patients continue to report that their chronic pain is not well
7 controlled.² At the same time, serious adverse drug effects (ADEs) associated with
8 opioids, including fatal respiratory depression, continue to be a serious concern.³ In
9 patients chronically taking opioids, tolerance to respiratory depression is incomplete⁴ and
10 such patients have an increased risk of overdose and death compared to non-opioid
11 users.⁵ Analgesic alternatives to opioids are, therefore, highly desirable.

12 One such alternative during the perioperative period is ketamine, an
13 N-methyl-D-aspartate (NMDA) receptor antagonist. A role of the NMDA receptor in the
14 development of opioid tolerance was suggested by studies from several decades ago.^{6,7}
15 Ketamine is a potent analgesic that does not cause respiratory depression, and may
16 improve postoperative analgesia while reducing opioid consumption.^{8,9} As an additional
17 potential benefit, recent evidence suggests that intravenous (IV) ketamine may decrease
18 the incidence of persistent postsurgical pain (PPSP).¹⁰ Ketamine infusions at our
19 institution have been used both intraoperatively by the anesthesia team and
20 postoperatively by the acute pain management service (APMS) as part of an
21 opioid-sparing strategy in complex, opioid-tolerant patients.

22 We had three primary objectives of the current study: 1) to identify factors
23 associated with current decisions by anesthesiologists to initiate ketamine during spine

1 surgery; 2) to identify specific **spine** procedures in which ketamine has been used most
2 frequently; and 3) to identify ADEs associated with **all** postoperative ketamine infusions
3 and which ADEs most frequently led to discontinuation. This information is necessary
4 for us to design prospective, randomized clinical trials comparing IV ketamine to placebo
5 in opioid-tolerant patients. Spine surgery was chosen to study in detail because it is
6 typically associated with moderate to severe postoperative pain, over 50% of our
7 ketamine use has been in spine surgery, and this is a high-volume service at our
8 institution. We retrospectively analyzed the preoperative medications and surgical details
9 of all patients over a three-year period who underwent any type of elective spine surgery
10 **and also examined a sample of postoperative patients from all surgical specialties who**
11 **received ketamine infusions for the presence of ADEs and classified them.**

12

1 **Methods**

2
3 This study was approved by the Thomas Jefferson University institutional review
4 board without requirement for written patient consent.

5 All patients admitted on the day of surgery following elective spine surgery by an
6 orthopedic surgeon or neurosurgeon under general anesthesia between January 1, 2012
7 and March 21, 2015 at Thomas Jefferson University Hospital (TJUH) were studied.
8 TJUH is a major academic medical center, as well as a regional spinal cord trauma center
9 **with a high spine surgery volume.** Patients undergoing microdiscectomy at our hospital
10 are usually cared for on an outpatient basis, and therefore most of those patients were not
11 included, **since their post-operative data were not available.** Demographic data,
12 preoperative medications, and dose and timing data related to ketamine infusions were
13 retrieved from the hospital's anesthesia information management system (Innovian[®],
14 Dräger, Telford, PA) and from the pharmacy information system database (Pyxis[®],
15 CareFusion, San Diego, CA). Patients who received intraoperative ketamine boluses but
16 not an infusion were excluded. Data elements analyzed included the date of surgery, age
17 in years, gender, weight, body mass index, American Society of Anesthesiologists (ASA)
18 physical status, scheduled duration of surgery, primary surgical service, and preoperative
19 medications. Preoperative opioids were classified as being taken on a "**scheduled**" or "as
20 needed" basis (Table 2). Planned procedures (using locally defined, procedure-specific
21 codes) were queried from the operating room case scheduling system (ORSOS[®],
22 McKesson, San Francisco, CA). **All planned, elective spine surgeries requiring hospital
23 admission (but not emergencies or cases booked as "add-ons") were included for
24 analysis. The decision to start an intraoperative ketamine infusion at our hospital was**

1 made by the attending anesthesiologist for the case. Data retrieved from the pharmacy
2 information system were aligned with patient anesthesia records to determine if
3 postoperative ketamine infusions had been started intraoperatively or initiated after the
4 patient left the OR.

5 ADE data were retrieved from the daily notes recorded contemporaneously by the
6 Acute Pain Management Service (APMS) nurses on a consecutive sample of 321 patients
7 who received a postoperative ketamine infusion while on the APMS from January 1,
8 2011 through December 31, 2013. Patients from all surgical subspecialties were included
9 for the ADE analysis, not just those undergoing spine procedures. All APMS nurses had
10 undergone training on the management of ketamine infusions, including the recognition
11 of side effects. No special monitoring, such as telemetry or intensive care, has been
12 required at our institution for patients receiving ketamine. Criteria for discontinuation of
13 ketamine infusions included the patient requesting discontinuation due to ADEs or the
14 patient's primary service requesting discontinuation.

15 Data were extracted and prepared for analysis using SQL Server 2008 R2
16 (Microsoft, Redmond, WA). Odds ratios (OR) were computed using the function
17 *oddsRatio* in the R *mosaic* library, Pearson's chi-square test (with Yate's continuity
18 correction), two-group Student t tests (with the Satterthwaite approximation), and local
19 polynomial regression fits using the functions *chisq.test*, *t.test*, and *loess*, respectively, in
20 the R *stats* library (R v3.2.0, The R Foundation for Statistical Computing, Vienna,
21 Austria).

22

23

1 **Results**

2 *Demographics and Preoperative Medications Associated with Ketamine Administration*

3 There were 4958 patients who underwent elective spine surgery under general
4 anesthesia during the study interval, 4748 of which were entered into our electronic
5 preoperative anesthesia system and had data available for analysis, and 211 of whom
6 received an intraoperative infusion of ketamine. Among all patients, those receiving
7 ketamine were younger (difference = -4.4 years, 95% CI -2.2 to -6.0 years, $P < 10^{-6}$), had
8 a higher ASA Physical Status ($P < 10^{-6}$), and were scheduled for surgeries of longer
9 estimated duration (difference = 72 minutes, 95% CI 60 to 84 minutes, $P < 10^{-6}$), (Table
10 1). There were no significant differences in weight or BMI between the two groups.
11 Males and females were represented equally (difference = -0.40%, 95% CI -3.1% to
12 2.3%, $P = 0.77$).

13 Medication factors at the time of the preoperative evaluation associated with a
14 greater likelihood of receiving an intraoperative ketamine infusion were taking vs. not
15 taking a scheduled opioid (OR 16.09, 95% CI 11.98 to 21.59), taking any opioid vs. no
16 opioid (OR 10.25, 95% CI 7.13 to 14.75), and taking vs. not taking an anti-depressant
17 (OR 2.69, 95% CI 2.02 to 3.57) (Table 2). Patients who were taking both a scheduled
18 opioid and an anti-depressant were more likely to receive ketamine than those taking just
19 a scheduled opioid (OR 1.64, 95% CI 1.11 to 2.46; Table 2).

20 Among the 552 patients who were taking a scheduled opioid at the time of the
21 preoperative evaluation, those receiving a ketamine infusion were younger (difference
22 = -2.8 years, 95% CI -0.6 to -5.1 years, $P = 0.012$), had a higher ASA Physical Status (P
23 = 0.01), and were scheduled for surgeries of longer estimated duration (difference = 49

1 minutes, 95% CI 32 to 67 minutes, $P < 10^{-6}$), (Table 3). There were no significant
2 differences in weight or BMI between the two groups. There was a higher proportion of
3 females (difference = 13.3%, 95% CI 4.3 to 22.3%, $P = 0.005$).

4

5 *Surgical Procedures Associated with Ketamine Administration*

6 There were 20 distinct spine procedures (12 primary spine procedures and 8
7 revision spine procedures) identified in the database (Table 4). Of these, there were 10
8 procedures that had $\geq 5\%$ prevalence of intraoperative ketamine administration. The three
9 most commonly performed of these 10 were posterior thoracic/lumbar fusion (N = 148
10 cases), anterior thoracic/lumbar fusion (N = 136 cases), and anterior/posterior cervical
11 fusion (N = 137 cases). Specific spine procedures are displayed in Table 4 according to
12 primary or revision status.

13

14 *Side Effects of Low-Dose Ketamine Infusions*

15 There were 31.8% of patients who experienced at least one ADE (Table 5). The
16 most frequent ADE was CNS excitation (16.2%), followed by sedation (9.4%) and visual
17 disturbances (3.1%). Some patients experienced more than one ADE (Table 5). Thirty-
18 seven patients (36.3% of all patients with an ADE) experienced ADEs severe enough to
19 have resulted in discontinuation of the ketamine infusion. The reasons for infusion
20 discontinuation are described in Table 5. Sedation was the ADE most likely to result in
21 ketamine discontinuation. Of the 37 patients whose infusions were discontinued, 35 of
22 them reported resolution of symptoms after the infusion was stopped. Twenty-six patients
23 received benzodiazepines, commonly used for treatment of side effects at our hospital,

1 while 11 patients did not. To view the TJUH ketamine infusion guidelines, see Appendix
2 A.

3 A postoperative infusion rate above 20 mg/hr was not associated with an
4 increased chance of having the infusion stopped compared to patients receiving ≤ 20
5 mg/hr (OR 0.71, 95% CI 0.34 to 1.5). The chance of discontinuation was also not
6 increased with a threshold of 10 mg/hr (OR 0.69, 95% CI 0.32 to 1.5).

7

1 Discussion

2 In this observational study, we found that patients who received intraoperative
3 ketamine infusions tended to be younger, sicker, and undergoing spine procedures of
4 longer duration (i.e., more complex) than those who did not. Patients who received
5 ketamine infusions were more likely to be taking preoperative opioids, and this increased
6 if patients were taking opioids on a **scheduled** basis.

7 We found that patients who were most likely to receive ketamine were those
8 undergoing the most complex spine procedures, often involving both an anterior and
9 posterior component **or a revision procedure** (Table 4). Previous studies examining
10 ketamine in spine **surgery have yielded conflicting results regarding postoperative opioid**
11 **consumption.**^{8,11-13} **The lack of consistent findings may have resulted, in part, from**
12 **combining potentially different procedures, listed as “lumbar fusion,”⁸ “lumbar or**
13 **thoracolumbar laminectomy and fusion,”¹² or “elective spine surgery.”¹³ Our data,**
14 however, suggest that anesthesiologists viewed patients undergoing longer, complex
15 procedures differently than less complex cases. For example, one of the most common
16 procedures, primary posterior lumbar fusion, had a low prevalence of ketamine
17 administration, suggesting that providers believed conventional opioid analgesia was
18 adequate. Patients who may benefit most from ketamine should be targeted for future
19 studies examining important long-term outcomes of interest. **Those patients undergoing**
20 **presumably less painful procedures with lower incidence of persistent postsurgical pain,**
21 **such as primary posterior lumbar fusion or anterior cervical fusion, may be able to be**
22 **managed with conventional therapy. Consistent with previous studies,⁹ the more complex**

1 and painful spine procedures were the ones most likely to be associated with ketamine
2 administration as a continuous infusion.

3 The apparent increased prevalence of ketamine use in patients taking both anti-
4 depressants and scheduled opioids was not surprising as the link between chronic pain,
5 anti-depressants, opioids, and spine surgery has been described.¹⁴ Although it is
6 impossible to determine retrospectively if this combined therapy factored into the
7 decision-making process by the anesthesia team, anti-depressant use is a potential
8 confounder that should be controlled for in future studies comparing ketamine to other
9 therapies.

10 Our results also confirm the tolerability of ketamine's ADEs in a clinical setting
11 (Table 5). This agrees with clinical trials in which up to 0.25 mg/kg/hr have been tried
12 without major ADEs.¹³ Our observed prevalence of central nervous system ADEs
13 (16.2%) is similar to the 22% retrospectively described by Rasmussen.¹⁵ Thus,
14 consideration of a variable-rate postoperative ketamine infusion in a treatment arm of a
15 randomized clinical trial is reasonable. Our ADE data confirm those from Mayo Clinic,
16 Jacksonville¹⁶ but go a step further in describing the specific ADEs, their prevalence, and
17 the rate of discontinuation in a daily clinical practice in which the infusion rates are
18 adjusted frequently.

19 Discontinuation of ketamine due to ADEs was unrelated to the maximum dose, a
20 somewhat surprising finding. Several factors may have played a role in this, including
21 ADEs resulting from simultaneous administration of opioids and benzodiazepines, as
22 well as variable patient sensitivity to ketamine, some of which may be related to
23 individual changes at the cellular level.¹⁷ There may be other unidentified factors as well.

1 Because 35 out of the 37 patients had resolution of symptoms once the infusions were
2 stopped, this suggests that the side effects were at least partially due to ketamine.

3 Using the data from this observational study, we are currently designing a
4 prospective, randomized trial comparing intra- and postoperative ketamine infusions to
5 placebo in opioid-tolerant patients undergoing complex spine surgery with particular
6 focus on long-term outcomes. Only complex spine procedures that are more likely to
7 result in severe pain will be included (i.e. anterior/posterior procedures or procedures
8 involving two or more spine areas, such as thoracic and lumbar). We will use the
9 frequency of ADEs encountered to guide the process of obtaining informed consent. The
10 ADE data could also be used as a guide for any hospitals considering starting a ketamine
11 service on the general medical floors.

12 Our study may have limited generalizability in that the decision to start an
13 intraoperative ketamine infusion was made at the discretion of the attending
14 anesthesiologist for the case. Thus, our findings may not apply to all practices. **Second,**
15 some patients may have been inappropriately placed in the “scheduled” opioid group
16 rather than the “as needed” group or vice versa due to documentation issues or patients
17 misrepresenting their opioid use.

18 In conclusion, we confirmed that postoperative ketamine infusions may be given
19 safely on general medical floors without special monitoring or intensive care, and
20 intraoperative infusions tend to be started for patients taking opioids, especially
21 scheduled opioids. Our data provide guidance both for hospitals considering the use of
22 ketamine infusions and for the design of future prospective, randomized clinical trials
23 looking at long-term benefits of ketamine or its ADEs.

References

1. Sullivan MD, Edlund MJ, Fan M, DeVries A, Braden JB, Martin BC. Trends in use of opioids for non-cancer pain conditions from 2000-2005 in Commercial and Medicaid insurance plans: The TROUP study. *Pain* 2008;138:440-449
2. Fredheim OMS, Mahic M, Shurtveit S, Dale O, Romundstad P, Borchgrevink PC. Chronic pain and use of opioids: A population-based pharmacoepidemiological study from the Norwegian Prescription Database and the Nord-Trondelag Health Study. *Pain* 2014;155:1213-1221
3. Benyamin R, Trescot AM, Datta S, et al. Opioid Complications and Side Effects. *Pain Physician* 2008;11:S105-S120
4. White JM, Irvine RJ. Mechanisms of fatal opioid overdose. *Addiction* 1999;94:961-972
5. Dunn KM, Saunders KW, Rutter CM, et al. Opioid Prescriptions for Chronic Pain and Overdose. *Ann Int Med* 2010;152:85-92
6. Trujillo KA, Akil H. Inhibition of Morphine Tolerance and Dependence by the NMDA Antagonist MK-801. *Science* 1991;251:85-87
7. Elliott K, Minami N, Kolesnikov YA, Pasternak GW, Inturrisi CE. The NMDA receptor antagonists, LY274614 and MK-801, and the nitric oxide synthase inhibitor, NG-nitro-L-arginine, attenuate analgesic tolerance to the mu-opioid morphine but not to kappa opioids. *Pain* 1994;56:69-75

8. 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;113:639-646
9. Laskowski K, Stirling A, McKay WP, Lim HJ. A systematic review of intravenous ketamine for postoperative analgesia. *Can J Anesth* 2011;58:911-923
10. McNicol ED, Schumann R, Haroutounian S. A systematic review and meta-analysis of ketamine for the prevention of persistent post-surgical pain. *Acta Anaesthesiol Scand* 2014;58:1199-1213
11. 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. *HSSJ* 2008;4:62-65
12. Subramaniam K, Akhouri V, Glazer PA, et al. Intra- and Postoperative Very Low Dose Intravenous Ketamine Infusion Does Not Increase Pain Relief after Major Spine Surgery in Patients with Preoperative Narcotic Analgesic Intake. *Pain Med* 2011;12:1276-1283
13. Garg N, Panda NB, Gandhi KA, et al. Comparison of Small Dose Ketamine and Dexmedetomidine Infusion for Postoperative Analgesia in Spine Surgery – A Prospective Randomized Double-blind Placebo Controlled Study. *J Neurosurg Anesthesiol* 2015;May 14 epub ahead of print
14. Walid MS, Zaytseva NV. Prevalence of mood-altering and opioid medication use among spine surgery candidates and relationship with hospital cost. *J Clin Neurosci* 2010;17:597-600

15. Rasmussen KG. Psychiatric side effects of ketamine in hospitalized medical patients administered subanesthetic doses for pain control. *Acta Neuropsychiatr* 2014;26:230-233
16. Porter SB, McClain RL, Howe BL, et al. Perioperative Ketamine for Acute Postoperative Analgesia: The Mayo Clinic-Florida Experience. *J Perianesth Nurs* 2015;30:189-195
17. Villasenor A, Ramamoorthy A, Silva dos Santos M, et al. A pilot study of plasma metabolomic patterns from patients treated with ketamine for bipolar depression: evidence for a response-related difference in mitochondrial networks. *Br J Pharmacol* 2014;171:2230-2242.