Intravenous Acetaminophen in Spine Surgery Patients: A Large Retrospective Analysis

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Abstract

Purpose: Opioids are the standard of care and often the most effective in treatment of pain. New intravenous formulations of Acetaminophen (IVA) may help decrease the number of opioids needed in the post-operative period therefore decreasing potential opioid side effects.

Methods: In a case control model, patients receiving intravenous acetaminophen and opioids were compared to patients receiving opioids alone. The patient groups were further broken down based on patients’ preoperative opioid usage. Morphine equivalent (ME) doses were then compared as well as length of stay (LOS), intensive care length of stay (ICU LOS), adverse drug reactions to opioids and IV acetaminophen, and surgical complications.

Results: Morphine equivalent doses were not different between IVA and opioid alone cohorts (83.56±7.64 ME vs. 93.01±9.02 ME [p=0.45]). LOS was also not significantly different (86.36±7.64 ME vs. 122.97±15.96 ME [p=0.10]). ICU LOS was also found to be nonsignificant.

Conclusions: Addition of intravenous acetaminophen did not decrease the usage of opioids and was not found to significantly decrease ICU stay. A more formalized randomized trial may help to find cohorts which would benefit from intravenous acetaminophen usage.

Introduction

Post-operative pain management in spinal surgery is often a challenge. This is further complicated by the large proportion of patients with chronic pain and the significant number of patients using opioid analgesics preoperatively [1]. Post-operative inadequate pain control is associated with increased lengths of stay and increased morbidity [1] including the development of chronic pain [2]. The cornerstone of post-operative analgesia has long been opioids, however, despite their potent analgesic properties, opioids are plagued by a host of side effects. Opioids cause gastrointestinal and respiratory depression and, of added importance in the neurosurgical patient, sedation [3]. These adverse drug effects may result in increased lengths of stay and more costly hospital stays.

A possible solution to the problems of post-operative analgesia is multimodal pain management and the use of supplemental non-narcotic analgesics such as NSAIDs [4]. In 2010, intravenous acetaminophen was approved in the United States and has been studied in a variety of surgical populations with mixed results [5-7].

Although the mechanism of action of acetaminophen is incompletely understood, it is thought to act through inhibition of COX-1 and COX-2 dependent pathways. Furthermore, there is now evidence that other mechanisms involving COX-3 inhibition and activation of capsaicin and cannabinoid receptors may also play a role in its analgesic properties [8,9]. Acetaminophen is not associated with the side effects of opioids and NSAIDs and compared to oral or rectal formulations; intravenous forms boast superior pharmacokinetics, with faster onset of action and more predictable plasma and CSF concentrations [5,10,11]. However, the intravenous formulation is costly, and as such its effectiveness with regards to both its analgesic qualities and its role in reducing opioid consumption and opioid related side
effects must be closely examined [7,12-17]. Although initial studies repeatedly demonstrated that IV acetaminophen results in lower visual analog pain scores (VAS) than placebo, its effect on opioid consumption has been less clear [14,17]. While some studies have shown IV acetaminophen use has reduced post-operative opioid use, many studies in a variety of non-spine surgical populations have demonstrated no difference in opioid consumption between patients receiving IV acetaminophen and those not [13,15,16,18-21]. The literature in spinal patients has shown mixed results with regards to the effect of IV acetaminophen on opioid consumption and opioid related side effects [7,12,22-24]. In the current study, we seek to help build a consensus on the role of IV acetaminophen in the management of post-operative pain following spinal surgery.

Materials and Methods

Study Design

This was a single center retrospective case-control study. The electronic medical records of patients included in the study were examined after proper approval from the institutional IRB and the relevant data was obtained and analyzed.

Subjects

All patients undergoing first time spinal surgery by a single surgeon at Temple University Hospital between 12/2011 and 8/2015 were included in the preliminary subject list. Patients were excluded if they had previous spinal surgery within the prior thirty days, if they were discharged on the same day as their surgery, or were on extremely high levels (Morphine Equivalent Doses greater than four standard deviations above the mean) of chronic opioid narcotics prior to surgery for cancer palliation. Furthermore, patients undergoing minimally invasive surgery were excluded because these patients only rarely received IV acetaminophen and therefore the treatment group was underpowered for any analysis. Patients were retrospectively divided into two groups for analysis: those who received intravenous IV acetaminophen post-operatively and those who did not. IV acetaminophen was added to our institution’s formula in late 2013. Prior to that date, no patients received IV acetaminophen and following its addition to our formulary, the majority of patients received IV acetaminophen following spinal surgery. Patients in the experimental group received 1g IV acetaminophen every six hours for either twenty four or forty eight hours. Patients discharged prior to forty eight hours post-op received only twenty four hours of treatment with IV acetaminophen, whereas patients that stayed longer than forty eight hours received forty eight hours of IV acetaminophen. For subset analysis, patients were further divided into those receiving surgery on the cervical, thoracic or lumbar spine. For procedures which spanned more than one vertebral level, the patient was placed in the cohort corresponding to the vertebral region in which the majority of their case was performed. Baseline demographics were collected including age, sex, and pre-operative opioid use.

Endpoints

The primary study endpoint was opioid use over the first twenty four hours post-operatively. To assess and standardize opioid use, doses of all opioids administered were converted into their morphine equivalents (ME, mg/day). For all oral opioids, doses were converted to an equivalent intravenous dose based on the drugs bioavailability. The morphine equivalents of all opioids administered within twenty-four hours of the conclusion of the surgery were totaled and recorded. The study examined several secondary endpoints. These were opioid use during twenty four to forty-eight hours post-op, length of stay (LOS), intensive care length of stay (ICU LOS), adverse drug reactions to opioids and IV acetaminophen, and surgical complications.

Statistical Analysis

For all continuous variables, significance was determined by a two-tailed Student's t-test. For categorical values, a Chi-square test was performed. For binomial variables, a z-test of proportions was performed.

Results

Baseline Patient Demographics

A total of three hundred and eleven patients were included in the final analysis. One hundred and twenty-three patients were in the experimental group and one hundred and eighty-nine patients were in the control group (Table 1). In the experimental cohort, the average age was fifty-seven years old (57.76±1.21) and forty six percent were female (46.34%). Forty four percent (44.71%) of patients were taking opioids preoperatively. In the control cohort, the average age was fifty-six (56.06±1.13) and thirty seven percent (37.04%) were female. Twenty six percent (26.46%) of the patients were taking opioids preoperatively. The baseline difference in the proportion of patients taking opioids preoperatively was deemed to be significant (44.71% vs. 26.46% [p<0.001]) (Table 1). A total of one hundred and sixty-five patients underwent surgery to the cervical spine. Sixty-three patients were in the treatment group with an average age of fifty-six years old (56.86±1.68). Forty four percent (44.11%) were female. One hundred and two patients were in the control group with an average age of fifty-seven years old (57.92±1.51).

Thirty percent (30.22%) were female. In the treatment group, thirty eight percent of patients were taking opioids preoperatively compared to nineteen percent in the control group (38.10% vs. 18.62% [p=0.006]) (Table 1). Cervical procedures included anterior discectomy and fusions (ACDF), posterior laminectomy and fusions, posterior fusions alone, and posterior laminectomy and fusions. The case breakdown among the acetaminophen and non-acetaminophen users was not significant (Table 2). A total of fifty-six patients underwent surgery to the thoracic spine. Nineteen patients were in the treatment group with an average age of sixty
years old (60.00±2.39). Forty seven percent were female (47.37%). Thirty-seven patients were in the control group with an average age of fifty-five (55.57±2.51). Fifty one percent were female (51.35%). In the treatment group, forty two percent of patients were taking opioids preoperatively compared to thirty percent in the control group (42.11% vs. 29.73% [p=0.35]) (Table 1). Thoracic procedures included posterior laminectomy and fusions, posterior laminectomies alone, posterior fusions alone, and posterior corpectomies. There was not a large difference in case breakdown between groups (Table 2).

Table 1.

| Region | Chronic Opioid Users | Opioid Naive |
|--------|----------------------|--------------|
|        | IVA + Opioids        | Opioids Alone|
| Cervical |                     |              |
|         | # of patients (n)    | Age          | P-value | Gender(%) Female | P-value |
|        | 24                   | 54.88±2.62   | 0.62    | 54.17            | 0.14    |
|        | 19                   | 57.00±3.37   |         | 31.58            |         |
|        | 39                   | 58.08±2.20   | 0.98    | 38.46            | 0.23    |
|        | 83                   | 58.01±1.70   |         | 27.71            |         |
| Thoracic |                     |              |
|         | IVA + Opioids        | Opioids Alone|
|         | 8                    | 60.00±2.39   | 0.07    | 50.00            | 0.14    |
|         | 11                   | 55.57±2.51   |         | 81.81            |         |
|         | 11                   | 56.36±4.21   | 0.65    | 45.45            | 0.69    |
|         | 26                   | 58.83±3.08   |         | 38.46            |         |
| Lumbar |                     |              |
|         | IVA + Opioids        | Opioids Alone|
|         | 22                   | 56.13±3.18   | 0.87    | 50.00            | 0.52    |
|         | 20                   | 55.40±2.97   |         | 60.00            |         |
|         | 18                   | 56.00±3.91   | 0.18    | 50.00            | 0.25    |
|         | 30                   | 49.59±2.79   |         | 33.33            |         |
| Total  |                     |              |
|         | IVA + Opioids        | Opioids Alone|
|         | 55                   | 56.84±1.85   | 0.91    | 49.09            | 0.48    |
|         | 50                   | 56.56±1.90   |         | 56.00            |         |

Table 2.

| Region | ACDF | Post. Lami/Fusion | Post. Fusion | Post. Lami | Corpectomy w/Fusion |
|--------|------|-------------------|--------------|------------|---------------------|
| Cervical | IVA + Opioids | 8 (3.3)           | 12 (50)      | 3(12.5)    | 1(4.2)              | 0(0) |
|         | Opioids Alone | 6 (3.1)           | 9 (47.4)     | 2(10.5)    | 1(5.3)              | 1(5.3)|
|         | IVA + Opioids | 10 (25.6)         | 21 (53.8)    | 4 (10.3)   | 4(10.3)             | 0(0) |
|         | Opioids Alone | 23 (25.8)         | 41 (49.4)    | 13(14.6)   | 4(4.5)              | 2(2.2)|
| Thoracic | IVA + Opioids | 1 (1.25)          | 2 (25)       | 3(47.5)    | 1 (12.5)            | 0(0) |
|         | Opioids Alone | 3 (27.3)          | 2 (18.2)     | 4 (36.4)   | 1 (9.1)             | 0(0) |
|         | IVA + Opioids | 3 (27.3)          | 2 (18.2)     | 4 (36.4)   | 1 (9.1)             | 1(9.1)|
|         | Opioids Alone | 10 (38.5)         | 3 (11.5)     | 9 (34.6)   | 2 (7.7)             | 0(0) |
| Lumbar | IVA + Opioids | 4 (18.2)          | 2 (9.1)      | 1 (4.5)    | 4 (18.2)            | 3 (13.6)|
|         | Opioids Alone | 4 (20.0)          | 3 (15.0)     | 0 (0)      | 3 (15.0)            | 2 (10.0)|
|         | IVA + Opioids | 4 (22.2)          | 3 (16.7)     | 0 (0)      | 2 (11.1)            | 3 (16.7)|
|         | Opioids Alone | 6 (20.0)          | 4 (13.3)     | 1 (3.3)    | 6 (20.0)            | 5 (16.7)|

A total of ninety patients underwent surgery to the lumbar spine. Forty patients were in the treatment group with an average age of fifty-five (55.28±2.38). Fifty percent (50.00%) were female. Fifty patients were in the control group with an average age of fifty-two (52.84±2.32). Forty four percent were female (44.00%). Fifty five percent of patients in the treatment group were taking opioids preoperatively compared to forty percent of patients in the control group (55% vs. 40% [p=0.16]) (Table 1). Lumbar procedures
included posterior laminectomies, posterior laminectomy, posterior lumbar interbody with posterior fusion, transforaminal lumbar interbody fusion, direct lateral interbody fusion, and kyphoplasties. The case breakdown was not significant amongst groups (Table 2).

**Primary Endpoint**

The primary study endpoint was opioid use in the first twenty-four hours postoperatively. No significant difference was observed between the control and treatment cohorts when pooling all subsets (83.56±7.64 ME vs. 93.01±9.02 ME [p=0.45]). However, to control for the significantly higher proportion of patients on preoperative opioids in the treatment group, patients were further subdivided into those who were opioid naïve and those on opioid narcotics preoperatively. In both subsets, the difference was not significant (Chronic opioid users: 110.07±12.06 vs. 103.25±13.16 [p=0.71], Opioid naïve: 62.11±7.08 vs. 89.32±11.22 [p=0.11]). This finding was echoed in all subsets. Among cervical, thoracic and lumber patients, no significant difference was observed in the primary endpoint (58.00±6.31 vs. 65.15±9.47 [p=0.58], 81.16±12.36 vs. 93.93±19.29 [p=0.65], 124.03±16.30 vs. 149.16±22.44 [p=0.39] respectively). However, to again control for the significantly higher proportion of cervical spine patients taking preoperative opioids in the treatment group, patients were again subdivided into those who were opioid naïve and those on opioid narcotics preoperatively. Similarly, in both subsets, the difference was not significant (Chronic opioid users: 104.63±20.84 vs. 57.09±18.14 [p=0.10], Opioid naïve: 64.09±13.58 vs. 93.10±23.57, [p=0.43]). Although the proportion of patients taking preoperative opioids in both the lumbar and thoracic subsets were not significantly different, identical analysis based on pre-operative opioid use was performed with similar results (Figure 1).

**Secondary Endpoints**

Several secondary endpoints were examined. The first was post-operative opioid use for the period of twenty-four to forty-eight hours post-op. No significant difference was observed between patients receiving IV acetaminophen and those receiving opioids alone (86.36±7.64 ME vs. 122.97±15.96 ME [p=0.10]). When controlling for the higher proportion of patients in the experimental cohort on pre-operative narcotics, there was again no significant difference (Chronic opioid users: 93.61±10.86 vs. 91.79±12.57 [p=0.91], Opioid naïve: 78.96±10.74 vs. 133.19±20.74 [p=0.12]). This finding was echoed for cervical and thoracic patients. Among lumbar patients, post-operative opioid use in chronic opioid users was not significant (137.10±25.23 vs. 132.2±23.11 [p=0.89]). Among Opioid naïve patients, those patients receiving IV acetaminophen trended toward lower post-operative opioid use over this time period, however this trend failed to reach significance (89.77±21.52 vs. 277.67±71.49 [p=0.06]) (Figure 2). Length of stay and intensive care length of stay were also examined. For both chronic opioid users and patients who were opioid naïve, no significant difference between length of stay or intensive care length of stay was observed (LOS: Chronic opioid users: 4.07±0.40 days vs. 3.86±0.45 days [p=0.72], opioid naïve: 6.01±0.66 vs. 7.40±0.77 [p=0.25], ICU LOS: Chronic opioid users: 2.50±0.34 vs. 1.76±1.11 [p=0.21], opioid naïve: 3.84±1.15 vs. 4.94±1.36 [p=0.57]). The same was found when looking at cervical, thoracic and lumbar patient subgroups (Figure 3). Adverse drug reactions and surgical and post-operative complications were also collected and examined. Among those reactions and complications observed were urinary retention, wound breakdown, fevers, deep venous thrombosis, gastrointestinal bleeding, ileus, pneumonia and wound infections (Table 3).
**Figure 2:** Morphine equivalent doses for patients 24-48 hours post op. Cohort break down and spinal segment of surgery is shown.

**Figure 3:** Length of stay shown by cohort. ICU length of stay also included.

**Table 3:** Adverse Drug Reactions and Complications Observed.

| Reaction/Complication                | IV Acetaminophen + Opioids | Opioids Alone |
|--------------------------------------|----------------------------|---------------|
| Urinary Retention                    | 2                          | 2             |
| Return to OR                         | 8                          | 6             |
| Wound Breakdown                      | 2                          | 0             |
| Fevers                               | 3                          | 1             |
| PE                                   | 2                          | 1             |
| DVT                                  | 1                          | 0             |
| GI Bleed                              | 1                          | 0             |
| Ileus                                 | 0                          | 2             |
| Seizure                               | 1                          | 1             |
| Pneumonia                             | 0                          | 1             |
| Operative Site Infection             | 0                          | 2             |
| Urinary Tract Infection              | 0                          | 2             |
| 30 Day Mortality                     | 2                          | 0             |
Discussion

The role of IV acetaminophen in the management of postoperative pain in patients undergoing spinal surgery is still be fully evaluated. In our study, we observed that IV acetaminophen did not significantly affect post-operative opioid use in either the first twenty-four hours or the subsequent twenty-four. Furthermore, no difference was observed in LOS or ICU LOS. This trend was repeatedly observed in each subset of patients examined, even when controlling for the long-term use of opioid narcotics prior to surgery. Similarly, no significant difference was observed between the treatment cohort and control patients with regard to adverse drug effects and surgical complications. Our study is the largest and most comprehensive to date to examine IV acetaminophen in spinal patients. Although each subset of patients was of moderate size, the same findings were repeatedly demonstrated across each population examined. This provides evidence that IV acetaminophen may not be medically or economically worthwhile as a supplement to traditional opioid narcotics for post-operative pain. However, it should be noted that our study did not examine subjective pain such as VAS scores.

In studies where VAS scores have been assessed, it has been observed that the addition of IV acetaminophen to opioid narcotics results in lower subjective pain, even when the opioid consumption has not been different [7,12,14,16,20,22]. In a time when patient satisfaction is increasingly important, the effects of IV acetaminophen on subjective pain should not be overlooked. Furthermore, there is evidence that inadequate pain control during the postoperative period may play a role in the development of chronic pain syndromes [2]. This suggests that even in the absence of an objective measure of post-operative pain such as opioid consumption, it may be beneficial to supplement analgesia with IV acetaminophen. Our study was limited by its retrospective nature, and by the fact that post-operative opioids were not standardized. By this, it is meant that patients received different narcotic medications as well as different formulations. We attempted to control for this by converting all intravenous formulations to their oral equivalents based on their bioavailability. We then converted all narcotics to morphine equivalents based on their known potencies. And while this is, admittedly, not a perfect system, it allows for an adequate comparison. In conclusion, our study demonstrates that IV acetaminophen is ineffective in reducing postoperative opioid consumption in spinal patients. Furthermore, it does not reduce length of stay or the incidence of adverse drug effects associated with opioids. However, there may still be a role for IV acetaminophen according to evidence that it improves subjective pain.

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