Complications following regional anesthesia versus general anesthesia for the treatment of distal radius fractures

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Received: 13 January 2021 / Accepted: 13 May 2021 / Published online: 29 May 2021
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Abstract

Purpose Open reduction and internal fixation (ORIF) are commonly utilized for the repair of distal radius fractures (DRF). While general anesthesia (GA) is typically administered for ORIF, recent studies have also demonstrated promising results with the usage of regional anesthesia (RA) in the surgical treatment of distal radius fractures. This study will compare complication rates between the use of RA versus GA for ORIF of DRFs.

Methods A multi-institutional surgical registry was utilized to identify patients who had undergone ORIF for DRFs from 2005 to 2018—these patients were stratified into GA and RA cohorts. Patients were matched utilizing coarsened-exact-matching (CEM) to compare postoperative outcomes and rates of 30-day complications were compared between the two cohorts.

Results Upon CEM-matching, 1191 patients receiving RA were matched to 9250 patients who had received GA, with a multivariate imbalance measure (L1) statistic of < 0.001. In the matched-cohort analysis, no significant differences were observed in rates of any complication (all $p \geq 0.083$). On multivariate regression analyses, RA was not associated with increased risk for any complication ($p = 0.445$), minor complications ($p = 0.093$), major complications ($p = 0.758$), unplanned reoperations ($p = 0.355$), unplanned readmissions ($p = 0.799$), or mortality ($p = 0.579$).

Conclusion With similar safety profiles, RA is a safe and reasonable alternative to GA when managing DRFs surgically. RA may be the preferred anesthetic technique for ORIF of DRFs in patients at high risk with GA, such as those with reactions to GA in the past or with significant cardiopulmonary risk factors.

Keywords Distal radius fracture · Regional anesthesia · Complications · General anesthesia · Peripheral nerve block

Introduction

Distal radius fractures (DRF) are the most common long bone fracture and are typically caused by a fall on an outstretched hand [1]. DRFs account for up to 18% of all fractures in the elderly population and up to 20% of all fractures treated in an emergency department setting [1–3]. Open reduction and internal fixation (ORIF) have recently become a mainstay of treatment in these cases as outcomes are very favorable with respect to healing and articular congruency [2, 4]. However, ORIF of DRFs are not without complications. With DRFs being one of the most common orthopaedic injuries, it is important to try and mitigate any other complications that could be prevented.

One way to try and minimize complications is by utilizing regional anesthesia (RA). RA has been shown to have beneficial effects including decreased perioperative opioid consumption, decreased length of stay (LOS), and decreased

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nausea in various orthopaedic procedures [5–8]. In addition, the use of RA in ORIF of DRFs has been reported to contribute to decreased pain and increased range of motion earlier in the post-operative period potentially expediting rehabilitation. [7] Although the use of RA and outcomes following DRF have been previously explored, only single-rehabilitation. [7] Although the use of RA and outcomes following ORIF of DRFs have been reported to contribute to decreased pain and increased range of motion, the use of RA in ORIF of DRFs has been reported to contribute to decreased pain and increased range of motion, the use of RA in ORIF of DRFs has been reported to contribute to decreased pain and increased range of motion, the use of RA in ORIF of DRFs has been reported to contribute to decreased pain and increased range of motion, the use of RA in ORIF of DRFs has been reported to contribute to decreased pain and increased range of motion, the use of RA in ORIF of DRFs has been reported to contribute to decreased pain and increased range of motion.

To the author’s knowledge, no study has been conducted with a large patient cohort analyzing outcomes following RA versus general anesthesia (GA) for ORIF of DRFs. This present study seeks to utilize the American College of Surgeons National Quality Improvement Program (ACS-NSQIP) database to compare outcomes following ORIF of DRFs with a large nationwide patient population to verify previous reports’ support for the role of RA. In doing so, surgeons can help mitigate complications for a very common injury and can better counsel patients on which type of anesthesia to utilize for the procedure.

Materials and methods

Patient selection

The ACS-NSQIP database was queried to identify all patients who had undergone surgery for the treatment of distal radius fracture from 2005 to 2018. Patients were selected based on their Current Procedure Terminology (CPT) codes, corresponding with CPT 25607, 25608, and 25609. Patients were then stratified into two cohorts based on the type of anesthesia they received: GA or RA. Patients receiving other methods of anesthesia, including epidural, spinal, and monitored anesthesia care/intravenous (MAC/IV) sedation, were excluded from this study. The ACS-NSQIP database classifies specific cases based on the primary method of anesthesia utilized. As such, if RA or epidural anesthesia was utilized with MAC or IV sedation, the primary method of anesthesia recorded in the ACS-NSQIP database would be classified as MAC/IV sedation. In addition, if RA was utilized at first but GA had to be utilized, then the primary method of anesthesia recorded would be GA. As such, the present study only compares ORIF cases for DRF in which RA was the sole method of anesthesia utilized versus GA. A total of 14,503 patients met inclusion criteria, of which 1348 (9.29%) had received regional anesthesia and 13,135 (90.71%) had received GA.

Variables

The ACS-NSQIP database contains over 150 variables for each patient, providing a comprehensive medical history and detailed reporting on postoperative complications. Demographic information was collected to better characterize this patient cohort; demographic variables included age, gender, race, and BMI, which was calculated with the given height (cm) and weight (kg) values. Preoperative comorbidities were also analyzed for differences between obesity classes, including smoking history, diabetes mellitus, dyspnea, ventilator dependence, chronic obstructive pulmonary disease (COPD), ascites, congestive heart failure (CHF), hypertension requiring medication management, acute renal failure, dialysis, disseminated cancer, open wounds/wound infections, chronic steroid use, significant weight loss of more than 10% of total body weight within the six-month preoperative period, hematologic disorders (i.e. hemophilias, coagulopathies, vitamin deficiencies), preoperative blood transfusions (within 72 h of the start of surgery), systemic sepsis, and functional dependence. Perioperative variables, such as ASA (American Society of Anesthesiologists) classification and type of anesthetic administered, were also included for analysis.

Medical complications occurring within the 30-postoperative period were reported by the ACS-NSQIP database, in addition to administrative parameters such as length of hospital stay and discharge information. This retrospective study included complications that were present in all available years of the ACS-NSQIP database to minimize the confounding effects of missing and incomplete data. The postoperative complications included in this study were superficial surgical site infections (SSI), deep SSI, organ/ space SSI, wound dehiscence, pneumonia, unplanned intubation, pulmonary embolisms, ventilator dependence (> 48 h), progressive renal insufficiency, acute renal failure, urinary tract infections (UTI), cerebrovascular accidents (CVA)/strokes, cardiac arrest requiring cardiopulmonary resuscitation, myocardial infarction, transfusions, deep venous thromboembolisms (DVT), systemic sepsis, septic shock, unplanned reoperations, unplanned readmissions, and mortality.

Statistical analysis

Patient demographic factors, preoperative comorbidities, and postoperative complication rates were analyzed for differences based on the primary anesthetic that they had received. Pearson’s chi-squared tests and Fischer’s exact tests (when expected cell sizes were less than 5 and variance could not be assumed to be normally distributed) were used to assess for differences in categorical variables, which are reported as number of subjects (n) and incidence/prevalence rates (%). All continuous variables, such as age and time parameters, are expressed as mean values with their respective standard deviations. All continuous variables were analyzed for differences in mean with one-way analyses of variance (ANOVA).

To control for differences in patient characteristics and comorbidities, the present study implements a widely
accepted method called coarsened exact matching (CEM). CEM has been used previously in the literature with large patient cohorts and has been recognized as a superior method of matching to the ones most commonly used in large-scale studies, such as propensity score matching [13–17]. CEM temporarily coarsens the data to produce exact matches based on the entered variables, allowing for less statistical assumptions and model dependence when attempting to estimate the effect of a specific treatment (regional vs. general anesthesia in the present study). Patients receiving RA were matched to those receiving GA, dropping cases that were not matched exactly on the factors that were entered for controlling. Patients were matched based on age, gender, race, BMI, smoking status, diabetes mellitus, dyspnea, ventilator dependence, COPD, ascites, CHF, hypertension, acute renal failure, dialysis, disseminated cancer, open wounds/wound infections, chronic steroid use, significant weight loss, hematologic disorders, preoperative transfusions, sepsis, functional dependence, and ASA classification. This exact matching method ensures the inclusion of patients who are healthily comparable with the most similar baseline characteristics. These matched cohorts were all analyzed for differences in patient characteristics, as well as complication rates.

A multivariate logistic regression model was implemented with the weights provided by the CEM matching to analyze the independent effect of regional anesthesia on risks for complications in comparison to those receiving GA. Outcomes analyzed in the multivariate logistic regression models included any complication, minor complications, major complications, unplanned reoperation, unplanned readmission, and mortality. All statistical findings with \( p \) values less than or equal to 0.05 were considered significant for this study. Variance was not assumed to be equal for any variable with the exception of age. All statistical analyses were performed using the IBM SPSS Version 25 Software (IBM Corp., Armonk, NY) and R© Version 3.3.3.

### Results

A total of 14,503 patients met inclusion criteria, of which 1348 (9.29%) had received RA and 13,135 (90.71%) had received GA. In the unmatched analysis, the RA cohort was significantly older (\( \bar{x} = 59.82 \) years, SD 15.805) than the GA cohort (\( \bar{x} = 56.32 \) years, SD 16.402; \( p < 0.001 \)). The RA cohort was also comprised of a larger proportion of females (78.12 vs. 72.93%; \( p < 0.001 \)) than the GA cohort (\( \bar{x} = 77.24 \) min, SD 45.505; \( p = 0.001 \) (Table 2). No differences were observed in perioperative parameters, with the exception of total operative time. On average, the RA cohort experienced longer operative times (\( \bar{x} = 81.68 \) min, SD 39.248) than the GA cohort (\( \bar{x} = 56.32 \) years, SD 16.402; \( p < 0.001 \)). (Supplementary Table 1). No differences were observed in perioperative parameters, with the exception of total operative time. On average, the RA cohort experienced longer operative times (\( \bar{x} = 81.68 \) min, SD 39.248) than the GA cohort (\( \bar{x} = 56.32 \) years, SD 16.402; \( p < 0.001 \)). (Supplementary Table 1).

In the matched-cohort analysis, no significant differences were observed in rates of any complication (all \( p \geq 0.083 \); Table 3). On multivariate regression analyses, RA was not associated with increased risk for any complication (\( p = 0.445 \)), minor complications (\( p = 0.093 \)), major complications (\( p = 0.758 \)), unplanned reoperations (\( p = 0.355 \)), unplanned readmissions (\( p = 0.799 \)), or mortality (\( p = 0.579 \)). (Table 4)

### Discussion

The use of anesthetic techniques has revolutionized medicine since its inception in the nineteenth century. GA provides intraoperative amnesia and has been used successfully in a plethora of both orthopaedic and non-orthopaedic procedures as it can be continuously administered with quick reversibility in the event of unexpected complications [18]. However, GA is typically inadequate in its own at providing local pain control in the immediate post-operative period and can suppress normal autonomic function requiring increased surveillance/monitoring [18]. Less common side effects of GA and intubation include, but are not limited to, aspiration, nausea, pruritis, and hoarseness [18]. In comparison to GA, RA avoids these aforementioned side effects and has been associated with improved postoperative pain control, decreased opioid use, and reduced recovery times in certain orthopedic procedures [18–22]. RA with laryngeal mask airway has also been reported to be associated with decreased rates of intraoperative hemodynamic instability and postoperative cognitive function compared to GA with endotracheal intubation in elderly patients for intertrochanteric fracture management [23]. RA has demonstrated other advantages including preservation of autonomic function, improved muscle relaxation, avoidance of airway manipulation, decreased nausea/vomiting, and reduced post-anesthesia care unit stay [18, 24, 25].

This current study demonstrates no significant differences in rates of any complications between the matched
Table 1  Demographics and comorbidities in CEM-matched general vs. regional anesthesia

| Demographics                          | General ($n=9250$) | Regional ($n=1191$) | $P$ value |
|---------------------------------------|--------------------|---------------------|-----------|
| Age (Mean ± SD)$^a$                   | 55.68 ± 16.738     | 55.88 ± 16.811      | 0.691     |
| Age (years)                           |                    |                     | 1.000     |
| x ≤ 40                                | 1906               | 20.61%              | 20.57%    |
| 40 < x ≤ 50                           | 1006               | 10.88%              | 10.92%    |
| 50 < x ≤ 60                           | 2249               | 24.31%              | 24.35%    |
| 60 < x ≤ 70                           | 2408               | 26.03%              | 26.03%    |
| 70 < x                                | 1681               | 18.17%              | 18.14%    |
| Sex                                   |                    |                     | 0.976     |
| Female                                | 7079               | 76.53%              | 76.49%    |
| Male                                  | 2171               | 23.47%              | 23.51%    |
| Race/Ethnicity                        |                    |                     | 0.997     |
| American Indian or Alaska Native      | 0                  | 0.00%               | 0.00%     |
| Asian or Pacific Islander             | 222                | 2.40%               | 2.43%     |
| Black or African American             | 102                | 1.10%               | 1.09%     |
| Hispanic                              | 0                  | 0.00%               | 0.00%     |
| White or Caucasian                    | 8926               | 96.50%              | 96.47%    |
| Body mass index (kg/m²)               |                    |                     | 1.000     |
| Normal                                | 3432               | 37.10%              | 37.11%    |
| Overweight                            | 3300               | 35.68%              | 35.68%    |
| Class I obese                         | 1643               | 17.76%              | 17.80%    |
| Class II obese                        | 544                | 5.88%               | 5.88%     |
| Class III obese                       | 331                | 3.58%               | 3.61%     |
| Pre-operative comorbidities           |                    |                     |           |
| Diabetes mellitus                     |                    |                     | 0.996     |
| No diabetes mellitus                  | 9008               | 97.38%              | 97.40%    |
| Non-insulin dependent                 | 189                | 2.04%               | 2.02%     |
| Insulin dependent                     | 53                 | 0.57%               | 0.59%     |
| Smoking history                       | 1420               | 15.35%              | 15.37%    | 0.990    |
| Dyspnea                               |                    |                     | 0.934     |
| No dyspnea                            | 9227               | 99.75%              | 99.75%    |
| Moderate exertion                     | 22                 | 0.24%               | 0.25%     |
| At rest                               | 1                  | 0.01%               | 0.00%     |
| Ventilator dependence                 | 0                  | 0.00%               | 0.00%     | –        |
| COPD                                  | 71                 | 0.77%               | 0.76%     | 0.965    |
| Ascites                               | 0                  | 0.00%               | 0.00%     | –        |
| Congestive heart failure              | 1                  | 0.01%               | 0.00%     | 1.000    |
| Hypertension                          | 2572               | 27.81%              | 27.79%    | 0.992    |
| acute renal failure                   | 0                  | 0.00%               | 0.00%     | –        |
| Dialysis                              | 0                  | 0.00%               | 0.00%     | –        |
| Disseminated cancer                   | 0                  | 0.00%               | 0.00%     | –        |
| Open wounds/wound infections          | 15                 | 0.16%               | 0.17%     | 1.000    |
| Chronic steroid use                   | 32                 | 0.35%               | 0.34%     | 1.000    |
| Weight loss                           | 0                  | 0.00%               | 0.00%     | –        |
| Bleeding disorders                    | 39                 | 0.42%               | 0.42%     | 0.993    |
| Transfusions (within 72 h preop)      | 0                  | 0.00%               | 0.00%     | –        |
| Systemic sepsis                       | 1                  | 0.01%               | 0.00%     | 1.000    |
| Functional status                     |                    |                     | 0.868     |
| Independent                           | 9208               | 99.55%              | 99.58%    |
| Partially dependent                   | 42                 | 0.45%               | 0.42%     |
| Totally dependent                     | 0                  | 0.00%               | 0.00%     |
RA and GA cohorts. The lack of differences in complication rates implicates that RA is a reasonable and safe alternative to GA for ORIF of DRFs. RA has been demonstrated to be efficacious and safe in other orthopaedic procedures as well. Helwani et al. previously reported reductions in post-operative deep SSI rates, hospital LOS, and major complications in patients undergoing total hip arthroplasty when utilizing RA compared to GA [8]. Liu et al. similarly reports a reduced risk of pneumonia and systemic infectious complications following total knee arthroplasty with neuraxial anesthesia compared to GA [23]. While the present study did not demonstrate a reduction in these adverse events and did not analyze the effects of neuraxial anesthesia, no significant differences were exhibited in terms of any complications, unplanned reoperations, or mortality when comparing RA and GA for ORIF of DRFs. The lack of differences in adverse events demonstrated in this study further supports the safety profile of RA and its role as a reasonable alternative to GA in ORIF of DRFs. Certain patient demographics and comorbidities, such as obesity, greater age, and frailty, are known to increase the risk of adverse events with GA [26]. For example, GA has been associated with an increased risk of postoperative cognitive dysfunction in elderly patients when compared to other forms of anesthesia [27]. The current findings suggest that, without any differences in complication rates, RA for DRF is a reasonable/safe alternative, and may be the preferred mode of anesthesia for high-risk patients—future work exploring the safety profile of RA in ORIF of DRFs in these high-risk patients is warranted. However, as with all other aspects of a patient’s care, the options and alternatives of anesthesia available should be thoroughly discussed with the patient so that factors such as anxiety and comfort can be taken into account. In doing so, the patient can make an informed decision with the perioperative care team’s recommendations.

There is currently debate within the literature as to whether RA improves functional outcomes following ORIF of DRFs in comparison to GA. However, brachial plexus blockade in upper extremity surgery has generally demonstrated reduced systemic anesthetic requirements, lower perioperative opioid consumption, decreased nausea, and shortened postoperative stays [5, 7, 8, 28, 29]. In a randomized control trial, Hadzic et al. reported lower pain scores for patients undergoing hand/wrist surgery who had received RA compared to GA in the perioperative period [30]. Ego et al. reported significantly decreased pain and improved function at three and six-month follow-up in patients who had received RA compared to GA in the perioperative period [30]. Ego et al. compared the use of hematoma blocks versus GA and found

### Table 1 (continued)

| ASA classification | General (n=9250) | Regional (n=1191) | P value |
|--------------------|-----------------|------------------|---------|
| 1-No disturb       | 1700            | 219              | 0.996   |
| 2-Mild disturb     | 5796            | 746              | 0.626   |
| 3-Severe disturb   | 1744            | 225              | 0.189   |
| 4-Life threat      | 10              | 1                | 0.087   |
| 5- Moribund        | 0               | 0                | 0.000   |
| None               | 0               | 0                | 0.000   |

All other values expressed as (%) and N

COPD Chronic obstructive pulmonary disease

*Values expressed as Mean ± Standard Deviation (SD)*

### Table 2 Perioperative parameters in CEM-matched general vs. regional anesthesia cohorts

| Parameter                                | General (n=9250) | Regional (n=1191) | P value |
|------------------------------------------|-----------------|------------------|---------|
| Days to operation from admission        | 0.22 ± 6.258    | 0.06 ± 0.774     | 0.377   |
| Total operating time (Minutes)          | 77.24 ± 45.505  | 81.68 ± 39.248   | 0.001   |
| Total hospital stay length (Days)       | 0.52 ± 5.847    | 0.34 ± 2.213     | 0.304   |
| Days from operation to discharge        | 0.33 ± 2.056    | 0.28 ± 2.035     | 0.463   |
| Wound classification                     |                 |                  | 0.084   |
| 1-Clean                                  | 9018            | 1174             | 98.57%  |
| 2-Clean/contaminated                     | 137             | 11               | 0.92%   |
| 3-Contaminated                           | 69              | 6                | 0.50%   |
| 4-Dirty/infected                         | 26              | 0                | 0.000   |
Table 3 Postoperative complication rates in CEM-matched general vs. regional anesthesia cohorts

| Postoperative complications         | General (n = 9250) | Regional (n = 1191) | P value |
|-------------------------------------|--------------------|---------------------|---------|
| Any complication                    | 187                | 28                  | 0.451   |
| Minor complication                  | 44                 | 10                  | 0.127   |
| Superficial incisional SSI          | 11                 | 4                   | 0.083   |
| Wound disruption                    | 2                  | 1                   | 0.305   |
| Pneumonia                           | 6                  | 0                   | 1.000   |
| Progressive renal insufficiency     | 1                  | 0                   | 1.000   |
| Urinary tract infection             | 15                 | 4                   | 0.264   |
| Blood transfusions                  | 9                  | 1                   | 1.000   |
| Major complication                  | 151                | 18                  | 0.755   |
| Deep incisional SSI                | 6                  | 0                   | 0.000   |
| Organ/space SSI                     | 2                  | 0                   | 0.000   |
| Unplanned intubation                | 4                  | 0                   | 0.000   |
| Pulmonary embolism                  | 3                  | 0                   | 0.000   |
| Ventilator dependence (> 48 h)     | 0                  | 0                   |         |
| Acute renal failure                 | 0                  | 0                   |         |
| CVA/stroke                          | 3                  | 2                   | 0.103   |
| Cardiac arrest                      | 2                  | 0                   | 0.000   |
| Myocardial infarction               | 1                  | 1                   | 0.215   |
| DVT                                 | 1                  | 0                   | 0.000   |
| Systemic sepsis                     | 2                  | 0                   | 0.000   |
| Septic shock                        | 0                  | 0                   |         |
| Unplanned reoperation               | 85                 | 8                   | 0.393   |
| Unplanned readmission               | 81                 | 11                  | 0.868   |
| Mortality                           | 5                  | 1                   | 0.517   |

SSI Surgical Site Infection, CVA Cerebral Vascular Accident, DVT Deep Venous Thromboembolism

Table 4 Multivariate logistic regression model analyzing regional anesthesia as a risk factor for complications

| Postoperative complications | OR    | 95% CI   | P value |
|-----------------------------|-------|----------|---------|
| Any complication            | 1.170 | 0.782    | 1.751   | 0.445   |
| Minor complication          | 1.803 | 0.907    | 3.587   | 0.093   |
| Major complication          | 0.925 | 0.565    | 1.516   | 0.758   |
| Unplanned reoperation       | 0.706 | 0.337    | 1.477   | 0.355   |
| Unplanned readmission       | 1.085 | 0.580    | 2.030   | 0.799   |
| Mortality                   | 1.789 | 0.230    | 13.912  | 0.579   |

similar results—patients who were given the hematoma block demonstrated significantly decreased pain intensity in the initial hours following fracture reduction and had reduced rates of intensive care unit admissions [12]. The present study, however, was unable to analyze functional outcome and pain scores at long-term follow-up as the ACS-NSQIP database does not record this information. The lack of long-term data and functional outcomes thus represents a limitation inherent to the database.

Although RA may avoid the risks of GA and provide adequate pain control in the immediate postoperative period, rebound pain is a known phenomenon associated with RA following various orthopaedic procedures [5, 31, 32]. Galos et al. report no differences in adverse events between patients who had received RA versus GA in the treatment of DRF [5]. However, the authors demonstrate that although GA is associated with greater immediate postoperative pain, peripheral nerve blocks are often associated with delayed rebound pain in the postoperative period [5]. In a retrospective quality improvement analysis, Sunderland et al. report increased rates of unplanned physician office visits following ORIF of DRFs when utilizing RA due to delayed rebound pain compared to GA [10]. As such, although RA may provide benefits by avoiding risks associated with GA in high-risk patient populations, thorough postoperative/ discharge analgesic medications should be implemented to curb rebound pain. As the ACS-NSQIP database does not record pain scores, the lack of data regarding patient pain levels is another limitation inherent to this study when comparing RA and GA for operative outcomes of DRFs.

Regarding peri-operative variables, the RA group had shorter days to operation from admission, total hospital stay length, and days from operation to discharge—however, these results were not statistically significant. Of note, RA had a significantly longer operative time (81.68 min) compared to GA (77.24 min). The cause for a longer operative time in the RA group is most likely multifactorial. Although the present study was able to control for differences in patient characteristics/comorbidities via CEM, only the variables/comorbidities that are maintained by the ACS-NSQIP database were controlled for. As this is a nationwide database, the granularity of information provided is limited. As such, the longer operative time in the RA group may be due to a sicker/more complex patient population that opted for RA who had variables that were unable to be controlled for (such as different grades of congestive heart failure or ascites). It is plausible that if RA was used for sicker/more complex patients, a longer operative time would be expected.

There are several additional limitations to this retrospective study. A limitation inherent to this study is the granularity of information provided by the ACS-NSQIP database. Although the database records the main mode of anesthetic utilized as discussed in the materials and methods, there is no data available as to what or if secondary/supplemental modes of anesthesia were utilized. As such, this represents a potential confounding variable that this study could not account for. In addition, the ACS-NSQIP database does not differentiate between types of RA, such as single-shot or continuous infusion of anesthetics. This could thus represent a systemic confounding variable that this study could...
Conclusion

The present study demonstrated no differences in adverse events/complications between RA and GA for ORIF of DRFs. With similar safety profiles, RA is a safe and reasonable alternative to GA when managing DRFs surgically. RA may be the preferred anesthetic technique for ORIF of DRFs in patients at high risk with GA. However, the current literature indicates an increase in instances of rebound pain when utilizing RA following ORIF of DRFs that this study could not comment on. Proper analgesic protocols must be put in place for postoperative/discharge planning in anticipation of delayed rebound pain with the use of RA to adequately control patients’ pain.

Supplementary Information  The online version contains supplementary material available at https://doi.org/10.1007/s00068-021-01704-1.

Funding  No benefits in any form have been received or will be received from a commercial party related directly or indirectly to the subject of this article.

Data availability  Publicly available through the American College of Surgeons National Surgical Quality Improvement Program Database.

Declarations

Conflict of interest  The authors declare that they have no conflict of interest.

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