Effects of Opioid-Limiting Legislation in the State of Ohio on Opioid Prescriptions After Shoulder Arthroscopy

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Background: Recent studies have shown that legislation regulating opioid prescriptions in the United States has been successful in reducing the morphine milligram equivalent (MME) prescribed after certain orthopaedic procedures.

Purpose: To (1) determine the effect of Ohio's legislation limiting opioid prescriptions after shoulder arthroscopy and (2) identify risk factors associated with prolonged opioid use and increased postoperative opioid dosing.

Study Design: Cohort study; Level of evidence, 3.

Methods: We reviewed the data of patients who underwent shoulder arthroscopy between January 1, 2016, and March 31, 2020. Patients were classified according to the date of legislation passage (August 31, 2017) as before legislation (PRE) or on/after legislation (POST). Patients were also classified based on the number of opioid prescriptions filled within 30 days of surgery as opioid-tolerant (at least 1 prescription) or opioid-naïve (zero prescriptions). We recorded patient characteristics, medical comorbidities, and surgical details, as well as the number of opioid prescriptions, MME per prescription from 30 days preoperatively to 90 days postoperatively, and the number of gamma-aminobutyric acid (GABA) analogues and benzodiazepine prescriptions from 30 days preoperatively to the date of surgery. Differences between cohorts were compared with the Fisher exact test and Wilcoxon test. A covariate-adjusted regression analysis was used to evaluate risk factors associated with increased postoperative opioid dosing.

Results: Overall, 279 patients (n = 97 PRE; n = 182 POST; n = 42 opioid-tolerant; n = 237 opioid-naïve) were included in the final analysis. There was a significant reduction in the cumulative MME prescribed in the immediate (0-7 days) postoperative period (PRE, 450 MME vs POST, 315 MME), the first 30 postoperative days (PRE, 590 MME vs POST, 375 MME), and the first 90 postoperative days (PRE, 600 MME vs POST, 420 MME) (P < .001 for all). The opioid-tolerant cohort had higher MME at every time point in the postoperative period (P < .001). Consumption of preoperative opioid ($\beta = 1682.5$; P < .001), benzodiazepine ($\beta = 468.09$; P < .001), and GABA analogue ($\beta = 251.37$; P = .04) was associated with an increase in the cumulative MME prescribed.

Conclusion: Opioid prescription–limiting legislation in Ohio significantly reduced the cumulative MME prescribed in the first 30 days postoperatively for both opioid-naïve and opioid-tolerant patients after shoulder arthroscopy. Consumption of opioids, benzodiazepines, and GABA analogues preoperatively was associated with increased postoperative opioid dosage.

Keywords: legislation; monitoring; opioids; postoperative; risk factors; shoulder arthroscopy

The use of opioids for nonmedical and recreational purposes has fueled a public health emergency in the United States over the past few decades. Between 2002 and 2011, approximately 25 million people throughout the country used opioid pain medications for nonmedical use.⁷ In 2016 alone, 11 million people in the United States misused opioids.⁴ This upward trend in consumption has led to a predictable increase in the mortality rate attributed to prescription opioid medications, which quadrupled⁶ between 2000 and 2014. Although many overdose cases are

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due to the consumption of illicit substances, including fentanyl and heroin, the prevalence of prescription opioid medications has also contributed to this issue.⁶

Policymakers in the United States have instituted reasonable prescriber practice changes in many states to reduce the prevalence of circulating opioids. Many states have installed prescription drug monitoring programs,¹³ while state-level legislation has been passed that regulates how opioids can be prescribed postoperatively. Ohio passed such legislation¹² on August 31, 2017. In Ohio, this legislation states that opioid prescriptions for acute pain, such as acute postoperative pain, cannot exceed a maximum dose per day of 30 morphine milligram equivalents (MMEs) and that they cannot be prescribed for more than 7 days for adults. Previous studies from various states have demonstrated that opioid prescription-limiting legislation is associated with significant reductions in the number and cumulative dosage of opioids prescribed after numerous orthopaedic procedures.^{3,13-17}

Although the success of opioid prescription-limiting legislation has been exhibited in the context of numerous orthopaedic procedures, few studies have evaluated the effect of such legislation in the context of shoulder arthroscopy. It is important to ascertain this effect for numerous reasons. Perioperative and postoperative opioid consumption have been associated with relatively poor functional outcomes after shoulder arthroscopy. In a retrospective review of 1242 patients who underwent shoulder arthroscopy. Lu et al⁹ identified preoperative opioid use as a significant predictor of reduced achievement of the Patient Acceptable Symptom State and increased likelihood of endorsing persistent pain. They also demonstrated that the risk of revision surgery increased with the consumption of high-dose opioids during the perioperative period. Likewise, Farley et al⁸ revealed that any patient using >1 oral morphine equivalent per day had increased rates of 3-year revision surgery, reoperations, hospital readmission, emergency department encounters, and infections after arthroscopic rotator cuff repair. In addition, in contrast to many arthroscopic procedures, arthroscopic shoulder surgery is associated with considerable postoperative pain.^{1,11,19}

The purpose of the present study was twofold. The first purpose was to determine the effect of Ohio's opioid prescription-limiting legislation on opioid prescribing practices after shoulder arthroscopy. The second purpose was to identify patient factors that are associated with long-term (ie, after 30 days postoperatively) opioid use and increased opioid dosing postoperatively. We hypothesized that there would be a decrease in the number of postoperative opioid prescriptions filled, a decrease in the MME prescribed in the immediate postoperative period (ie, prescriptions filled within 7 days postoperatively) and in the first 30 postoperative days but no significant difference in the MME prescribed in the first 90 postoperative days.

METHODS

Study Design and Data Collection

Approval for the study protocol was obtained from our institutional review board and the State of Ohio Board of Pharmacy. This retrospective cohort study included patients who underwent arthroscopic shoulder surgery between January 1, 2016, and March 31, 2020, by a single surgeon (R.J.G.) at an academic tertiary referral center. The Current Procedural Terminology codes 29805, 29806, 29807, 29822, 29823, 29825, 29826, and 23430 were used to identify potentially eligible participants. Patients who underwent arthroscopic rotator cuff repair were excluded because the procedure is associated with considerable postoperative pain when compared with other arthroscopic shoulder procedures.^{1,19} Likewise, we excluded arthroscopic distal clavicle excision from our cohort of patients because it is routinely performed as an open procedure at our institution. In addition, patients were excluded from the study if their information in the electronic medical record or the Ohio Automated Rx Reporting System (OARRS), the State of Ohio's Board of Pharmacy prescription drug monitoring program, was absent or incomplete.

Patients were categorized into 1 of the 2 cohorts based on their date of surgery. Those patients who underwent shoulder arthroscopy before August 31, 2017, (the date the opioid prescription–limiting legislation was passed) were placed into the prelegislation (PRE) cohort, while patients who underwent surgery on or after August 31, 2017, were placed into the postlegislation (POST) cohort. In addition, patients were classified as opioid-naïve if they did not have an opioid prescription prescribed within 30 days before surgery and as opioid-tolerant if they had at least 1 opioid prescription prescribed within the 30 days before surgery. This is in accordance with previously published studies¹⁴⁻¹⁷ that used the 30-day cutoff as per the

Ethical approval for this study was obtained from Case Western Reserve University, Cleveland, Ohio, USA (No. STUDY20200623).

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regulations of the Department of Health in the State of Rhode Island.

The patients' electronic medical records were reviewed, and information on age, sex, body mass index (BMI), laterality of the procedure, medical comorbidities, and preoperative medication/substance use were extracted (ie, tobacco, alcohol). In addition, a query was performed on the OARRS database, which is a 5-year rolling database; thus, data before the 5-year period were automatically expunged daily from the system. Data regarding controlled substance prescriptions were obtained for all patients from 30 days preoperatively to 90 days postoperatively. Information was obtained regarding the type and dose of oral opioids, the number of prescriptions, and the number of pills per prescription. To account for the relative potency of each opioid, the dosage was converted into MME using the following multipliers¹⁰: oxycodone, number of milligrams \times 1.5; tramadol, number of milligrams \times 0.1; and hydromorphone, number of milligrams \times 4. No multipliers were applied to morphine or hydrocodone. The new dosing value was multiplied by the number of tablets within each prescription to obtain the MME for each prescription. The cumulative MME, or the sum of the MME for each prescription, prescribed in the following postoperative periods were calculated: immediate (first 7 days); 0 to 30 days; 31 to 60 days; and 61 to 90 days. In addition, data were collected regarding the number of benzodiazepines (eg, diazepam, alprazolam, and chlordiazepoxide) and gammaaminobutvric acid (GABA) analogue (eg. gabapentin, pregabalin) prescriptions that were prescribed within 30 days before surgery.

Statistical Analysis

The patients' sociodemographic variables were stratified by time with respect to the passage of legislation (PRE vs POST) using frequency and percentage for categorical variables and medians and interquartile ranges (IQRs) for continuous variables. Differences between PRE and POST data were compared using the Fisher exact test for categorical variables and Wilcoxon test for continuous variables. The Wilcoxon test was also used to distinguish intracohort measures for the opioid-tolerant versus opioid-naïve groups. The Kruskal-Wallis equality of populations rank test was applied to assess the interaction between both opioid use and legislation.

An average treatment effect (ATE) regression model was used to determine the effect of the opioid prescription-limiting legislation on the cumulative MME prescribed postoperatively, with inverse-probability weighting to correct the imbalance in the distributions of the patient characteristics and surgical procedures on the potential outcomes before and after the enactment of the legislation. We performed a covariate-adjusted regression analysis to evaluate risk factors associated with increased postoperative opioid dosing. In all regression models, the variables of interest were adjusted or controlled for age,

TABLE 1Patient Characteristics According toPre- and Postlegislation Cohorts $(N = 279)^a$

Variable	PRE $(n = 97)$	POST (n = 182)	P
Age, y	47 [33-57]	49.50 [32-56.8]	.819
Male sex	45 (46.4)	94 (51.6)	.451
BMI, kg/m ²	27.46 [23.8-34.4]	29.30 [24.7-33.6]	.281
Right shoulder involved	59 (60.8)	96 (52.7)	.208
Diabetes			
Controlled by diet	5 (5.2)	2(1.1)	.052
On oral medication	8 (8.2)	9 (4.9)	.299
On insulin	2(2.1)	14 (7.7)	.061
Alcohol use	18 (18.6)	68 (37.4)	.001
Tobacco use	25 (25.8)	47 (25.8)	>.999
History of chronic pain	8 (8.2)	9 (4.9)	.299
Hypertension	32 (33)	59 (32.4)	>.999
Heart disease	18 (18.6)	20 (11)	.099
Pulmonary disease	20 (20.6)	33 (18.1)	.633
GABA-analogue	6 (6.2)	14 (7.7)	.809
Benzodiazepine	10 (10.3)	12 (6.6)	.351
Opioid-tolerant	17 (17.5)	25 (13.7)	.482

^aData are reported as median [interquartile range] or n (%). The bold *P* value indicates a statistically significant difference between groups (P < .05). BMI, body mass index; GABA, gamma-aminobutyric acid; POST, postlegislation; PRE, prelegislation.

sex, BMI, and date of legislation passage (August 31, 2017). Finally, a post hoc power analysis was conducted, which was powered to the cumulative MME in the first 30 days based on our ATE analysis.

All statistical tests were 2-sided, and statistical significance was defined as P < .05. The statistical analyses were performed using Stata 15.0 and R 3.4.1.

RESULTS

Overall, 279 patients were selected for this study, with 97 and 182 patients placed into the PRE and POST cohorts, respectively (Table 1). The percentage of patients who reported that they consumed any amount of alcohol was the only significant difference between the 2 cohorts (PRE, 18.6% vs POST, 37.4%; P = .001).

Effect of Legislation on Postoperative Opioid Prescription

There was a significant reduction in the median [IQR] cumulative MME prescribed in the immediate postoperative period (PRE, 450 [450-600] MME vs POST, 315 [210-450] MME; P < .001), the first 30 postoperative days (PRE, 590 [450-915] MME vs POST, 375 [210-546] MME; P < .001), and first 90 postoperative days (PRE, 600 [450-1200] MME vs POST, 420 [210-660] MME; P < .001) (Table 2). There was also a significant reduction in the median number of opioid prescriptions within the first 30 postoperative days between the PRE and POST cohorts

TABLE 2
Cumulative Opioid Prescriptions According to Pre- and Postlegislation Cohorts ^a

	-		
PRE $(n = 97)$	POST (n = 182)	Р	
2 [1-2]	1 [1-2]	<.001	
	015 [010 (50]	0.01	
450 [450-600]	315 [210-450]	<.001	
590 [450-915]	375[210-546]	<.001	
450 [300-810.1]	475 [300-900]	.762	
450 [262.5-615]	490 [210-1031.3]	.520	
600 [450-1200]	420 [210-660]	<.001	
	$\begin{array}{c} 2 \ [1-2] \\ 450 \ [450-600] \\ 590 \ [450-915] \\ 450 \ [300-810.1] \\ 450 \ [262.5-615] \end{array}$	2 [1-2] 1 [1-2] 450 [450-600] 315 [210-450] 590 [450-915] 375 [210-546] 450 [300-810.1] 475 [300-900] 450 [262.5-615] 490 [210-1031.3]	

^{*a*}Data are reported as median [interquartile range]. Bold *P* values indicate statistically significant differences between groups (P < .05). MME, morphine milligram equivalent; POST, postoperative; PRE, preoperative.

TABLE 3
Average Treatment Effects of the Opioid Prescription-
Limiting Legislation on MME and Prescriptions ^a

	ATE	Robust SE	P
No. of 30-day postop prescriptions Cumulative MME	-0.38	0.11	.001
Immediate postop	-149.34	26.27	<.001
0-30 days postop	-248.93	59.18	<.001
31-60 days postop	-104.22	222.40	.63
61-90 days postop	170.41	144.98	.24
0-90 days postop	-343.43	134.50	.01

^{*a*}Bold *P* values indicate statistical significance (P < .05). ATE, average treatment effect; MME morphine milligram equivalent; Postop, postoperative.

(PRE, 2 [1-2] prescriptions vs POST, 1 [1-2] prescription; P < .001).

The ATEs of the opioid prescription–limiting legislation on cumulative postoperative MMEs are shown in Table 3. The legislation was associated with a significant reduction in the cumulative MME prescribed in the immediate postoperative period (ATE, -149.34 MME; P < .001), 0-30 days postoperative (ATE, -248.93 MME; P < .001), and 0-90 days postoperative (ATE, -343.43 MME; P = .01). There was a significant reduction in the number of opioid prescriptions in the 30-day postoperative period (ATE, -0.38prescriptions; P = .001).

Effect of Preoperative Opioid Tolerance Status on Postoperative Opioid Prescriptions

Table 4 illustrates the effects of the preoperative opioid use status (ie, opioid-naïve vs opioid-tolerant) on postoperative opioid prescriptions. The median [IQR] cumulative MME was significantly less in the opioid-naïve cohort compared with the opioid-tolerant cohort in the immediate postoperative period (tolerant, 450 [375-600] MME vs naïve, 337.5 [210-450] MME; P < .001), the first 30 postoperative days (tolerant, 1031.3 [595-2025] MME vs naïve, 435 [300-600] MME; P < .001), 31-60 postoperative days (tolerant, 840 [450-1800] MME vs naïve, 300 [175-450] MME;

P < .001), 61-90 postoperative days (tolerant, 703.5 [450-1350] MME vs naïve, 275 [210-451.8] MME; P < .001), and 0-90 postoperative days (tolerant, 2218.5 [1012.6-4162.5] MME vs naïve, 450 [300-600] MME; P < .001).

Combined Effect of Legislation and Preoperative Opioid Status on Postoperative Opioid Prescriptions

Opioid prescription patterns stratified by the legislation period and preoperative opioid tolerance status are shown in Table 5. Within the opioid-naïve cohort, 80 patients were placed into the PRE subcohort, and 157 patients were placed into the POST subcohort. In the opioid-tolerant cohort, 17 patients were placed into the PRE subcohort, and 25 patients were placed into the POST subcohort. There was a significant reduction in the median [IQR] cumulative MME prescribed in the immediate postoperative period (PRE, 450 [450-600] MME vs POST, 315 [210-420] MME; P < .001), first 30-days postoperatively (PRE, 450 [450-750] MME vs POST, 315 [210-450] MME; P < .001), and 0-90 days postoperatively (PRE, 525 [450-843.8] MME vs POST, 315 [210-450] MME; P < .001) between the PRE and POST opioid-naïve cohorts. Likewise, there was a significant reduction in the median cumulative MME prescribed in the immediate postoperative period (PRE, 525 [450-600] MME vs POST, 435 [315-450] MME; P = .014) and first 30 days postoperatively (PRE, 1500 [975-2722.5] MME vs POST, 855 [450-1800] MME; P = .021) between the PRE and POST opioidtolerant cohorts.

Risk Factors for Increased Postoperative Opioid Use

The risk factors for increased postoperative opioid dosing are shown in Table 6. Preoperative opioid tolerance was an independent risk factor for increased cumulative MME at 0-30 postoperative days ($\beta = 705.33$; P < .001), 0-90 postoperative days ($\beta = 1682.53$; P < .001), and 31-90 postoperative days (odds ratio,16.67; P < .001). In addition, preoperative GABA analogue prescription was a risk factor for increased cumulative MME in the 0-90 days postoperatively ($\beta = 251.37$; P = .04), while preoperative benzodiazepine prescription was a risk factor for increased cumulative MME in the 0-90 days postoperatively ($\beta = 251.37$; P = .04), while preoperative benzodiazepine prescription was a risk factor for increased cumulative MME in

Opio	spiola Trescription Data According to Opiola Tolerance			
	Opioid Naïve (n = 237)	Opioid Tolerant (n = 42)	Р	
No. of 30-day postop prescriptions	1 [1-2]	2 [1-4]	<.001	
Cumulative MME				
Immediate postop	337.5 [210-450]	450 [375-600]	<.001	
0-30 days postop	435 [300-600]	1031.3 [595-2025]	<.001	
31-60 days postop	300 [175-450]	840 [450-1800]	<.001	
61-90 days postop	275[210-451.8]	703.5 [450-1350]	<.001	
0-90 days postop	450 [300-600]	2218.5 [1012.6-4162.5]	<.001	

^{*a*}Data are reported as median [interquartile range]. Bold *P* values indicate statistically significant differences between groups (P < .05). MME, morphine milligram equivalent; postop, postoperative.

TABLE 5	
Opioid Prescription Data According to Both Opioid Tolerance an	nd Legislation ^a

	Opioid-Naïve			Opioid-Tolerant		
	PRE $(n = 80)$	POST (n = 157)	Р	PRE (n = 17)	POST $(n = 25)$	Р
No. of 30-day postop prescriptions Cumulative MME	1 [1-2]	1 [1-2]	.009	3 [2-4]	2 [1-3]	.036
Immediate postop	450 [450-600]	315 [210-420]	<.001	525 [450-600]	435 [315-450]	.014
0-30 days postop	450 [450-750]	315 [210-450]	<.001	1500 [975-2722.5]	855 [450-1800]	.021
31-60 days postop	300 [150-450]	307.5 [250-537.5]	.663	600 [465-1800]	870 [450-1800]	.885
61-90 days postop	262.5 [145-450]	293.8 [210-494.9]	.328	615 [465 - 1349.8]	900 [450-1575]	.797
0-90 days postop	525 [450-843.8]	315 [210-450]	<.001	2775.1 [1760-4972.5]	1659.9 [900-3890.1]	.098

^{*a*}Data are reported as median [interquartile range]. Bold P values indicate statistically significant differences between groups (P < .05). MME, morphine milligram equivalent; postop, postoperative.

TABLE 6 Results of Covariate-adjusted Regression Analysis Evaluating Independent Risk Factors for Postoperative Opioid Prescribing^a

	MME 0-30 Days $Postop^b$		MME 0-90 Days Postop ^{b}		MME 31-90 Days $Postop^c$	
	β (SE)	Р	β (SE)	P	OR (SE)	Р
Opioid-tolerant	705.33 (59.44)	<.001	1682.53 (115.12)	<.001	16.67 (7.36)	<.001
Preop GABA analogue prescription	130.11 (89.50)	.15	$251.37\ (123.53)$.04	2.19 (1.08)	.11
Preop benzodiazepine prescription	$277.14\ (85.28)$.001	468.09 (106.8)	<.001	2.39 (1.16)	.07

 a Bold P values indicate statistical significance (P < .05). GABA, gamma-aminobutyric acid; MME, morphine milligram equivalent; OR, odds ratio; Postop, postoperative; Preop, preoperative.

^bCoefficients from quantile regression analysis for the outcome of interest: MME intake up to first 30 days; MME intake 0 to 90 days. ^cOR from logistic regression analysis of prolonged opioid use (yes or no) in the 31- to 90-day postoperative period. In all regression models,

the variables of interest were adjusted or controlled for age, sex, body mass index, and date of legislation passage.

the 0-30 days postoperatively (β = 277.14; *P* = .001) and 0-90 days postoperatively (β = 106.8; *P* < .001).

Results of Post Hoc Power Analysis

Based on our ATE analysis, the difference in the cumulative MME in the first 30 days is 248.93 ± 59.18 . Using this MME value, an alpha of .05, and a 2-sided 2-sample equal variance z test, we determined that the sample sizes of 97 and 182 for the PRE and POST cohorts, respectively, would achieve a power of >99%.

DISCUSSION

The major findings of our study demonstrate that there were significant reductions in the cumulative MME

prescribed in the immediate (PRE, 450 MME; POST, 315 MME), first 30-day (PRE, 590 MME; POST, 375 MME), and first 90-day (PRE, 600 MME; POST, 420 MME) postoperative periods after shoulder arthroscopy between the prelegislation and postlegislation cohorts, respectively (P < .001 for all). However, the reduction in cumulative MME prescribed in the 0-90 days postoperative periods was largely attributed to the reduction observed in the 0-30 days postoperative periods because no significant decreases were observed in the 31-60 days and 61-90 days postoperative periods. We also identified preoperative opioid, GABA-analogue, and benzodiazepine consumption as independent risk factors for increased cumulative MME prescribed at various time points throughout the postoperative period. Interestingly, the POST cohort had a significantly higher number of patients who consumed any amount of alcohol relative to the PRE cohort (PRE, 18.6% vs POST 37.4%; P = .001), yet the POST cohort received fewer cumulative MME postoperatively relative to the PRE cohort within the first 90 days postoperatively. Therefore, it is certainly possible that these groups are not entirely comparable because of this significant difference.

Our results are congruent with previously published studies that examined the effect of similar legislation on opioid-prescribing practices after shoulder surgery.^{14,18} In a retrospective review of 446 patients who underwent arthroscopic rotator cuff repair in Rhode Island, Shah et al¹⁸ highlighted a significant reduction in the number of opioid prescriptions filled within the first 30 days postoperatively and the cumulative MME prescribed in the first postoperative prescription as well as the first 30 days postoperatively between their PRE and POST cohorts. Similarly, in an analysis of 334 patients who underwent shoulder arthroplasty in Ohio, Raji et al¹⁴ demonstrated significant reductions in the cumulative MME prescribed in the 0-7 days and 0-30 days postoperative periods when comparing their PRE and POST cohorts.

We could not demonstrate a significant decrease in the cumulative MME prescribed in the 31-60 days or 61-90 days postoperative periods. These results are comparable to previously published studies. Reid et al were able to show significant decreases in the cumulative MME prescribed during the first 30 days after lumbar spine surgery¹⁵ and total joint arthroplasty¹⁷ but not within 30 to 90 days after these procedures in Rhode Island. Together, these findings suggest the laws in Ohio and Rhode Island are not likely to decrease chronic opioid use. However, these laws still provide a larger societal benefit by reducing the number of circulating opioids and thus potentially decreasing the risk of opioid diversion, a practice in which legal prescription drugs are distributed illegally for use not intended by the prescriber.

Reducing the prevalence and accessibility to opioid medications after orthopaedic surgery is critical in the effort to curtail the opioid epidemic in the United States as well as to improve patient outcomes postoperatively. Increased opioid consumption after hip arthroscopy for femoroacetabular impingement syndrome is associated with significantly worse Harris Hip scores and visual analog satisfaction and pain scores.^{2,20} With respect to shoulder arthroscopy, preoperative opioid use is a significant predictor of worse outcomes postoperatively,⁹ while the consumption of ≥ 1 oral morphine equivalent was associated with increased rates of revision surgery, reoperations, hospital readmission, emergency department encounters, and infection.⁸ Future studies are warranted to examine the impact of opioid prescription–limiting legislation on clinical outcomes, including reoperations, hospital readmissions, and emergency department utilization.

Limitations

The outcomes of this study must be interpreted within its limitations. First, although the post hoc power analysis demonstrated that our study was adequately powered with our given sample sizes, we acknowledge that our patient population was relatively small. We attribute this to an inherent flaw of the OARRS database. Although the OARRS database tracks the prescription of all scheduled medications within Ohio, the database can be incomplete and/or inaccurate. Second, the results of this study ostensibly reflect the experience of a single surgeon at a single, academic, tertiary referral center. Third, the OARRS database tracks written and filled opioid prescriptions, but it does not track opioid consumption. It is reasonable to conclude that patients consumed less opioids than what was originally prescribed and either disposed of or diverted any unused medication. However, a recent survey of patients receiving opioids for pain management reported that only 33% of patients disposed of their unused opioid medication.⁵ Therefore, we believe that filled opioid prescriptions serve as a valid metric for opioid consumption. Fourth, the OARRS database does not track inpatient opioid prescription or consumption patterns. Although many, if not all, patients within this cohort underwent surgery on an outpatient basis, they still received opioids intraoperatively and during their brief stay in the postanesthesia care unit.

CONCLUSION

Opioid prescription-limiting legislation in Ohio significantly reduced the cumulative MME prescribed in the first 30 days postoperatively for both opioid-naïve and opioidtolerant patients after shoulder arthroscopy. Consumption of opioids, benzodiazepines, and GABA analogues preoperatively was associated with a larger cumulative MME prescribed postoperatively. The results of this study demonstrate that such legislation in Ohio is associated with a significant reduction in the amount of MME prescribed after shoulder arthroscopy.

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