Minimizing Opioids After Joint Operation: Protocol to Decrease Postoperative Opioid Use After Primary Total Knee Arthroplasty

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Background: For decades, opioids have been the mainstay in pain management after total joint arthroplasty despite evidence that their use should be curtailed. To limit unnecessary prescribing of opioids, the US Department of Veterans Affairs (VA) Portland Health Care System Total Joints Service implemented the Minimizing Opioids After Joint Operation (MOJO) postoperative pain protocol in 2018 to reduce opioid use following total knee arthroplasty (TKA). This protocol included reductions of inpatient and outpatient opioid prescribing, preoperative optimization, use of perioperative nerve blocks, and surgery without a tourniquet.

Methods: We performed a retrospective chart review that compared the first 20 consecutive patients undergoing TKA using the MOJO protocol with the last 20 patients using the prior routine. Outcomes included total inpatient opioid use, daily opioid use, emergency department (ED) visits or readmissions within

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or decades, opioids have been a mainstay in the management of pain after total joint arthroplasty. In the past 10 years, however, opioid prescribing has come under increased scrutiny due to a rise in rates of opioid abuse, pill diversion, and opioid-related deaths.^{1,2} Opioids are associated with adverse effects, including nausea, vomiting, constipation, apathy, and respiratory depression, all of which influence arthroplasty outcomes and affect the patient experience. Although primary care groups account for nearly half of prescriptions written, orthopedic surgeons have the third highest per capita rate of opioid prescribing of all medical specialties.^{3,4} This puts orthopedic surgeons, particularly those who perform routine procedures, in an opportune but challenging position to confront this problem through novel pain management strategies.

Approximately 1 million total knee arthroplasties (TKAs) are performed in the US every year, and the US Department of Veterans Affairs (VA) health system performs about 10,000 hip and knee joint replacements.^{5,6} There is no standardization of opioid prescribing in the postoperative period following these procedures, and studies have reported a wide varia-

90 days, phone calls for pain or medication refills, length of stay (LOS), and pain during inpatient hospital stay.

Results: There were significant differences between the pre-MOJO and the MOJO groups with regard to daily inpatient morphine equivalent dose (MED) (82 mg vs 31 mg, P < .01) and total inpatient MEDs (306 mg vs 33 mg, P < .01). There was less self-reported pain on postoperative day 1 in the MOJO group (5.5 vs 4.1, P = .01), decreased LOS (4.4 days vs 1.1 days, P < .01), fewer total ED visits (6 vs 2, P < .07), and fewer discharges to skilled nursing facilities (12 vs 0, P < .01).

Conclusions: The MOJO protocol reduced postoperative opioid use after TKA in the VA setting without compromising pain control or increasing ED visits. The framework and routines described are potentially applicable to other institutions and surgical specialties.

tion in prescribing habits even within a single institution for a specific surgery.⁷ Patients who undergo TKA are at particularly high risk of long-term opioid use if they are on continuous opioids at the time of surgery; this is problematic in a VA patient population in which at least 16% of patients are prescribed opioids in a given year.⁸ Furthermore, veterans are twice as likely as nonveterans to die of an accidental overdose.⁹ Despite these risks, opioids remain a cornerstone of postoperative pain management both within and outside of the VA.¹⁰

In 2018, to limit unnecessary prescribing of opioid pain medication, the total joint service at the VA Portland Health Care System (VAPHCS) in Oregon implemented the Minimizing Opioids after Joint Operation (MOJO) postoperative pain protocol. The goal of the protocol was to reduce opioid use following TKA. The objectives were to provide safe, appropriate analgesia while allowing early mobilization and discharge without a concomitant increase in readmissions or emergency department (ED) visits. The purpose of this retrospective chart review was to compare the efficacy of the MOJO protocol with our historical experience and report our preliminary results.

METHODS

Institutional review board approval was obtained to retrospectively review the medical records of patients who had undergone TKA surgery during 2018 at VAPHCS. The MOJO protocol was composed of several simultaneous changes. The centerpiece of the new protocol was a drastic decrease in routine prescription of postoperative opioids (Table 1). Other changes included instructing patients to reduce the use of preoperative opioid pain medication 6 weeks before surgery with a goal of no opioid consumption, perform daily sets of preoperative exercises, and attend a preoperative consultation/ education session with a nurse coordinator to emphasize early recovery and discharge. In patients with chronic use of opioid pain medication (particularly those for whom the medication had been prescribed for other sources of pain, such as lumbar back pain), the goal was daily opioid use of \leq 30 morphine equivalent doses (MEDs). During the inpatient stay, we stopped prescribing prophylactic pain medication prior to physical therapy (PT).

We encouraged preoperative optimization of muscle strength by giving instructions for 4 to 8 weeks of daily exercises (Appendix). We introduced perioperative adductor canal blocks (at the discretion of the anesthesia team) and transitioned to surgery without a tourniquet. Patients in both groups received intraoperative antibiotics and IV tranexamic acid (TXA); the MOJO group also received topical TXA.

Further patient care optimization included providing patients with a team-based approach, which consisted of nurse coordinators, physician assistants and nurse practitioners, residents, and the attending surgeon. Our team reviews the planned pain management protocol, perioperative expectations, criteria for discharge, and anticipated surgical outcomes with the patient during their preoperative visits. On postoperative day 1, these members round as a team to encourage patients in their immediate postoperative recovery and rehabilitation. During rounds, the team assesses whether the patient meets the criteria for discharge, adjusting the pain management protocol if necessary.

Changes in surgical technique included arthrotomy with electrocautery, minimizing

TABLE 1 Comparison of Postoperative PainManagement Protocols Before and After Implementationof the MOJO Protocol

Medication Types	Postoperative I Before MOJO	Pain Protocol MOJO				
Opioids	 Oxycodone 10 mg or hydrocodone 10 mg by mouth every 4-6 h as needed for 6-8 wk; start day 0 Hydromorphone 0.2-0.4 mg or morphine 2.5 mg IV for break- through pain while inpatient only; start day 0 	Hydrocodone 10/325 mg by mouth every 8 h as needed for 14 d; start day 0				
Nonopioids	Tylenol 650 mg by mouth every 6 h for 8-12 wk; start day 0	 Tylenol 975 mg by mouth 3 times a day; start day 0 for 14 d Tramadol 50-100 mg by mouth every 8 h as needed for 14 d; start day 0 Celebrex 200 mg by mouth twice daily; start day 1 for 14 d 				
Muscle relaxants	Diazepam 10 mg by mouth every 8 h; start day 0 with 2 doses	 Diazepam 10 mg by mouth every 8 h, 2 doses; start day 0 Cyclobenzaprine 10 mg by mouth every 8 h as needed; start day 1 at 6 am for 14 d 				

Abbreviation: MOJO, Minimizing Opioids After Joint Operation protocol.

traumatic dissection or resection of the synovial tissue, and intra-articular injection of a cocktail of ropivacaine 5 mg/mL 40 mL, epinephrine 1:1,000 0.5 mL, and methylprednisolone sodium 40 mg diluted with normal saline to a total volume of 120 mL.

The new routine was gradually implemented beginning January 2017 and fully implemented by July 2018. This study compared the first 20 consecutive patients undergoing primary TKA after July 2018 to the last 20 consecutive patients undergoing primary TKA prior to January 2017. Exclusion criteria included bilateral TKA, death before 90 days, and revision as the indication for surgery. The senior attending surgeon performed all surgeries using a standard midline approach. The majority of surgeries were performed using a cemented Vanguard total knee system (Zimmer Biomet); 4 patients in the historical group had a NexGen knee system, cementless monoblock tibial components (Zimmer Biomet); and 1 patient had a Logic knee system (Exactech). Surgical selection criteria for patients did not differ between groups.

Electronic health records were reviewed

TABLE 2 Patient Characteristics

Characteristics	Before MOJO	MOJO	P Value
Age, y	65.3	71.9	.04
Male gender, No. (%)	17 (85)	18 (90)	.99
Body mass index, mean (SD)	34.2 (4.2)	30.2 (4.4)	.006
American Society of Anesthesiologists score, mean (SD)	2.8 (0.4)	2.9 (0.3)	.39
Risk Assessment and Predictive Tool score, mean (SD)	9.5 (2.5)	9.9 (1.4)	.53
Operating room time, mean (SD), min	141.6 (25.3)	130.1 (17.2)	.11

Abbreviation: MOJO, Minimizing Opioids After Joint Operation protocol.

and data were abstracted. The data included demographic information (age, gender, body mass index [BMI], diagnosis, and procedure), surgical factors (American Society of Anesthesiologists score, Risk Assessment and Predictive Tool score, operative time, tourniquet time, estimated blood loss), hospital factors (length of stay [LOS], discharge location), postoperative pain scores (measured on postoperative day 1 and on day of discharge), and postdischarge events (90-day complications, telephone calls reporting pain, reoperations, returns to the ED, 90-day readmissions).

The primary outcome was the mean postoperative daily MED during the inpatient stay. Secondary outcomes included pain on postoperative day 1, pain at the time of discharge, LOS, hospital readmissions, and ED visits within 90 days of surgery. Because different opioid pain medications were used by patients postoperatively, all opioids were converted to MED prior to the final analysis. Collected patient data were de-identified prior to analysis.

Power analysis was conducted to determine whether the study had sufficient population size to reject the null hypothesis for the primary outcome measure. Because practitioners controlled postoperative opioid use, a Cohen's d of 1.0 was used so that a very large effect size was needed to reach clinical significance. Statistical significance was set to 0.05, and patient groups were set at 20 patients each. This yielded an appropriate power of 0.87. Population characteristics were compared between groups using *t* tests and χ^2 tests as appropriate. To analyze the primary outcome, comparisons were made between the 2 cohorts using 2-tailed *t* tests. Secondary outcomes were compared between groups using *t* tests or χ^2 tests. All statistics were performed using R version 3.5.2. Power analysis was conducted using the package pwr.¹¹ Statistical significance was set at *P* < .05.

RESULTS

Forty patients met the inclusion criteria, evenly divided between those undergoing TKA before and after instituting the MOJO protocol (Table 2). A single patient in the MOJO group died and was excluded. A patient who underwent bilateral TKA also was excluded. Both groups reflected the male predominance of the VA patient population. MOJO patients tended to have lower BMIs (34 vs 30, P < .01). All patients indicated for surgery with preoperative opioid use were able to titrate down to their preoperative goal as verified by prescriptions filled at VA pharmacies. Twelve of the patients in the MOJO group received adductor canal blocks.

Results of *t* tests and χ^2 tests comparing primary and secondary endpoints are listed in Table 3. Differences between the daily MEDs given in the historical and MOJO groups are shown. There were significant differences between the pre-MOJO and MOJO groups with regard to daily inpatient MEDs (82 mg vs 29 mg, P < .01) and total inpatient MEDs (306 mg vs 32 mg, *P* < .01). There was less self-reported pain on postoperative day 1 in the MOJO group (5.5 vs 3.9, P < .01), decreased LOS (4.4 days vs 1.2 days, P < .01), a trend toward fewer total ED visits (6 vs 2, P = .24), and fewer discharges to skilled nursing facilities (12 vs 0, P < .01). There were no blood transfusions in either group.

There were no readmissions due to uncontrolled pain. There was 1 readmission for shortness of breath in the MOJO group. The patient was discharged home the following day after ruling out thromboembolic and cardiovascular events. One patient from the control group was readmitted after missing a step on a staircase and falling. The patient sustained a quadriceps tendon rupture and underwent primary suture repair.

DISCUSSION

Our results demonstrate that a multimodal approach to significantly reduce postoperative opioid use in patients with TKA is possible without increasing readmissions or ED visits for pain control. The patients in the MOJO group had a faster recovery, earlier discharge, and less use of postoperative opioid medication. Our approach to postoperative pain management was divided into 2 main categories: patient optimization and surgical optimization.

Patient Selection

Besides the standard evaluation and optimization of patients' medical conditions, identifying and optimizing at-risk patients before surgery was a critical component of our protocol. Managing postoperative pain in patients with prior opioid use is an intractable challenge in orthopedic surgery. Patients with a history of chronic pain and preoperative use of opioid medications remain at higher risk of postoperative chronic pain and persistent use of opioid medication despite no obvious surgical complications.⁸ In a sample of > 6,000 veterans who underwent TKA at VA hospitals in 2014, 57% of the patients with daily use of opioids in the 90 days before surgery remained on opioids 1 year after surgery (vs 2 % in patients not on long-term opioids).⁸ This relationship between pre- and postoperative opioid use also was dose dependent.12

Furthermore, those with high preoperative use may experience worse outcomes relative to the opioid naive population as measured by arthritis-specific pain indices.¹³ In a well-powered retrospective study of patients who underwent elective orthopedic procedures, preoperative opioid abuse or dependence (determined by the International Classification of Diseases, Ninth Revision diagnosis) increased inpatient mortality, aggregate morbidity, surgical site infection, myocardial infarction, and LOS.14 Preoperative opioid use also has been associated with increased risk of ED visits, readmission, infection, stiffness, and aseptic revision.¹⁵ In patients with TKA in the VA specifically, preoperative opioid use (> 3 months in the prior year) was associated with increased revision rates that were even higher than those for patients with diabetes mellitus.¹⁶

TABLE 3 Comparison of Primary and Secondary Endpoints in Treatment Groups

Variables	Before MOJO	MOJO	P Value	
Morphine equivalent daily dose, mean (SD)	81.7 (44.3)	28.7 (24.7)	< .001	
Length of stay, mean (SD), d	4.4 (2.8)	1.2 (0.4)	< .001	
Pain at baseline, mean (SD)	5.5 (1.7)	3.9 (1.8)	.006	
Pain at discharge, mean (SD)	5.1 (1.9)	3.9 (1.9)	.06	
Discharged to skilled nursing facilities, No. (%)	12 (60)	0 (0)	< .001	
Phone call for pain or pain medication, No. (%)	1 (5)	1 (5)	.99	
Readmission in 90 d, No. (%)	1 (5)	1 (5)	.99	
ED visit for pain, No. (%)	1 (5)	1 (5)	.99	
ED visit for any reason, No. (%)	6 (30)	2 (10)	.24	

Abbreviations: ED, emergency department, MOJO, Minimizing Opioids After Joint Operation protocol.

Patient Education

Based on this evidence, we instruct patients to reduce their preoperative opioid dosing to zero (for patients with joint pain) or < 30 MED (for patients using opioids for other reasons). Although preoperative reduction of opioid use has been shown to improve outcomes after TKA, pain subspecialty recommendations for patients with chronic opioid use recommend considering adjunctive therapies, including transcutaneous electrical nerve stimulation, cognitive behavioral therapy, gabapentin, or ketamine.17,18 Through patient education our team has been successful in decreasing preoperative opioid use without adding other drugs or modalities.

Patient Optimization

Preoperative patient optimization included 4 to 8 weeks of daily sets of physical activity instructions (prehab) to improve the musculoskeletal function. These instructions are given to patients 4 to 8 weeks before surgery and aim to improve the patient's balance, mobility, and functional ability (Appendix). Metaanalysis has shown that patients who undergo preoperative PT have a small but statistically significant decrease in postoperative pain at

Do 10 repetitions of each exercise 3 times each day.	2 = able to do 3 sets of 10 1 = able to do 3 sets of 5 0 = doing less than 3 sets of 5						
Week No.		т	w	Th	F	Sat	Sun
1. Turn in bed and sit up. Do it on the right and left sides							
2. Get out of bed and stand up from your bed							
3. Walk (use walker/cane if needed) for 50 yards (150 feet)							
4. Sit down and stand up from a chair							
5. Climb 3 stairs and use the railing							
6. Stand 3 minutes at a time (use walker/cane if needed)							
Total							

APPENDIX Prehabilitation Exercises Before Surgery

4 weeks, though this does not persist beyond that period. $^{19}\,$

We did note a lower BMI in patients in the MOJO group. Though this has the potential to be a confounder, a study of BMI in > 4,000 patients who underwent joint replacement surgery has shown that BMI is not associated with differences in postoperative pain.²⁰

Surgeon and Surgical-Related Variables

Patients in the MOJO group had increased use of adductor canal blocks. A 2017 metaanalysis of 12,530 patients comparing analgesic modalities found that peripheral nerve blocks targeting multiple nerves (eg, femoral/sciatic) decreased pain at rest, decreased opioid consumption, and improved range of motion postoperatively.²¹ Also, these were found to be superior to single nerve blocks, periarticular infiltration, and epidural blocks.²¹ However, major nerve and epidural blocks affecting the lower extremity may increase the risk of falls and prolong LOS.^{22,23} The preferred peripheral block at VAPHCS is a single shot ultrasound-guided adductor canal block before the induction of general or spinal anesthesia. A randomized controlled trial has demonstrated superiority of this block to the femoral nerve block with regard to postoperative quadriceps strength, conferring the theoretical advantage of decreased fall risk and ability to participate in immediate PT.²⁴ Although we are unable to confirm an association between anesthetic

modalities and opioid burden, our clinical impression is that blocks were effective at reducing immediate postoperative pain. However, among MOJO patients there were no differences in patients with and without blocks for either pain (4.2 vs 3.8, P = .69) or opioid consumption (28.8 vs 33.0, P = .72) after surgery, though our study was not powered to detect a difference in this restricted subgroup.

Patients who frequently had reported postoperative thigh pain prompted us to make changes in our surgical technique, performing TKA without use of a tourniquet. Tourniquet use has been associated with an increased risk of thigh pain after TKA by multiple authors.^{25,26} Postoperative thigh pain also is pressure dependent.²⁷ In addition, its use may be associated with a slightly increased risk of thromboembolic events and delayed functional recovery.^{28,29}

Because postoperative hemarthrosis is associated with more pain and reduced joint recovery function, we used topical TXA to reduce postoperative surgical site and joint hematoma. TXA (either oral, IV, or topical) during TKA is used to control postoperative bleeding primarily and decrease the need for transfusion without concomitant increase in thromboembolic events.^{30,31} Topical TXA may be more effective than IV, particularly in the immediate postoperative period.³² Although pain typically is not an endpoint in studies of TXA, a prospective study of 48 patients showed evidence that its use may be associated with decreased postoperative pain in the first 24 hours after surgery (though not after).³³ Finally, the use of intraarticular injection has evolved in our clinical practice, but literature is lacking with regard to its efficacy; more studies are needed to determine its effect relative to no injection. We have not seen any benefits to using cryotherapy in our practice; considering the costs for equipment and health care provider time, cryotherapy was not included in our new protocol.

Limitations

This is a nonrandomized retrospective single-institution study. Our study population is composed of mostly males with military experience and is not necessarily a representative sample of the general population eligible for joint arthroplasty. Our primary endpoint (reduction of opioid use postoperatively) also was a cornerstone of our intervention. To account for this, we set a very large effect size in our power analysis and evaluated multiple secondary endpoints to determine whether postoperative pain remained well controlled and complications/readmission minimized with our interventions. Because our intervention was multimodal, our study cannot make conclusions about the effect of a particular component of our treatment strategy. We did not measure or compare functional outcomes between both groups, which offers an opportunity for further research.

These limitations are balanced by several strengths. Our cohort was well controlled with respect to the dose and type of drug used. There is staff dedicated to postoperative telephone follow-up after discharge, and veterans are apt to seek care within the VA health care system, which improves case finding for complications and ED visits. No patients were lost to follow-up. Moreover, our drastic reduction in opioid use is promising enough to warrant reporting, while the broader orthopedic literature explores the relative impact of each variable.

CONCLUSIONS

The MOJO protocol has been effective for reducing postoperative opioid use after TKA without compromising effective pain management. The drastic reduction in the postoperative use of opioid pain medications and LOS have contributed to a cultural shift within our department, comprehensive team approach, multimodal pain management, and preoperative patient optimization. Further investigations are required to assess the impact of each intervention on observed outcomes. However, the framework and routines are applicable to other institutions and surgical specialties.

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