

# UCSF

## UC San Francisco Previously Published Works

### Title

Lifetime Medical Costs of Knee Osteoarthritis Management in the United States: Impact of Extending Indications for Total Knee Arthroplasty

### Permalink

<https://escholarship.org/uc/item/49w7j4fm>

### Journal

Arthritis Care & Research, 67(2)

### ISSN

2151-464X

### Authors

Losina, Elena  
Paltiel, A David  
Weinstein, Alexander M  
[et al.](#)

### Publication Date

2015-02-01

### DOI

10.1002/acr.22412

Peer reviewed



Published in final edited form as:

*Arthritis Care Res (Hoboken)*. 2015 February ; 67(2): 203–215. doi:10.1002/acr.22412.

## Lifetime medical costs of knee osteoarthritis management in the United States: Impact of extending indications for total knee arthroplasty

Elena Losina, PhD<sup>1,2,3</sup>, A. David Paltiel, PhD<sup>4</sup>, Alexander M. Weinstein, BA<sup>1</sup>, Edward Yelin, PhD<sup>5</sup>, David J. Hunter, MBBS, PhD<sup>6</sup>, Stephanie P. Chen, BS<sup>1</sup>, Kristina Klara, BS<sup>1</sup>, Lisa G. Suter, MD<sup>4,7</sup>, Daniel H. Solomon, MD, MPH<sup>2,8</sup>, Sara A. Burbine, BA<sup>1</sup>, Rochelle P. Walensky, MD, MPH<sup>2,9,10,11</sup>, and Jeffrey N. Katz, MD, MSc<sup>1,2,8</sup>

<sup>1</sup>Orthopaedic and Arthritis Center for Outcomes Research, Department of Orthopedic Surgery, Brigham and Women's Hospital, Boston, MA

<sup>2</sup>Harvard Medical School, Boston, MA

<sup>3</sup>Department of Biostatistics, Boston University School of Public Health, Boston, MA

<sup>4</sup>Yale School of Medicine, New Haven, CT

<sup>5</sup>University of California, San Francisco, San Francisco, CA

<sup>6</sup>Kolling Institute and Royal North Shore Hospital, University of Sydney, Sydney, Australia

<sup>7</sup>Veterans Affairs Medical Center, West Haven, CT

<sup>8</sup>Division of Rheumatology, Immunology and Allergy, Brigham and Women's Hospital, Boston, MA

<sup>9</sup>Division of Infectious Disease, Brigham and Women's Hospital, Boston, MA

<sup>10</sup>Division of Infectious Disease, Massachusetts General Hospital, Boston, MA

<sup>11</sup>Division of General Medicine and Medical Practice Evaluation Center, Massachusetts General Hospital, Boston, MA

### Abstract

**Objective**—The impact of increasing utilization of total knee arthroplasty (TKA) on lifetime costs in persons with knee OA is under-studied.

**Methods**—We used the Osteoarthritis Policy Model to estimate total lifetime costs and TKA utilization under a range of TKA eligibility criteria among US persons with symptomatic knee OA. Current TKA utilization was estimated from the Multicenter Osteoarthritis Study and calibrated to Health Care Utilization Project (HCUP) data. OA treatment efficacy and toxicity were drawn from published literature. Costs in 2013 USD were derived from Medicare

---

Correspondence: Elena Losina, PhD, Orthopaedic and Arthritis Center for Outcomes Research, Brigham and Women's Hospital, 75 Francis Street, BC 4-016, Boston, MA 02115, Phone: 617-732-5338, Fax: 617-525-7900, [elosina@partners.org](mailto:elosina@partners.org).

No other financial interests or disclosures.

reimbursement schedules and Red Book Online®. Time costs were derived from published literature and the US Bureau of Labor Statistics.

**Results**—Estimated average discounted (3%/year) lifetime costs for persons diagnosed with knee OA were \$140,300. Direct medical costs were \$129,600, with \$12,400 (10%) attributable to knee OA over 28 years. OA patients spent, on average, 13 (SD 10) years waiting for TKA after failing non-surgical regimens. Under current TKA eligibility criteria, 54% of knee OA patients underwent TKA over their lifetimes. Estimated OA-related discounted lifetime direct medical costs ranged from \$12,400 (54% TKA uptake) when TKA eligibility was limited to K-L 3 or 4 to \$16,000 (70% TKA uptake) when eligibility was expanded to include symptomatic OA with a lesser degree of structural damage.

**Conclusion**—Due to low efficacy of non-surgical regimens, knee OA treatment-attributable costs are low, representing a small portion of all costs for OA patients. Expanding TKA eligibility increases OA-related costs substantially for a population, underscoring the need for more effective non-operative therapies.

### Keywords

knee osteoarthritis; utilization of TKA; direct medical costs; lifetime costs

---

## INTRODUCTION

Symptomatic knee osteoarthritis (OA) is a chronic, painful condition affecting an estimated 9.3 million adults 45 years and older in the US (1). In addition to lowering quality of life, knee OA is a major economic burden (2–6). The American College of Rheumatology (ACR) (7), Osteoarthritis Research Society International (OARSI) (8,9), and American Academy of Orthopaedic Surgeons (AAOS) (10,11) have published clinical practice guidelines for the management of knee OA. However, these guidelines do not consider costs and do not provide guidance on indications for TKA (7–11). Based upon the number of TKAs performed annually and the estimated number of individuals in the US with OA, we determined that surgery is currently performed in persons in pain and Kellgren-Lawrence (K-L) grades 3 or 4 (12,13).

Since OA often occurs with multiple concomitant comorbidities, it is important to understand what portion of total direct medical costs is attributable to OA. The majority of prior economic evaluations regarding knee OA have focused on individual treatments (14–23). However, knee OA management includes a combination of treatments over a long time horizon. To date, there have been no published studies estimating the *lifetime* costs incurred by persons affected by symptomatic knee OA in the US.

Furthermore, TKA utilization has doubled over the last decade, an increase that was not entirely explained by population growth and the obesity epidemic (24). The growth in utilization is partially due to expanding eligibility criteria and greater willingness of symptomatic OA patients to undergo TKA, regardless of radiographic severity (24). While patients below 65 years of age comprised just a quarter of TKA recipients in 1997 (25), almost 40% of TKAs are now done prior to age 65 (24).

In the absence of treatment guidelines linking clinical choices to data on the economic burden of disease, we sought to estimate the lifetime resource use (including direct, OA-specific costs; direct, non-OA costs; and time costs due to productivity losses) associated with alternative TKA eligibility criteria.

## METHODS

### Analytic overview

We used the Osteoarthritis Policy (OAPol) Model (26,27) and published data on costs, utilization, efficacy, and toxicity of OA treatments to project lifetime direct medical costs (costs due to OA as well as all other conditions), knee OA-related costs, and time costs due to productivity losses in persons with diagnosed symptomatic knee OA. For comparison, we also estimated lifetime costs for knee OA-free individuals with similar demographic and clinical characteristics. While guidelines are largely prescriptive for non-surgical OA treatments, those for TKA are less detailed. We therefore conducted analysis across five sets of TKA eligibility criteria based on pain that is not relieved by non-surgical treatment and: 1) 100% K-L grade 4 (most conservative, limited to end-stage disease), 2) 50% K-L 3 and 100% K-L 4; 3) 100% K-L 3 or greater (defined as the “current TKA eligibility” criterion); 4) 50% K-L 2, 100% K-L 3 or greater; and 5) 100% K-L 2 or greater (least conservative). Lifetime cost estimates in real (i.e. inflation-adjusted) 2013 USD are reported both undiscounted and discounted at an annual rate of 3%, as recommended by the Panel on Cost-Effectiveness in Health and Medicine (28). In additional analyses, we added estimates of time costs due to absenteeism among employees diagnosed with knee OA (29,30).

Under these ranging TKA eligibility criteria, we estimated the proportion of individuals with knee OA that received each treatment, the proportion that received TKA before age 65, the mean duration of each treatment, and the mean age of undergoing TKA.

### OAPol Model structure

The OAPol Model is a validated, state-transition, computer simulation model of the natural history and management of knee OA (24,26,27,31,32). In the model, individuals transition among health states defined by structural severity of knee OA (K-L grades 0 to 4) and symptom status (33). Symptomatic knee OA is defined as radiographic knee OA (K-L grades 2 to 4) accompanied by pain on most days. In the beginning of each simulation, each hypothetical patient is assigned a K-L grade, age, sex, and BMI.

In addition to capturing the incidence and progression of knee OA, the OAPol Model tracks the development of other chronic conditions prevalent in persons with knee OA. Large cohorts are followed until death, which is determined in a probabilistic manner using US life tables adjusted with disease-specific relative risks of mortality (34–36). Additional descriptions of the OAPol Model structure have been published previously (26,32,37).

### Treatment strategies

We considered two treatment strategies: 1) care limited to occasional analgesics (for comparison purposes); and 2) guideline-concordant care. Occasional analgesic use was

modeled as acetaminophen, over-the-counter NSAIDs (ibuprofen or naproxen), or one of three opioids (codeine, hydrocodone, or oxycodone) prescribed by a general practitioner.

Consistent with recommendations published by ACR, OARSI, and AAOS, guideline-concordant care was modeled as four regimens offered sequentially in order of increasing intensity, with analgesics used as needed between treatments (Table 1) (7–11,38). Under guideline-concordant care, all individuals with diagnosed symptomatic knee OA received Non-Surgical Regimen 1, comprised of physical therapy, knee braces, acetaminophen, and NSAIDs. Upon failure of Non-Surgical Regimen 1, individuals were offered Non-Surgical Regimen 2, consisting of corticosteroid injections. Failure of Non-Surgical Regimen 2 led to offering of TKA (Surgical Regimen 3) for all those whose pain was not relieved with non-surgical treatment and who had structural changes due to OA evident on plain radiographs. Acceptance of primary TKA varied by age and sex and was derived from data from two large cohort studies (39,40). Individuals with primary TKA failure received revision TKA (Surgical Regimen 4). Those with pain, including patients who were in pain despite undergoing TKA, were assumed to use analgesics and accrue time costs due to productivity loss. Subjects whose pain was relieved by TKA had lower OA-related healthcare costs. These patients were also assumed to have lower pain-driven time costs.

## Input data

**Demographics and clinical characteristics**—The average age of diagnosis of symptomatic knee OA was assumed to be 54 (standard deviation (SD) 14) years (Table 1) (32). Data on sex, race, and obesity in persons with knee OA were derived from the 2012 National Health Interview Survey (41). Prevalence and incidence of comorbidities (cancer, coronary heart disease, chronic obstructive pulmonary disease, diabetes mellitus, and other musculoskeletal diseases) by sex and race/ethnicity were derived from the 2009–2010 National Health and Nutrition Examination Survey (NHANES) (42). Annual rates of knee OA progression, stratified by sex, obesity, and K-L grade, varied from 1.3% to 12.3% (26).

**Knee OA treatment efficacy**—The efficacy of non-surgical regimens in relieving pain was derived from published clinical trials, with Non-Surgical Regimen 1 efficacy weighted according to 2009 Medicare Current Beneficiary Survey (MCBS) data on NSAID utilization (43–45). Efficacy of pain relief in the first year for Non-Surgical Regimen 1 was 63% and for Non-Surgical Regimen 2 was 64% for individuals with early-stage knee OA. This efficacy was sustained in 76% of persons on Regimen 1 and in 81% on Regimen 2 in each subsequent year. For TKA, pain relief the first year after surgery was 86% (46). Mechanical failure of primary TKA leading to revision was stratified by age and ranged from 0.3% per year (for those 65 years and older) to 1.2% per year (for those younger than 65 years) (47) (see Technical Appendix for detailed data).

**Knee OA treatment toxicity**—Treatment toxicities were associated with specific costs, quality-of-life decrements, and risks of death. Estimated costs of toxicities were derived from 2011 HCUP data and converted to real (i.e. inflation-adjusted) 2013 USD (Table 1) (13,48).

## **Knee OA treatment costs**

**Occasional analgesic use:** We included an annual cost of occasional analgesic use for knee OA subjects either in pain but not on any regular regimen or when TKA fails to relieve pain (Table 1). This cost (\$102) was derived from the average annual cost of acetaminophen and opioids (49) weighted by the estimated utilization of each type of drug (21% - opioids; 79% - acetaminophen) (45). Additional details on cost derivation are presented in the Technical Appendix.

**Non-Surgical Regimen 1:** Overall, the cost of Non-Surgical Regimen 1 was estimated at \$684 in the first year and \$520 in each subsequent year. Costs of annual physician office visits, physical therapy, assistive devices, and laboratory tests were derived from 2012 Medicare reimbursement schedules (50,51). Costs of NSAIDs, acetaminophen, and gastro-protective agents were derived from Red Book Online® (49). Regimen component costs were weighted according to their utilization by OA patients, as derived from MCBS data. Further details are presented in the Technical Appendix.

**Non-Surgical Regimen 2:** Annual costs of Regimen 2 (\$494) included the cost of one physician office visit (\$99) (50,51) and an average of 2.5 corticosteroid knee injections per year (45). The estimated cost of each injection (\$157) was derived from the 2012 Medicare reimbursement schedule (averaged for hospital and non-hospital settings) for joint injections of lidocaine (10 mg) and methylprednisolone (40 mg) (50,51).

**Primary and revision TKA:** Total costs in 2013 USD for primary (\$20,293) and revision (\$26,388) TKA included the costs of surgery and rehabilitation derived from 2012 Medicare reimbursement schedules (50), the 2011 Medicare Hospital Inpatient Prospective Payment System (52), and literature (53). Annual follow-up costs (\$143) included a physician office visit and knee radiograph (\$44) (50,51). From a national survey of physicians' recommendations regarding post-TKA follow-up care, we estimated the annual likelihood of having a post-TKA follow-up visit to be 67% (54). The product of these two quantities yielded an annual average cost for post-TKA follow-up of \$95.

**Direct medical costs unrelated to knee OA—**We estimated average annual direct medical costs unrelated to knee OA, stratified by age and number of comorbidities, by weighting Centers for Medicare & Medicaid Services hierarchical condition categories (CMS-HCC) estimates according to aggregated data from NHANES 2009–10 (42). Costs were inflated to real (i.e., inflation-adjusted) 2013 USD using the Consumer Price Index (CPI) inflation calculator for Medical Care in the US (48). These costs varied from \$1,414 to \$8,202 per year for those with 0 to 1 comorbidities to \$14,291 to \$19,092 per year for those with greater than three comorbidities (Table 1).

**Time costs due to productivity losses—**We based time cost estimates (cost of lost productivity) on published data addressing work absenteeism among individuals with OA in the US work force (29). Our estimates are based entirely on the labor market and do not account for other types of productivity losses. The reported annual time costs for persons with OA who did not undergo TKA were estimated at \$1,474 per year (in 2007 USD)

among those in the workforce. Time costs for OA patients who had TKA while being in the workforce were estimated at \$7,104 (in 2007 USD) during the year of surgery. To properly account for differing levels of work absenteeism across age strata, we used employment data stratified by age provided by the US Bureau of Labor Statistics (30) (see the Technical Appendix for derivation details).

**Sensitivity analyses**—We conducted sensitivity analyses to address uncertainties in parameter estimates and to examine the effects of alternative assumptions. First, we varied the mean age of symptomatic knee OA diagnosis from 50 to 70 years. Second, we varied the distribution of background analgesics used from acetaminophen only to opioids only. Third, we varied K-L grade distribution at the time of diagnosis. Fourth, we evaluated the impact of using more effective and expensive analgesics in Regimens 1 and 2 on overall treatment utilization and cost. Lastly, we varied rates of knee OA progression based on obesity status.

## RESULTS

### Proportion of cohort receiving TKA and timing of treatments

Individuals spent several years on each non-surgical regimen before waiting an average of 13.3 (SD 10.4) years before receiving TKA, using analgesics as needed to manage pain prior to surgical treatment or death. Only about half the original cohort underwent primary TKA and a smaller percentage received revision TKA. Time spent on primary TKA was estimated at 16 years (Figure 1).

Utilization of primary TKA ranged from 19% of the total cohort when TKA eligibility was limited to K-L grade 4 to 70% with eligibility expanded to K-L grades 2 or above (Technical Appendix Figure 6-1). Under current TKA eligibility criteria, 38% of patients who underwent TKA received surgery prior to age 65. Overall, 21% of the knee OA population received primary TKA prior to age 65 (Technical Appendix Figure 6-1, third bar from the left, hatched black-and-white section). This estimate ranged from 6% with TKA eligibility limited to K-L 4 to 29% with eligibility set to K-L 2 or greater. Mean age at the time of primary TKA ranged from 65 (SD 12) years with TKA indications at K-L 2 or greater to 70 (SD 11) years with TKA indications limited to K-L 4 (Technical Appendix, Figure 6-1). The average time to revision or death for those TKA recipients under age 65 and for those ages 65 and over differed primarily due to competing mortality. When subjects were receiving TKA at age 65 or older, the time from primary to revision TKA or death was estimated at 13.0 years. For those receiving TKA before age 65, the average time from primary to revision TKA or death was estimated at 18.3 years.

### Lifetime direct medical costs in persons with symptomatic knee OA

When followed from an average age of knee OA diagnosis of 54 (SD 14) years, persons without knee OA who had similar demographic and clinical characteristics had average estimated per-person discounted lifetime medical costs (total, for all conditions) of \$117,500 (\$209,800 undiscounted) (Table 2). Under Treatment Strategy 1 (occasional analgesics only), those with symptomatic knee OA incurred average lifetime direct medical costs of

\$119,300 (\$212,700 undiscounted) per person. Under Treatment Strategy 2 (guideline-concordant care), this estimate was \$129,600 (\$228,600 undiscounted) per person.

### **Direct medical costs attributable to knee OA**

Under guideline-concordant care with current TKA eligibility criteria, direct lifetime medical costs attributable to knee OA were estimated at \$12,400 (\$19,600 undiscounted), 10% of the overall discounted lifetime direct medical costs for persons diagnosed with knee OA (Table 2). Knee OA-related direct medical costs ranged from \$6,600 (\$10,300 undiscounted) when only those with K-L grade 4 were eligible for TKA to \$16,000 (\$24,400 undiscounted) when those with K-L grades 2 or higher were eligible (Figure 2).

### **Distribution of knee OA-related costs by regimen**

Under current TKA eligibility criteria, the largest proportion of knee OA-related direct medical costs was attributable to primary TKA, followed by Non-Surgical Regimen 1, which consisted of NSAIDs, annual physician visits, physical therapy, assistive devices, acetaminophen, and gastro-protective agents (Figure 2). As eligibility criteria for TKA expanded, costs associated with surgical treatments comprised a greater proportion of OA-related costs, while those associated with analgesic use and non-surgical treatment comprised a smaller proportion.

### **Sensitivity analyses**

Under guideline-concordant care, knee OA-attributable per-person direct medical costs were greater when the mean age of symptomatic knee OA diagnosis was set to 50 years compared to 70 years. OA-related direct lifetime costs increased by about \$2,000 when the annual cost of intermittent analgesics was varied from \$71 (acetaminophen only) to \$218 (opioids only). Lifetime direct medical costs and OA-related costs were higher as the radiographic severity of knee OA at the time of diagnosis increased. Increasing the efficacy of Non-Surgical Regimens 1 and 2 had minimal impact on the utilization of TKA. When efficacy and costs were increased by 50%, TKA utilization decreased from 54% to 51%. Costs were somewhat more affected with total direct medical costs increasing by 2% and direct medical costs attributable to OA increasing by 21%. Finally, altering knee OA progression rates by obesity had very little effect on lifetime direct medical costs (Table 3).

Discounted time costs due to lost productivity decreased with expanding eligibility criteria for TKA, ranging from \$10,900 when TKA was restricted to symptomatic K-L grade 4 to \$10,400 for symptomatic OA regardless of structural severity (Figure 3). However, combining time costs with non-OA and OA-related direct medical costs increased total discounted lifetime medical costs with expanding TKA indications, with \$134,900 estimated for eligibility criteria of K-L grade 4 only to \$143,400 for eligibility at all K-L grades 2 and higher (Figure 3).

## **DISCUSSION**

Using a validated, published computer simulation model (26,27), we projected the lifetime direct medical and time costs for patients with symptomatic knee OA from disease onset

until death. The small proportion of lifetime direct medical costs (10%) attributable to knee OA treatment can be explained by the limited treatment options available for persons diagnosed with this debilitating condition. We found that persons with knee OA spend nearly 50% of their post-diagnosis life expectancy ‘in between’ treatments while waiting for TKA after exhausting available non-surgical options. We estimated that under current TKA eligibility criteria (pain unrelieved by non-surgical regimens and K-L grades 3 and 4), primary TKA is utilized by 54% of those with symptomatic knee OA and accounts for 61% of OA-related direct medical costs.

We found previously that the recent increase in TKA utilization is not explained entirely by population growth and the obesity epidemic (24), and is likely due, in part, to expanding indications and an increased willingness among patients to undergo surgery. Extending current eligibility criteria for TKA to any symptomatic disease, regardless of structural severity, led to an earlier average age of TKA, greater uptake of primary and revision TKA, and a 29% increase in lifetime direct medical costs attributable to knee OA. This finding – coupled with recent data suggesting greater failure rates and lower satisfaction rates among persons who underwent TKA in less advanced stages of OA – could prompt a stricter definition of TKA appropriateness. It also underscores the importance of finding alternative treatments for knee OA.

Our results extend the literature regarding the cost of OA management (29,55,56). Previous estimates of the annual direct medical costs of OA in the US vary widely (29,55,56), partly owing to heterogeneity in the methods used to distinguish OA-related care from all-cause health care utilization (2–6,55,57). Prior studies in this area have focused on the *annual* cost of medical care for persons with OA and report costs ranging from \$989 to \$10,313 in the US (56). Similar to other studies (58), our paper demonstrates that individuals with knee OA spend substantial financial resources on managing comorbidities.

Several findings from our study deserve special emphasis. First, guideline-concordant treatment represents a low but non-trivial cost burden for individuals with diagnosed symptomatic knee OA. Per-person lifetime costs of knee OA-related care (\$12,400) account for 10% of lifetime direct medical costs. Second, TKA is the only treatment for symptomatic knee OA with any substantial duration of sustained efficacy and represents the largest source of costs. While only half of those diagnosed with symptomatic knee OA ultimately underwent TKA in our model, costs related to primary and revision TKA constituted 69% of total direct medical costs attributable to OA. As eligibility criteria for TKA expanded, OA-related costs increased.

Third, our findings draw attention to the question of who is paying for these costly surgeries. We estimated that 38% of persons receiving primary TKA under current eligibility criteria underwent surgery prior to age 65, when most healthcare costs falls to private insurers. If eligibility criteria for TKA expand, OA-related costs will increase and a higher percentage of TKAs will occur prior to age 65. As a result, the number of revision surgeries after age 65 (when payment is likely to fall to Medicare) will increase (13). These findings highlight the dramatic shifts in cost burden among payers that could result from changes in TKA timing. Preventive measures focused, for example, on reducing obesity and knee injury could delay

knee OA onset, reducing the likelihood of eventual TKA. Although this has the potential to reduce the lifetime costs of the disease dramatically, attention must be paid to potential repercussions of shifting costs to Medicare.

Our study has certain limitations. First, we assumed that patients underwent treatments sequentially, failing prior treatments before undergoing new ones. While this sequence was based on multiple published treatment guidelines (7–11,38), some physicians may tailor treatments for individual patients, leading to variations in the time a patient spends on each regimen. Second, our lifetime cost estimates do not account for the high cost of nursing home care, suggesting a conservative estimate of costs. However, nursing home care is utilized only by a relatively small segment of the elderly population. Third, our projections of lifetime direct medical costs in patients affected by knee OA incorporate only currently available treatments and do not factor in the benefits of drugs and surgeries under development. Fourth, our estimates of time costs are likely to be conservative as other activities affected by pain or suffering (reduced work and home productivity, loss of leisure time) were not considered. Further examination of health-related utility – as a function of both K-L stages and age – might provide a more balanced perspective on the costs and benefits of changing TKA eligibility criteria. And finally, the data for background morbidity in persons with OA were derived from national data sources. These parameters do not change rapidly from year to year. To maintain data consistency, we occasionally use the previous release of the national data source. These restrictions do not affect any OA-related estimates we report.

This is the first study to report lifetime costs of knee OA and the first one to consider the effects of expanding eligibility criteria for TKA. TKA is the only long-term effective treatment for symptomatic knee OA, but it is also the most costly. Expanding eligibility criteria for TKA may contribute to patients receiving TKA earlier in life and thus increase knee OA-related costs for both private insurers and Medicare. Implementation of combined exercise and weight loss programs (shown to be efficacious in persons with knee OA (59)), efforts to prevent injuries, and the development of disease modifying osteoarthritis drugs (DMOADs) that reduce the progression of knee OA may delay or prevent the need for and use of TKA and later revision. On the other hand, DMOAD therapy may be very costly. Therefore it could potentially be more cost-effective to offer TKA earlier in OA progression. Some studies have shown that patients with worse pain and function prior to surgery have worse outcomes following TKA (60–62), suggesting that there may be some benefit to offering TKA to patients with less severe OA. However, studies have also indicated that patients with more severe OA tend to be more satisfied with the results of surgery (63–65). Many studies have also suggested that undergoing primary TKA at a younger age is associated with an increased risk of failure and revision surgery (66,67). Additionally, several other variables, such as mental health, patient expectations, obesity and other comorbidities, affect outcomes of TKA (68–70), making it difficult to distill the effects that expanding indications would have on outcomes. Taken together with these considerations, our findings may have major policy implications for the cost and treatment of knee OA and the budgetary impact on payers, including Medicare. Additional research is needed to fully explain the costs and/or benefits of earlier surgery.

## Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

## Acknowledgments

Supported by: NIH/NIAMS R01 AR064320, K24 AR057827

## References

1. Lawrence RC, Felson DT, Helmick CG, Arnold LM, Choi H, Deyo RA, et al. Estimates of the prevalence of arthritis and other rheumatic conditions in the united states. Part ii Arthritis Rheum. 2008; 58:26–35.
2. Gabriel SE, Crowson CS, Campion ME, O’Fallon WM. Direct medical costs unique to people with arthritis. J Rheumatol. 1997; 24:719–25. [PubMed: 9101508]
3. Gabriel SE, Crowson CS, O’Fallon WM. Costs of osteoarthritis: Estimates from a geographically defined population. J Rheumatol Suppl. 1995; 43:23–5. [PubMed: 7752127]
4. Lanes SF, Lanza LL, Radensky PW, Yood RA, Meenan RF, Walker AM, et al. Resource utilization and cost of care for rheumatoid arthritis and osteoarthritis in a managed care setting: The importance of drug and surgery costs. Arthritis Rheum. 1997; 40:1475–81. [PubMed: 9259428]
5. MacLean CH, Knight K, Paulus H, Brook RH, Shekelle PG. Costs attributable to osteoarthritis. J Rheumatol. 1998; 25:2213–8. [PubMed: 9818666]
6. Mapel DW, Shainline M, Paez K, Gunter M. Hospital, pharmacy, and outpatient costs for osteoarthritis and chronic back pain. J Rheumatol. 2004; 31:573–83. [PubMed: 14994407]
7. Hochberg MC, Altman RD, April KT, Benkhalti M, Guyatt G, McGowan J, et al. American college of rheumatology 2012 recommendations for the use of nonpharmacologic and pharmacologic therapies in osteoarthritis of the hand, hip, and knee. Arthritis Care Res. 2012; 64:465–74.
8. Zhang W, Moskowitz RW, Nuki G, Abramson S, Altman RD, Arden N, et al. Oarsi recommendations for the management of hip and knee osteoarthritis, part ii: Oarsi evidence-based, expert consensus guidelines. Osteoarthritis Cartilage. 2008; 16:137–62. [PubMed: 18279766]
9. Zhang W, Nuki G, Moskowitz RW, Abramson S, Altman RD, Arden NK, et al. Oarsi recommendations for the management of hip and knee osteoarthritis: Part iii: Changes in evidence following systematic cumulative update of research published through january 2009. Osteoarthritis Cartilage. 2010; 18:476–99. [PubMed: 20170770]
10. Richmond J, Hunter D, Irrgang J, Jones MH, Snyder-Mackler L, Van Durme D, et al. American academy of orthopaedic surgeons clinical practice guideline on the treatment of osteoarthritis of the knee. J Bone Joint Surg Am. 2010; 92:990–93. [PubMed: 20360527]
11. Richmond J, Hunter D, Irrgang J, Jones MH, Levy B, Marx R, et al. Treatment of osteoarthritis of the knee (nonarthroplasty). J Am Acad Orthop Surg. 2009; 17:591–600. [PubMed: 19726743]
12. Weinstein AM, Rome BN, Reichmann WM, Collins JE, Burbine SA, Thornhill TS, et al. Estimating the burden of total knee replacement in the united states. J Bone Joint Surg Am. 2013; 95:385–92. [PubMed: 23344005]
13. Nationwide inpatient sample. Agency for Healthcare Research and Quality; 2011. Healthcare cost and utilization project. <http://hcupnet.ahrq.gov/> [Accessed October 16, 2013 – November 14, 2013]
14. Coupe VM, Veenhof C, van Tulder MW, Dekker J, Bijlsma JW, Van den Ende CH. The cost effectiveness of behavioural graded activity in patients with osteoarthritis of hip and/or knee. Ann Rheum Dis. 2007; 66:215–21. [PubMed: 16880195]
15. Fioravanti A, Valenti M, Altobelli E, Di Orto F, Nappi G, Crisanti A, et al. Clinical efficacy and cost-effectiveness evidence of spa therapy in osteoarthritis. The results of “naiade” italian project. Panminerva Med. 2003; 45:211–7. [PubMed: 14618120]
16. Kamath CC, Kremers HM, Vanness DJ, O’Fallon WM, Cabanela RL, Gabriel SE. The cost-effectiveness of acetaminophen, nsais, and selective cox-2 inhibitors in the treatment of symptomatic knee osteoarthritis. Value Health. 2003; 6:144–57. [PubMed: 12641865]

17. Koskinen E, Eskelinen A, Paavolainen P, Pulkkinen P, Remes V. Comparison of survival and cost-effectiveness between unicompartmental arthroplasty and total knee arthroplasty in patients with primary osteoarthritis: A follow-up study of 50,493 knee replacements from the Finnish Arthroplasty Register. *Acta Orthop*. 2008; 79:499–507. [PubMed: 18766483]
18. Maetzel A, Krahn M, Naglie G. The cost effectiveness of rofecoxib and celecoxib in patients with osteoarthritis or rheumatoid arthritis. *Arthritis Rheum*. 2003; 49:283–92. [PubMed: 12794781]
19. Marshall JK, Pellissier JM, Attard CL, Kong SX, Marentette MA. Incremental cost-effectiveness analysis comparing rofecoxib with nonselective NSAIDs in osteoarthritis: Ontario Ministry of Health perspective. *Pharmacoeconomics*. 2001; 19:1039–49. [PubMed: 11735672]
20. Pope JE, Prashker M, Anderson J. The efficacy and cost effectiveness of n of 1 studies with diclofenac compared to standard treatment with nonsteroidal antiinflammatory drugs in osteoarthritis. *J Rheumatol*. 2004; 31:140–9. [PubMed: 14705233]
21. Reinhold T, Witt CM, Jena S, Brinkhaus B, Willich SN. Quality of life and cost-effectiveness of acupuncture treatment in patients with osteoarthritis pain. *Eur J Health Econ*. 2008; 9:209–19. [PubMed: 17638034]
22. Sevick MA, Miller GD, Loeser RF, Williamson JD, Messier SP. Cost-effectiveness of exercise and diet in overweight and obese adults with knee osteoarthritis. *Med Sci Sports Exerc*. 2009; 41:1167–74. [PubMed: 19461553]
23. Soohoo NF, Sharifi H, Kominski G, Lieberman JR. Cost-effectiveness analysis of unicompartmental knee arthroplasty as an alternative to total knee arthroplasty for unicompartmental osteoarthritis. *J Bone Joint Surg Am*. 2006; 88:1975–82. [PubMed: 16951114]
24. Losina E, Thornhill TS, Rome BN, Wright J, Katz JN. The dramatic increase in total knee replacement utilization rates in the United States cannot be fully explained by growth in population size and the obesity epidemic. *J Bone Joint Surg Am*. 2012; 94:201–7. [PubMed: 22298051]
25. Nationwide inpatient sample. Agency for Healthcare Research and Quality; 1997. Healthcare cost and utilization project. <http://hcupnet.ahrq.gov/> [Accessed December 12, 2013]
26. Holt HL, Katz JN, Reichmann WM, Gerlovin H, Wright EA, Hunter DJ, et al. Forecasting the burden of advanced knee osteoarthritis over a 10-year period in a cohort of 60–64 year-old US adults. *Osteoarthritis Cartilage*. 2011; 19:44–50. [PubMed: 20955807]
27. Losina E, Walensky RP, Reichmann WM, Holt HL, Gerlovin H, Solomon DH, et al. Impact of obesity and knee osteoarthritis on morbidity and mortality in older Americans. *Ann Intern Med*. 2011; 154:217–26. [PubMed: 21320937]
28. Weinstein MC, Siegel JE, Gold MR, Kamlet MS, Russell LB. Recommendations of the panel on cost-effectiveness in health and medicine. *JAMA*. 1996; 276:1253–8. [PubMed: 8849754]
29. Berger A, Hartrick C, Edelsberg J, Sadosky A, Oster G. Direct and indirect economic costs among private-sector employees with osteoarthritis. *J Occup Environ Med*. 2011; 53:1228–35. [PubMed: 22015547]
30. Employment status of the civilian population by sex, age, and disability status, not seasonally adjusted. Bureau of Labor Statistics, US Department of Labor; 2013. Economic news release. Table a-6. <http://www.bls.gov/news.release/empsit.t06.htm> [Accessed April 11, 2013]
31. Losina E, Walensky RP, Kessler CL, Emrani PS, Reichmann WM, Wright EA, et al. Cost-effectiveness of total knee arthroplasty in the United States: Patient risk and hospital volume. *Arch Intern Med*. 2009; 169:1113–21. [PubMed: 19546411]
32. Losina E, Weinstein AM, Reichmann WM, Burbine SA, Solomon DH, Daigle ME, et al. Lifetime risk and age at diagnosis of symptomatic knee osteoarthritis in the US. *Arthritis Care Res (Hoboken)*. 2013; 65:703–11. [PubMed: 23203864]
33. Kellgren JH, Lawrence JS. Radiological assessment of osteoarthrosis. *Ann Rheum Dis*. 1957; 16:494–502. [PubMed: 13498604]
34. National vital statistics system (accessed from Health Data Interactive). Centers for Disease Control and Prevention, and National Center for Health Statistics, US Department of Health and Human Services; 2010. <http://www.cdc.gov/nchs/hdi.htm> [Accessed March 13, 2014]
35. United States revised life tables. Centers for Disease Control, US Department of Health and Human Services; 2009. <http://www.cdc.gov/nchs/products/nvsr.htm#vol62> [Accessed April 29, 2014]

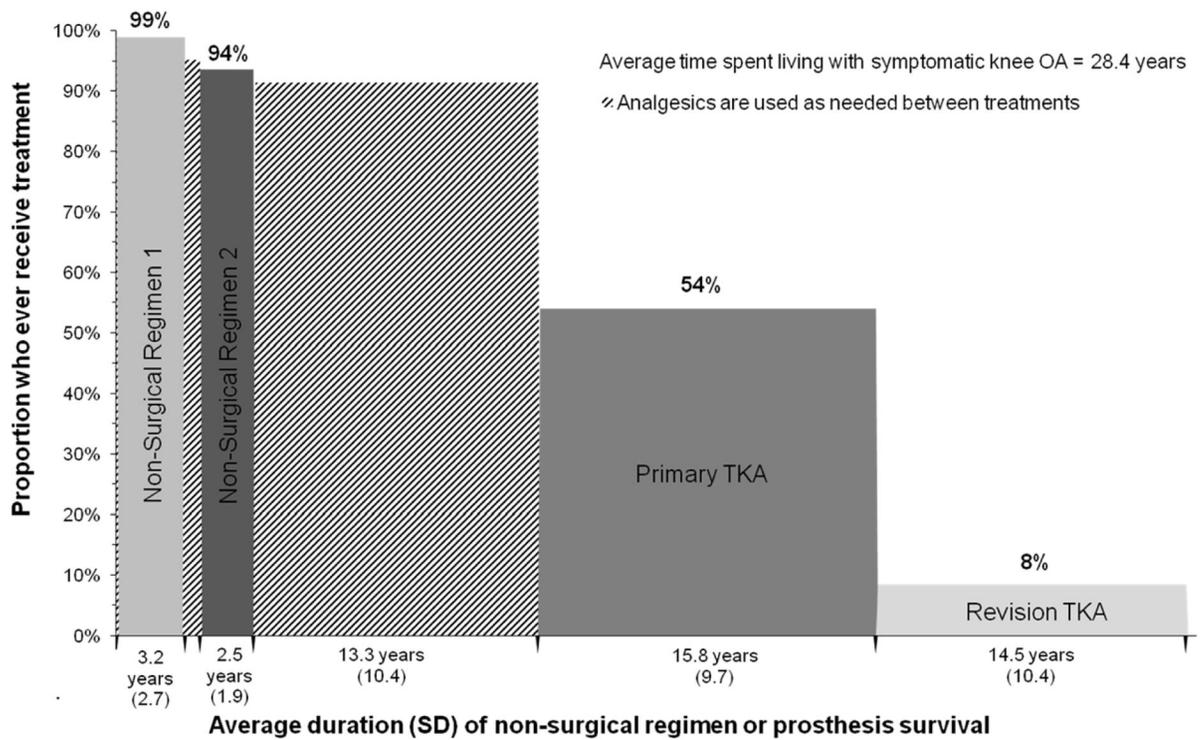
36. National health interview survey. Centers for Disease Control and Prevention, National Center for Health Statistics, US Department of Health and Human Services; 2010. <http://www.cdc.gov/nchs/nhis.htm> [Accessed March 13, 2014]
37. Suter LG, Paltiel AD, Rome BN, Solomon DH, Thornhill TS, Abrams SK, et al. Placing a price on medical device innovation: The example of total knee arthroplasty. *PLoS One*. 2013; 8:e62709. [PubMed: 23671626]
38. Recommendations for the medical management of osteoarthritis of the hip and knee: 2000 update. *Arthritis Rheum*. 2000; 43:1905–15. [PubMed: 11014340]
39. Wise BL, Felson DT, Clancy M, Niu J, Neogi T, Lane NE, et al. Consistency of knee pain and risk of knee replacement: The multicenter osteoarthritis study. *J Rheumatol*. 2011; 38:1390–5. [PubMed: 21498481]
40. Osteoarthritis initiative. Coordinating Center, University of California; San Francisco: 2010. <http://oai.epi-ucsf.org/datarelease/default.asp> [Accessed April 16, 2013]
41. National health interview survey. Centers for Disease Control and Prevention, National Center for Health Statistics, US Department of Health and Human Services; 2012. <http://www.cdc.gov/nchs/nhis.htm> [Accessed April 17, 2014]
42. National health and nutrition examination survey 2009–2010. Hyattsville, MD: Centers for Disease Control and Prevention, National Center for Health Statistics, US Department of Health and Human Services; 2010.
43. Raynauld JP, Buckland-Wright C, Ward R, Choquette D, Haraoui B, Martel-Pelletier J, et al. Safety and efficacy of long-term intraarticular steroid injections in osteoarthritis of the knee: A randomized, double-blind, placebo-controlled trial. *Arthritis Rheum*. 2003; 48:370–7. [PubMed: 12571845]
44. Scott DL, Berry H, Capell H, Coppock J, Daymond T, Doyle DV, et al. The long-term effects of non-steroidal anti-inflammatory drugs in osteoarthritis of the knee: A randomized placebo-controlled trial. *Rheumatology (Oxford)*. 2000; 39:1095–101. [PubMed: 11035129]
45. Medicare current beneficiary survey. Centers for Medicare & Medicaid Services; 2009. [http://www.cms.gov/LimitedDataSets/11\\_MCBS.asp](http://www.cms.gov/LimitedDataSets/11_MCBS.asp) [Accessed January 5, 2012]
46. Katz JN, Mahomed NN, Baron JA, Barrett JA, Fossel AH, Creel AH, et al. Association of hospital and surgeon procedure volume with patient-centered outcomes of total knee replacement in a population-based cohort of patients age 65 years and older. *Arthritis Rheum*. 2007; 56:568–74. [PubMed: 17265491]
47. Paxton EW, Namba RS, Maletis GB, Khatod M, Yue EJ, Davies M, et al. A prospective study of 80,000 total joint and 5000 anterior cruciate ligament reconstruction procedures in a community-based registry in the united states. *J Bone Joint Surg Am*. 2010; 92 (Suppl 2):117–32. [PubMed: 21123596]
48. [Accessed April 24, 2014] Consumer price index. 2014. <http://www.bls.gov/cpi/#data>
49. [Accessed March 24, 2014 – April 4, 2014] Red book online® Truven Health Analytics Inc. 2014. <http://www.micromedexsolutions.com/home/dispatch>
50. Medicare fee schedules. Centers for Medicare & Medicaid Services; 2012. <http://www.cms.gov/home/medicare.asp> [Accessed October 15, 2013]
51. Medicare hospital outpatient prospective payment system. Centers for Medicare & Medicaid Services; 2012. <http://www.cms.gov/HospitalOutpatientPPS/> [Accessed October 15, 2013]
52. Medicare hospital inpatient prospective payment system. Centers for Medicare & Medicaid Services; 2011. <http://www.cms.gov/Research-Statistics-Data-and-Systems/Statistics-Trends-and-Reports/MedicareFeeforSvcPartsAB/MEDPAR.html> [Accessed October 16, 2013]
53. Buntin, MB.; Deb, P.; Escarce, J.; Hoverman, C.; Paddock, S.; Sood, N. Comparison of medicare spending and outcomes for beneficiaries with lower extremity joint replacements. Arlington, VA: RAND Health, Medicare Payment Advisory Commission; 2005. WR-271-MEDPAC
54. Teeny SM, York SC, Mesko JW, Rea RE. Long-term follow-up care recommendations after total hip and knee arthroplasty: Results of the american association of hip and knee surgeons' member survey. *J Arthroplasty*. 2003; 18:954–62. [PubMed: 14658097]

55. White AG, Birnbaum HG, Janagap C, Buteau S, Schein J. Direct and indirect costs of pain therapy for osteoarthritis in an insured population in the united states. *J Occup Environ Med.* 2008; 50:998–1005. [PubMed: 18784547]
56. Xie F, Thumboo J, Li SC. True difference or something else? Problems in cost of osteoarthritis studies. *Semin Arthritis Rheum.* 2007; 37:127–32. [PubMed: 17350673]
57. Liang MH, Larson M, Thompson M, Eaton H, McNamara E, Katz R, et al. Costs and outcomes in rheumatoid arthritis and osteoarthritis. *Arthritis Rheum.* 1984; 27:522–9. [PubMed: 6721884]
58. Lee DW, Meyer JW, Clouse J. Implications of controlling for comorbid conditions in cost-of-illness estimates: A case study of osteoarthritis from a managed care system perspective. *Value Health.* 2001; 4:329–34. [PubMed: 11705300]
59. Messier SP, Loeser RF, Miller GD, Morgan TM, Rejeski WJ, Sevick MA, et al. Exercise and dietary weight loss in overweight and obese older adults with knee osteoarthritis: The arthritis, diet, and activity promotion trial. *Arthritis Rheum.* 2004; 50:1501–10. [PubMed: 15146420]
60. Fortin PR, Penrod JR, Clarke AE, St-Pierre Y, Joseph L, Belisle P, et al. Timing of total joint replacement affects clinical outcomes among patients with osteoarthritis of the hip or knee. *Arthritis Rheum.* 2002; 46:3327–30. [PubMed: 12483739]
61. Fortin PR, Clarke AE, Joseph L, Liang MH, Tanzer M, Ferland D, et al. Outcomes of total hip and knee replacement: Preoperative functional status predicts outcomes at six months after surgery. *Arthritis Rheum.* 1999; 42:1722–8. [PubMed: 10446873]
62. Desmeules F, Dionne CE, Belzile EL, Bourbonnais R, Champagne F, Fremont P. Determinants of pain, functional limitations and health-related quality of life six months after total knee arthroplasty: Results from a prospective cohort study. *BMC Sports Sci Med Rehabil.* 2013; 5:2. [PubMed: 23566925]
63. Keurentjes JC, Fiocco M, So-Osman C, Onstenk R, Koopman-Van Gemert AW, Poll RG, et al. Patients with severe radiographic osteoarthritis have a better prognosis in physical functioning after hip and knee replacement: A cohort-study. *PLoS One.* 2013; 8:e59500. [PubMed: 23573200]
64. Schnurr C, Jarrous M, Gudden I, Eysel P, König DP. Pre-operative arthritis severity as a predictor for total knee arthroplasty patients' satisfaction. *Int Orthop.* 2013; 37:1257–61. [PubMed: 23525526]
65. Dowsey MM, Nikpour M, Dieppe P, Choong PF. Associations between pre-operative radiographic changes and outcomes after total knee joint replacement for osteoarthritis. *Osteoarthritis Cartilage.* 2012; 20:1095–102. [PubMed: 22800770]
66. Dy CJ, Marx RG, Bozic KJ, Pan TJ, Padgett DE, Lyman S. Risk factors for revision within 10 years of total knee arthroplasty. *Clin Orthop Relat Res.* 2013
67. Julin J, Jansen E, Puolakka T, Kontinen YT, Moilanen T. Younger age increases the risk of early prosthesis failure following primary total knee replacement for osteoarthritis. A follow-up study of 32,019 total knee replacements in the finnish arthroplasty register. *Acta Orthop.* 2010; 81:413–9. [PubMed: 20809740]
68. Mahomed NN, Liang MH, Cook EF, Daltroy LH, Fortin PR, Fossel AH, et al. The importance of patient expectations in predicting functional outcomes after total joint arthroplasty. *J Rheumatol.* 2002; 29:1273–9. [PubMed: 12064846]
69. Gandhi R, Davey JR, Mahomed NN. Predicting patient dissatisfaction following joint replacement surgery. *J Rheumatol.* 2008; 35:2415–8. [PubMed: 19004032]
70. Dowsey MM, Liew D, Stoney JD, Choong PF. The impact of pre-operative obesity on weight change and outcome in total knee replacement: A prospective study of 529 consecutive patients. *J Bone Joint Surg Br.* 2010; 92:513–20. [PubMed: 20357327]
71. Pope GC, Kautter J, Ellis RP, Ash AS, Ayanian JZ, Lezzoni LI, et al. Risk adjustment of medicare capitation payments using the cms-hcc model. *Health Care Financ Rev.* 2004; 25:119–41. [PubMed: 15493448]
72. The use of medicines in the united states: Review of 2010. Parsippany, NJ: IMS Institute for Healthcare Informatics; 2011.
73. Medicaid drug price comparison: Average sales price to average wholesale price. Office of Inspector General, US Department of Health and Human Services; 2005. <http://oig.hhs.gov/oei/reports/oei-03-05-00200.pdf>

74. Grindrod KA, Marra CA, Colley L, Cibere J, Tsuyuki RT, Esdaile JM, et al. After patients are diagnosed with knee osteoarthritis, what do they do? *Arthritis Care Res.* 2010; 62:510–15.
75. Van der Esch M, Heijmans M, Dekker J. Factors contributing to possession and use of walking aids among persons with rheumatoid arthritis and osteoarthritis. *Arthritis Care Res.* 2003; 49:838–42.
76. Bensen WG, Fiechtner JJ, McMillen JJ, Zhao WW, Yu SS, Woods EM, et al. Treatment of osteoarthritis with celecoxib, a cyclooxygenase-2 inhibitor: A randomized controlled trial. *Mayo Clinic Proceedings.* 1999; 74:1095–105. [PubMed: 10560596]
77. Solomon SD, McMurray JJV, Pfeffer MA, Wittes J, Fowler R, Finn P, et al. Cardiovascular risk associated with celecoxib in a clinical trial for colorectal adenoma prevention. *N Engl J Med.* 2005; 352:1071–80. [PubMed: 15713944]
78. Goldstein JL, Silverstein FE, Agrawal NM, Hubbard RC, Kaiser J, Maurath CJ, et al. Reduced risk of upper gastrointestinal ulcer complications with celecoxib, a novel cox-2 inhibitor. *Am J Gastroenterol.* 2000; 95:1681–90. [PubMed: 10925968]
79. Silverstein FE, Graham DY, Senior JR, Davies HW, Struthers BJ, Bittman RM, et al. Misoprostol reduces serious gastrointestinal complications in patients with rheumatoid arthritis receiving nonsteroidal anti-inflammatory drugs. A randomized, double-blind, placebo-controlled trial. *Ann Intern Med.* 1995; 123:241–9. [PubMed: 7611589]
80. Yen Z-S, Lai M-S, Wang C-T, Chen L-S, Chen S-C, Chen W-J, et al. Cost-effectiveness of treatment strategies for osteoarthritis of the knee in taiwan. *J Rheumatol.* 2004; 31:1797–803. [PubMed: 15338503]
81. Ayril X. Injections in the treatment of osteoarthritis. *Best Pract Res Clin Rheumatol.* 2001; 15:609–26. [PubMed: 11567543]
82. Katz JN, Barrett J, Mahomed NN, Baron JA, Wright RJ, Losina E. Association between hospital and surgeon procedure volume and the outcomes of total knee replacement. *J Bone Joint Surg Am.* 2004; 86-A:1909–16. [PubMed: 15342752]

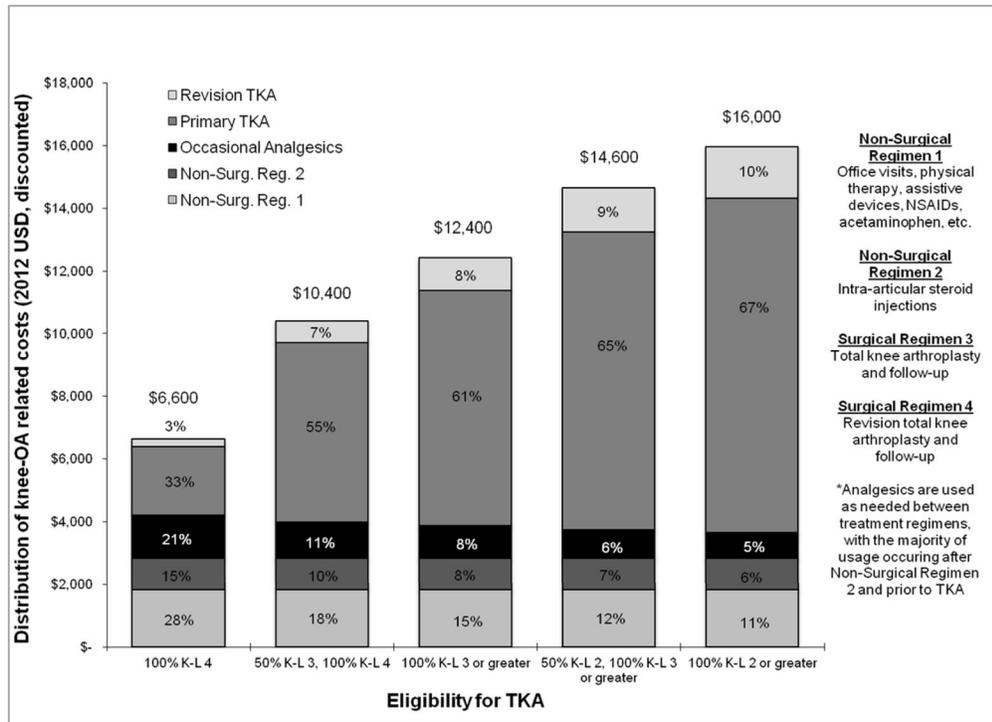
**SIGNIFICANCE AND INNOVATIONS**

- This is the first study to report lifetime costs of knee OA. Our estimates suggest that the cost of symptomatic knee OA in the US is relatively low (\$12,400 per-person), accounting for 10% of total direct medical costs for persons diagnosed with symptomatic knee osteoarthritis.
- Expanding TKA eligibility criteria may contribute to patients receiving TKA earlier in life and thus could increase knee OA-related costs by 29%.



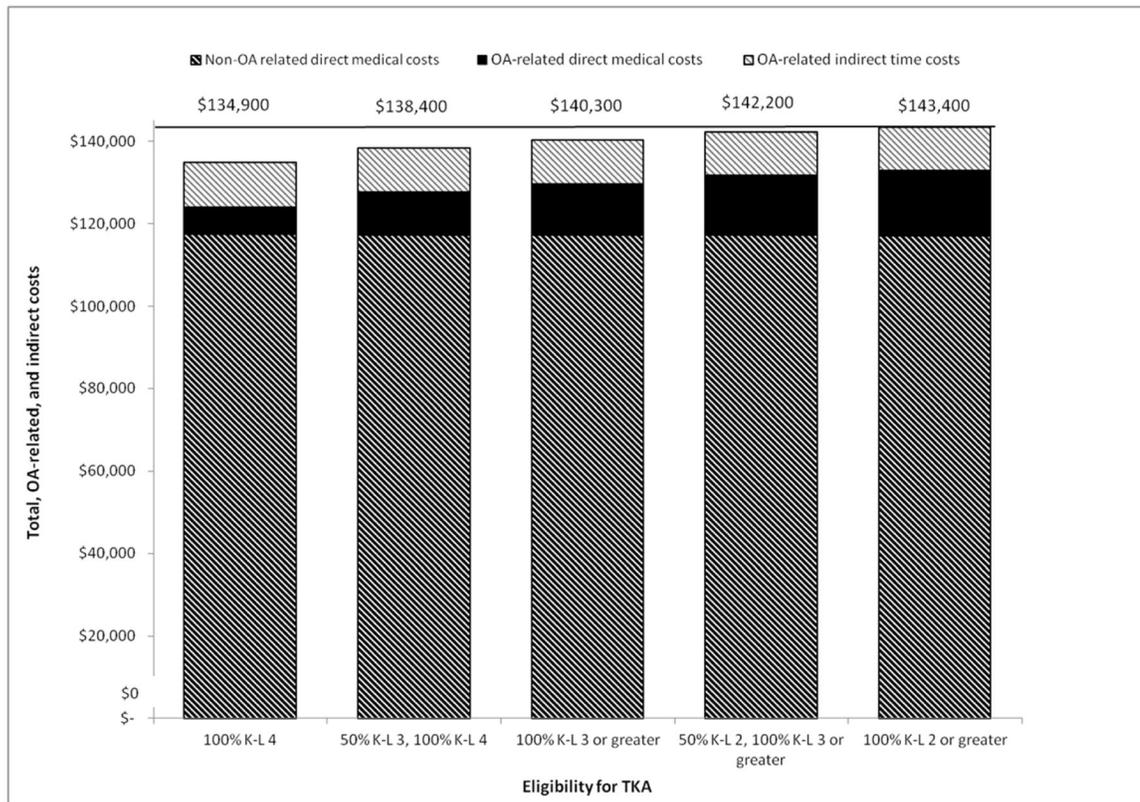
**Figure 1. Utilization and duration of Knee OA treatments under guideline-concordant care, assuming TKA eligibility criteria is defined by persistent pain, unrelieved by non-surgical regimens and evidence of advanced knee OA (K-L grades 3 or 4)**

Treatments under Treatment Strategy 2 (guideline-concordant care) are organized chronologically in order of increasing intensity along the x-axis. Intervening periods of occasional analgesic use are depicted with the hatched portions, with the longest duration of analgesic use occurring after Non-Surgical Regimen 2 and prior to primary TKA. The vertical height of each segment represents the proportion of those with symptomatic knee OA who ever received the treatment. The horizontal width of each segment represents the average duration (in years) of non-surgical regimens or prosthesis survival for those who ever received the treatment. The average time spent within each time period is written under each segment with the standard deviation, reported in years, in parenthesis. The average time spent living with symptomatic knee OA was 28.4 years. Primary and revision TKA were offered to patients with K-L grades 3 or 4.



**Figure 2. Varying eligibility criteria for TKA: Effects on distribution of OA-related, lifetime direct medical costs in patients with knee OA under guideline-concordant care**

The series of stacked columns compares the percent and dollar distribution of lifetime direct medical costs (discounted) attributable to the four treatments for knee OA as well as occasional analgesic use under Treatment Strategy 2 (guideline-concordant care). Columns are presented across expanding eligibility criteria for TKA (left to right). The total direct medical costs attributable to knee OA under guideline-concordant care are listed above the columns. The cost of occasional analgesic use is shown in black and positioned between regimens 2 and 3 because use of analgesics is greatest during this treatment phase. Non-surgical Regimen 1 consists of office visits, physical therapy, assistive devices, and typical pain relief medications such as NSAIDs, acetaminophen, etc. Non-surgical Regimen 2 consists of intra-articular steroid injections. Surgical Regimen 3 is total knee arthroplasty and follow-up appointments, and Regimen 4 is revision total knee arthroplasty and follow-up appointments. Analgesics are used as needed between treatment regimens, with the majority of usage occurring after Non-surgical Regimen 2 and prior to TKA.



**Figure 3. Varying eligibility criteria for TKA: Effects on lifetime direct and indirect medical costs for patients with knee OA**

The series of stacked columns compares the distribution of lifetime total medical costs in knee OA patients receiving guideline-concordant care across expanding eligibility criteria for TKA (left to right). The total costs for each TKA eligibility criterion are listed above the columns. OA-related direct medical costs are shown in between total indirect medical costs attributable to knee OA (above) and non-OA related direct medical costs (below). For comparison, the horizontal line represents the total cost under TKA eligibility K-L grade 2 or greater.

**Table 1**

OAPol model input parameters, cohort characteristics, and treatment parameters

Parameter	Overall Estimates
Mean Age (St. Dev)	53.5 (14.4)
% Female	53.2
Mean Body Mass Index (SD)	30.4 (6.8)
Prevalence of Knee Osteoarthritis	100% of population has Kellgren-Lawrence (K-L) grade 2 100% of population has knee pain associated with osteoarthritis
Progression estimates (annual probability, %)	Male Female
Progression from K-L 2 to K-L 3 (non-obese/obese)	5.6/12.3 4.0/8.9
Progression from K-L 3 to K-L 4 (non-obese/obese)	1.3/2.9 1.9/4.3
<b>Direct Medical Costs/Year (42,71) (in real (i.e. inflation-adjusted) 2013 USD)</b>	
Age	0-1 comorbidities 2-3 comorbidities >3 comorbidities
25-34	\$1,414.26 \$7,464.92 \$14,290.62
35-44	\$1,975.45 \$8,026.10 \$14,290.62
45-49	\$2,673.83 \$8,170.64 \$14,290.62
50-54	\$2,674.65 \$8,171.46 \$14,291.43
55-59	\$3,502.56 \$8,798.13 \$14,702.03
60-64	\$4,269.37 \$9,564.95 \$15,468.85
65-69	\$4,582.21 \$9,899.78 \$15,472.25
70-74	\$5,337.78 \$10,655.35 \$16,227.82
75-79	\$6,240.09 \$11,557.66 \$17,130.13
80+	\$8,201.93 \$13,519.49 \$19,091.96
<b>Treatment description Treatment parameters</b>	
Occasional analgesics <sup>a</sup> (acetaminophen, over-the-counter NSAIDs, opioids)	Regimen cost (annual, \$) <sup>b</sup> 102
Non-Surgical Regimen 1: office visits, devices, physical therapy, NSAIDs, acetaminophen	Regimen cost (annual, \$) <sup>b</sup> Subsequent years Initial year
	2012 Medicare reimbursement schedules; 2012 Medicare reimbursement schedules;

Treatment description	Treatment parameters			Source
	684	520		Red Book Online®; MCBS 2009; Van Der Esch et al, 2003; Grindrod et al, 2010; IMS Report; Office of Inspector General (45,49–51,72–75)
<b>Toxicities</b>	<b>Cost of toxicity (\$)<sup>b</sup></b>	<b>Risk of toxicity while on regimen (annual, %)</b>	<b>Risk of death due to toxicity (annual, %)(13)</b>	<b>Risk of toxicity</b>
General <sup>c</sup>	51	2.76 (initial year) 2.11 (subsequent years)	0.00	Bensen et al, 1999; Scott et al, 2000 (44,76)
Cardiovascular <sup>d,e</sup>	19,089	0.12	5.29	Solomon et al, 2005(77)
Gastrointestinal <sup>d,f</sup>	9,984	0.23	2.68	Goldstein et al, 2000; Silverstein et al, 1995(78,79)
<b>Non-Surgical Regimen 2: office visits, injections</b>	<b>Initial year</b>	<b>Subsequent years</b>	<b>Source</b>	
	494	494	2012 Medicare reimbursement schedules, MCBS 2009(45,50,51)	
<b>Toxicities</b>	<b>Cost of toxicity (\$)<sup>b</sup></b>	<b>Risk of toxicity while on regimen (annual, %)</b>	<b>Risk of death due to toxicity (annual, %)(13)</b>	<b>Risk of toxicity</b>
Skin flare	0	24.00	0.00	Yen et al, 2004(80) Ayrat et al, 2001(81)
Sepsis <sup>d</sup>	14,065	0.0013	8.35	HCUP 2011(13) Ayrat et al, 2001(81)
<b>Treatment 3: primary TKA and follow-up</b>	<b>Year of surgery</b>	<b>Years of follow-up</b>	<b>Year of surgery</b>	<b>Years of follow-up</b>
	20,293	95	2012 Medicare reimbursement schedules; Bunton et al, 2005; HCUP 2011(13,50,52,53)	Medicare reimbursement schedules; Teeny et al, 2003(50,51,54)
<b>Toxicities</b>	<b>Cost of toxicity (\$)<sup>b</sup></b>	<b>Risk of toxicity during year of surgery (%)(13)<sup>g</sup></b>	<b>Cost</b>	<b>Risk of toxicity</b>
Myocardial infarction	19,089	0.80	HCUP 2011(13)	Katz et al, 2004(82)

Treatment description		Treatment parameters			Source
	Pulmonary embolus	11,440	0.79	HCUP 2011(13)	Katz et al. 2004(82)
	Pneumonia	10,684	1.36	HCUP 2011(13)	Katz et al. 2004(82)
	Prosthetic joint infection <sup>d</sup>	23,488	0.70	HCUP 2011(13)	Paxton et al. 2010(47)
<b>Death from toxicity</b>	<b>Cost of death due to toxicity (\$)<sup>b</sup></b>	<b>Cost of death due to toxicity (\$)(13)<sup>g</sup></b>	<b>Risk of death due to toxicity during year of surgery (%)<sup>(13)</sup><sup>g</sup></b>	<b>Cost</b>	<b>Risk of death</b>
	15,149	26,388	0.63	HCUP 2011(13)	Katz et al. 2004(82)
<b>Treatment 4: revision TKA and follow-up</b>	<b>Treatment cost (annual, \$)<sup>b</sup></b>	<b>Year of surgery</b>	<b>Years of follow-up</b>	<b>Year of surgery</b>	<b>Years of follow-up</b>
		26,388	95	2012 Medicare reimbursement schedules; Teeny et al, 2003(50,51,54)	Medicare reimbursement schedules; Teeny et al, 2003(50,51,54)
<b>Treatment 4: revision TKA and follow-up</b>	<b>Toxicities</b>	Same as primary TKA			
<b>Death from toxicity</b>	<b>Cost of death due to toxicity (\$)<sup>b</sup></b>	<b>Cost of death due to toxicity during year of surgery (%)<sup>(13)</sup><sup>g</sup></b>	<b>Cost</b>	<b>Risk of death</b>	<b>Risk of death</b>
	15,149	0.96	HCUP 2011(13)	Katz et al. 2004 (82)	

<sup>a</sup>We did not account for toxicities related to occasional use of analgesics for pain management.

<sup>b</sup>All costs are reported in real (i.e. inflation-adjusted) 2013 USD.

<sup>c</sup>General toxicity includes the most common adverse events associated with the use of celecoxib for OA. These include headache, upper respiratory tract infection, dyspepsia, diarrhea, and abdominal pain.

<sup>d</sup>Occurrence of this toxicity in a given year causes the individual to be removed from the current regimen and evaluated for the subsequent regimen.

<sup>e</sup>Cardiovascular adverse events include myocardial infarction and thromboembolic events.

<sup>f</sup>Gastrointestinal adverse events include symptomatic upper gastrointestinal ulcer complications.

<sup>g</sup>For surgical regimens, we assume that any toxicity associated with the treatment will occur in the initial year. Thus, likelihood of toxicity or death due to toxicity in subsequent years is 0.

**Table 2**  
Average per-person lifetime direct medical costs and costs attributable to symptomatic knee OA

Treatment Strategy(or Cohort Description)	Total direct medical costs* (SD)		Direct medical costs* attributable to symptomatic knee OA (SD)		Proportion (%) of lifetime costs attributable to symptomatic knee OA	
	Discounted	Undiscounted	Discounted	Undiscounted	Discounted	Undiscounted
Matched control group**	\$117,500 (\$63,300)	\$209,800 (\$136,300)	-	-	-	-
1. Occasional analgesic use only	\$119,300 (\$63,800)	\$212,700 (\$137,600)	\$1,800 (\$700)	\$2,900 (\$1,600)	1.5%	1.4%
2. Guideline-concordant care	\$129,600 (\$66,800)	\$228,600 (\$144,800)	\$12,400 (\$9,400)	\$19,600 (\$16,200)	9.6%	8.6%

\* Costs are reported in real (i.e., inflation-adjusted) 2013 USD.

\*\* The matched control group had zero prevalence or incidence of symptomatic knee OA. Treatment Strategies 1 and 2 refer to cohorts with symptomatic knee OA.

**Table 3**

Sensitivity analyses, evaluating the effect on outcomes of varying natural history and intervention input parameters

Treatment strategy	Mean age of onset	Discounted lifetime direct medical costs	Undiscounted lifetime direct medical costs	Discounted OA-related costs	Undiscounted OA-related costs
<i>Matched control group</i>					
	50	\$117,000	\$217,900	-	-
	60	\$115,900	\$191,100	-	-
	70	\$106,300	\$156,400	-	-
<i>1. Occasional analgesic use only</i>					
	50	\$119,000	\$221,000	\$1,900	\$3,200
	60	\$117,500	\$193,600	\$1,600	\$2,400
	70	\$107,600	\$158,200	\$1,300	\$1,700
<i>2. Guideline-concordant care</i>					
	50	\$129,900	\$238,400	\$13,100	\$21,300
	60	\$126,400	\$206,700	\$10,800	\$16,100
	70	\$114,200	\$167,200	\$8,100	\$11,200
Treatment strategy	Percent of cohort using: opioids/acetaminophen (average cost of analgesic use)	Discounted lifetime direct medical costs	Undiscounted lifetime direct medical costs	Discounted OA-related costs	Undiscounted OA-related costs
<i>1. Occasional analgesic use only</i>					
	100% use acetaminophen (\$71)	\$118,800	\$211,900	\$1,300	\$2,000
	50% use opioids, 50% use acetaminophen (\$144)	\$120,100	\$214,000	\$2,600	\$4,100
<i>2. Guideline-concordant care</i>					
	100% use opioids (\$218)	\$121,400	\$216,000	\$3,900	\$6,200
	100% use acetaminophen (\$71)	\$129,200	\$228,000	\$12,000	\$19,000
	50% use opioids, 50% use acetaminophen (\$144)	\$130,200	\$229,500	\$12,900	\$20,400
	100% use opioids (\$218)	\$131,000	\$230,800	\$13,800	\$21,800
Treatment strategy	OA severity at time of diagnosis (K-L grade)	Discounted lifetime direct medical costs	Undiscounted lifetime direct medical costs	Discounted OA-related costs	Undiscounted OA-related costs
<i>2. Guideline-concordant care</i>					
	85% K-L 2, 10% K-L 3, 5% K-L 4	\$130,400	\$229,600	\$13,200	\$20,500
	60% K-L 2, 30% K-L 3, 10% K-L 4	\$131,700	\$231,200	\$14,500	\$22,100
Treatment Strategy Used	OA Progression Rate	Discounted Lifetime Direct Medical Costs*	Undiscounted Lifetime Direct Medical Costs*	Discounted OA-related costs*	Undiscounted OA-related Costs*
<i>Matched control group</i>	Current	\$117,500	\$209,800	-	-

Treatment Strategy *	OA Progression Rate Used	Discounted Lifetime Direct Medical Costs *	Undiscounted Lifetime Direct Medical Costs *	Discounted OA-related costs *	Undiscounted OA-related Costs *
	Non-obese	\$117,500	\$209,800	-	-
	Weighted	\$117,600	\$210,000	-	-
	Obese	\$117,500	\$209,800	-	-
<hr/>					
	Current	\$119,400	\$212,900	\$1,800	\$2,900
<i>1. Occasional analgesic use only</i>					
	Non-obese	\$119,400	\$212,900	\$1,800	\$2,900
	Weighted	\$119,300	\$212,700	\$1,800	\$2,900
	Obese	\$119,300	\$212,700	\$1,800	\$2,900
<hr/>					
	Current	\$258,100	\$455,500	\$12,400	\$19,600
<i>2. Guideline-concordant care</i>					
	Non-obese	\$255,200	\$451,100	\$11,000	\$17,400
	Weighted	\$258,500	\$456,200	\$12,700	\$20,100
	Obese	\$261,000	\$459,600	\$13,900	\$21,800

\* The matched control group had zero prevalence or incidence of symptomatic knee OA. Treatment Strategies 1 and 2 refer to cohorts with symptomatic knee OA.

\*\* Costs are reported in real (i.e., inflation-adjusted) 2013 USD.