Association Between Preoperative Radiographic Severity of Osteoarthritis and Patient-Reported Outcomes of Total Knee Replacement

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Background: The goal of this study was to investigate the association between preoperative radiographic severity of knee osteoarthritis (OA) and patient-reported outcomes following total knee replacement.

Methods: We used data from a prospective cohort study of individuals who underwent total knee replacement at a high-volume medical center. Patient-reported outcomes included the Western Ontario and McMaster Universities Osteoarthritis Index (WOMAC) pain score and the Knee injury and Osteoarthritis Outcome Score (KOOS) activities of daily living (ADL) subscore, assessed preoperatively and 2 years postoperatively. We measured preoperative radiographic OA severity using the Osteoarthritis Research Society International (OARSI) Atlas score, dichotomized at the median. We assessed the association between radiographic OA severity and postoperative patient-reported outcomes in bivariate analyses and in multivariable linear regression, with adjustment for age, sex, body mass index, and comorbidity score.

Results: The analytic cohort included 240 patients with a mean age at surgery of 66.6 years (standard deviation, 8 years); 61% were female. The median total OARSI radiographic severity score was 10 (range, 3 to 17). The cohort improved substantially at 2 years following total knee replacement, with WOMAC pain and KOOS ADL score improvements on the order of 30 points. We did not observe significant or clinically important differences in pain relief or functional improvement between patients with milder and more severe radiographic OA. Sensitivity analyses using other radiographic assessment measures yielded similar findings.

Conclusions: Total knee replacement offers substantial symptomatic relief and functional improvement regardless of preoperative radiographic OA severity.

Level of Evidence: Prognostic Level IV. See Instructions for Authors for a complete description of levels of evidence.

Millions of primary total knee replacement procedures have been performed worldwide, and >80% have been performed for a primary diagnosis of osteoarthritis (OA). As of 2010, approximately 4.7 million adults were living with a knee replacement in the United States. Although total knee replacement has a high success rate, is cost-effective, and improves quality of life for millions of individuals, up to 20% of patients experience suboptimal outcomes. The number of total knee replacement procedures performed annually continues to rise, highlighting the importance of continued attention to optimizing outcomes for patients following total knee replacement.

Knee radiographs are the mainstay of OA diagnosis. Understanding implications of preoperative radiographic findings for total knee replacement surgical outcomes may help guide the preoperative planning process and help set expectations prior to total knee replacement surgery. Previous investigations of the association between preoperative radiographic OA severity and

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postoperative patient-reported outcomes following total knee replacement have not reached definitive conclusions\textsuperscript{10-16}. Some studies have demonstrated better outcomes among individuals with radiographically severe OA compared with milder OA\textsuperscript{10-14}, while other studies have shown no difference in outcomes between the 2 groups\textsuperscript{15,16}. Most of these investigations utilized the Kellgren-Lawrence (KL) system, or a variant of this method, to grade radiographic OA severity. However, the KL system does not provide granular assessments of radiographic findings, such as osteophytes or joint-space narrowing, does not differentiate between medial and lateral tibiofemoral disease, and gives substantial weight to joint-space narrowing in the final score\textsuperscript{17}.

The Osteoarthritis Research Society International (OARSI) Atlas is a widely accepted and standardized radiographic assessment tool that takes into account osteophyte formation and joint-space narrowing in the medial and lateral tibiofemoral compartments\textsuperscript{9}. Distinct contributions of each osteophyte size and location and each joint-space width are graded and added together for a final, total OARSI score. Each radiographic element’s contribution is weighted equally, thus providing a more standardized and more granular assessment than the KL grade (Fig. 1).

The goal of this study was to investigate the association between preoperative radiographic OA severity, as measured by the OARSI score, and patient-reported outcomes following total knee replacement.

\section*{Materials and Methods}

\subsection*{Sample}

The Adding Value in Knee Arthroplasty (AViKA) Postoperative Care Navigation Trial is a 2-arm randomized controlled trial that evaluated motivational interviewing to enhance rehabilitation following total knee replacement\textsuperscript{17}. Three hundred and eight consecutive patients undergoing primary unilateral total knee replacement at a tertiary medical center were enrolled. Eligible patients were community-dwelling adults $\geq 40$ years of age who had a primary diagnosis of OA and spoke English. Excluded were patients with an underlying diagnosis other than OA (e.g., inflammatory arthritis), psychological issues precluding participation, or dementia, non-English speakers, nursing home residents, and patients with plans for additional elective surgery within 6 months. The primary outcome variable investigated in that study\textsuperscript{17} was improvement in function as measured by the change in Western Ontario and McMaster Universities Osteoarthritis Index (WOMAC) function score from preoperatively to 6 months postoperatively. Our final analysis included patients who had radiographs available at baseline as well as outcomes available at both baseline and 2 years.

\subsection*{Data Elements}

\subsection*{Radiographic Assessments}

The large majority of preoperative radiographs were posterior-anterior (PA) flexed weight-bearing views. We assessed

\begin{figure}[h]
\centering
\includegraphics[width=\textwidth]{fig1.png}
\caption{Figs. 1-A, 1-B, and 1-C The Osteoarthritis Research Society International (OARSI) score offers a more standardized and granular radiographic assessment than does Kellgren-Lawrence (KL) grade. All 3 knee radiographs demonstrate KL grade-4 osteoarthritis (OA) given definite osteophyte formation and severe joint-space narrowing. However, the OARSI score is different for each radiograph, demonstrating differences in the presence of osteophytes and differences in medial and lateral involvement. Fig. 1-A Advanced knee OA: KL grade 4, OARSI score of 17. Fig. 1-B Advanced knee OA: KL grade 4, OARSI score of 9. Fig. 1-C Advanced knee OA: KL grade 4, OARSI score of 8.}
\end{figure}
Kellgren and Lawrence grade was calculated on the basis of the method described by and compartment-specific analyses included KL grade, compartment-specific joint-space narrowing (JSN) scores. KL grade was calculated on the basis of the method described by Kellgren and Lawrence, with 0 representing no abnormalities and 4 representing the most severe arthritis possible. The compartment-specific OARSI score was calculated according to the OARSI method, but only included the score from a single tibiofemoral compartment, with 0 representing the least severe findings possible and 9 representing the most severe possible. The compartment-specific JSN score was calculated for individual compartments alone according to the OARSI method, with 0 representing no joint-space narrowing and 3 representing the most joint-space narrowing possible.

**Data Collection and Outcomes**
Participants completed a baseline questionnaire within 6 weeks prior to surgery and were asked to complete another questionnaire 2 years following total knee replacement. Primary outcomes were the WOMAC pain score and the Knee Injury and Osteoarthritis Outcome Score (KOOS) activities of daily living (ADL) subscore. WOMAC pain scores were converted to a 0-to-100 scale, with 100 indicating the worst pain possible, per convention. KOOS ADL subscores were converted to a 0-to-100 scale, with 0 indicating the greatest knee dysfunction possible, per convention.

**Covariates**
We assessed demographics and preoperative clinical characteristics of the cohort at baseline, including age, race, sex, body mass index (BMI), and Charlson Comorbidity Index (Table I). BMI was calculated from self-reported height and weight. We stratified BMI into 5 groups: <25.0, 25 to 29.9, 30 to 34.9, 35 to 39.9, and 40.0 kg/m². Charlson Comorbidity Index scores were grouped as follows: 0, 1, 2, and ≥3.

**Statistical Analysis**
Initially, we assessed the relationship between preoperative OARSI score and patient-reported outcomes with scatterplots. We assessed the association between OARSI score and patient-reported outcomes using Pearson (baseline and 2-year change) or Spearman (2-year outcomes, which were skewed) correlation coefficients and linear regression. We assessed the unadj usted association between the binary radiographic OA severity measure and postoperative patient-reported outcomes by a t test or Wilcoxon rank-sum test, as appropriate. We used multivariable linear regression to adjust for age, sex, BMI, and comorbidity score. For baseline and change in patient-reported outcomes, we used the untransformed outcome measure; due to the skewed nature of the 2-year outcome scores, we used the cube root transformation for statistical testing.

We performed sensitivity analyses to examine the association between outcomes and alternate measures of baseline OA severity. These included (1) KL grade (<4 versus 4), (2) compartment-specific (i.e., medial and lateral) OARSI scores, (3) compartment-specific (medial and lateral) JSN scores, and (4) 4-level OARSI score based on quartiles. We investigated KL grade because of its wide use in previous studies. We investigated compartment-specific OARSI scores because total knee replacement may be performed for unicompartmental disease. We investigated compartment-specific joint-space narrowing because it is one major driver of clinical indication for total knee replacement. For compartment-specific JSN scores, we performed an unadjusted analysis of variance (ANOVA) or Kruskal-Wallis test, as appropriate.

KL grade and compartment-specific JSN score subcategories indicating radiographically milder or more severe OA were created on the basis of the distribution of KL grades or compartment-specific JSN scores, respectively, within the cohort. Compartment-specific OARSI score subcategories of low (“milder OA”) and high (“more severe OA”) were created by dichotomizing compartment-specific OARSI scores within the cohort at the median for medial (median = 6) and lateral (median = 4) compartments individually. A 4-level compartment-specific OARSI score metric was created by describing each combination of subcategories as a separate entity: (1) low medial, low lateral; (2) low medial, high lateral; (3) high medial, low lateral; and (4) high medial, high lateral.

We compared the radiographic features between those who completed the 2-year survey and those who missed 2-year follow-up in order to assess the impact of missing data.

All analyses were conducted in SAS version 9.4. (SAS Institute).

**Results**
Of the 308 enrolled subjects, 304 had radiographs available for analysis (see “Radiographic Assessments” above). Of these 304 subjects, 240 (78.95%) completed baseline and 2-year outcome surveys (see “Data Collection and Outcomes” above). These 240 subjects represent our final analytic cohort.

The mean age at surgery was 66.6 years (standard deviation [SD], 8 years), and 61% of the subjects were female. The median preoperative OARSI radiographic severity score was 10 (range, 3 to 17); 102 (42.5%) of the subjects had an OARSI score of <10. Of note, OARSI scores were similar between subjects in the analytic cohort and those excluded because of missing questionnaire data (Table I). Prior to total knee replacement, study participants reported a mean WOMAC pain score of 39.1 (SD, 17). We did not observe a significant or
clinically relevant association between preoperative OARSI score and preoperative patient-reported pain and functional limitation (Table II). Scatterplots of the 2-year change in scores for WOMAC pain and KOOS ADL provide a clear depiction of the lack of linear association with OARSI score (Appendix Figure 1). The correlation between patient-reported outcomes...
and OARSI score ranged from −0.036 (95% confidence interval [CI], −0.162 to 0.091) for the WOMAC pain score at 2 years to 0.079 (95% CI, −0.048 to 0.204) for the KOOS ADL subscore at 2 years (Table II).

Demographic characteristics and radiographic OA severity scores were similar between subjects in the analytic cohort and those who were excluded from our analysis; however, those with complete data exhibited superior self-reported baseline pain, function, and mental health (Table I).

The cohort improved substantially over 2 years following total knee replacement, with improvements in WOMAC pain and KOOS ADL scores on the order of 30 points (0-to-100 scale) (Table III). Low postoperative WOMAC pain scores among the majority of the cohort were indicative of excellent outcomes. We did not observe a significant or clinically important difference in pain relief between subjects with milder and more severe radiographic OA (change in WOMAC pain score, −30.7 for those with an OARSI score of <10 versus −32.6 for those with an OARSI score of ≥10; difference, 1.9 [95% CI, −2.4 to 6.2]) (Table III). The results were confirmed in multivariable analyses that adjusted for age, sex, BMI, and comorbidity. Similarly, we did not observe a significant or clinically important difference in functional improvement, as assessed by change in KOOS ADL subscores, between subjects

### Table II: Association Between Patient-Reported Outcome Scores at Baseline and 2 Years Postoperatively and Continuous Preoperative Osteoarthritis Research Society International (OARSI) Score *

| Patient-Reported Outcome | Unadjusted | Adjusted |
|--------------------------|------------|----------|
|                          | Correlation Coefficient (95% CI) | Parameter Estimate (95% CI) | Parameter Estimate (95% CI) |
| WOMAC pain               |            |          |          |
| Baseline                 | 0.012 (−0.114, 0.139) | 0.08 (−0.74, 0.90) | −0.23 (−1.06, 0.59) |
| 2 yr                     | −0.036 (−0.162, 0.091) | −0.17 (−0.79, 0.44) | −0.37 (−1.00, 0.27) |
| Change from baseline to 2 yr | −0.041 (−0.166, 0.087) | −0.25 (−1.06, 0.55) | −0.13 (−0.98, 0.72) |
| KOOS ADL                 |            |          |          |
| Baseline                 | −0.004 (−0.131, 0.122) | −0.03 (−0.81, 0.76) | 0.23 (−0.57, 1.03) |
| 2 yr                     | 0.079 (−0.048, 0.204) | 0.41 (−0.25, 1.08) | 0.63 (−0.06, 1.32) |
| Change from baseline to 2 yr | 0.067 (−0.06, 0.192) | 0.44 (−0.39, 1.27) | 0.4 (−0.48, 1.29) |

*Unadjusted correlations are from Pearson (baseline, 2-year change) or Spearman (2-year outcome) correlation. Unadjusted parameter estimates are from linear regression, and adjusted estimates are from multivariable linear regression, adjusting for age, sex, BMI, and comorbidity score. The parameter estimate is interpreted as the increase in patient-reported outcome associated with a 1-unit increase in baseline OARSI score. WOMAC (Western Ontario and McMaster Universities Osteoarthritis Index) pain is scaled 0 to 100, with 100 being the worst pain.

### Table III: Patient-Reported Outcome Scores at Baseline and 2 Years Postoperatively, Stratified by Preoperative Osteoarthritis Research Society International (OARSI) Score *

| Patient-Reported Outcomes | OARSI Score <10 | OARSI Score ≥10 | Unadjusted Difference Between Groups (95% CI) | Adjusted Difference Between Groups (95% CI) |
|--------------------------|-----------------|-----------------|-----------------------------------------------|---------------------------------------------|
| WOMAC pain               |                 |                 |                                               |                                             |
| Baseline                 | 39.0 (16.9)     | 39.1 (17.5)     | 0.0 (−4.5, 4.4)                              | 0.68 (−3.75, 5.1)                          |
| 2 yr                     | 8.3 (14.8)      | 6.5 (11.2)      | 1.8 (−1.5, 5.2)                              | 0.18 (−0.15, 0.5)                          |
| Change from baseline to 2 yr | −30.7 (17.8) | −32.6 (16.1) | 1.9 (−2.4, 6.2) | 1.62 (−2.92, 6.17) |
| KOOS ADL                 |                 |                 |                                               |                                             |
| Baseline                 | 60.1 (15.8)     | 60.6 (17.0)     | −0.5 (−4.8, 3.7)                             | −1.11 (−5.4, 3.17)                         |
| 2 yr                     | 88.7 (16.7)     | 91.8 (11.4)     | −3.1 (−6.7, 0.5)                             | −0.1 (−0.2, 0.01)                          |
| Change from baseline to 2 yr | 28.6 (18.3) | 31.2 (17.0) | −2.6 (−7.1, 1.9) | −2.53 (−7.3, 2.24) |

*Patient-reported outcome scores are reported separately for individuals with radiographically milder osteoarthritis (OARSI score of <10) and more severe osteoarthritis (OARSI score of ≥10). Continuous variables are given as the mean, with the standard deviation in parentheses. “Difference between groups” indicates the difference in mean outcome scores between OARSI score categories as assessed by t test or Wilcoxon rank-sum test, as appropriate, for unadjusted analyses and linear regression for adjusted analyses. WOMAC (Western Ontario and McMaster Universities Osteoarthritis Index) pain is scaled 0 to 100, with 100 being the worst pain. KOOS ADL (Knee injury and Osteoarthritis Outcome Score activities of daily living) is scaled 0 to 100, with 100 being the best function. CI = confidence interval. No significant associations were found.
Change in Western Ontario and McMaster Universities Osteoarthritis Index (WOMAC) pain score from baseline to 2 years by preoperative radiographic osteoarthritis (OA) severity. Individuals with mild and severe radiographic arthritis, as defined by both Kellgren-Lawrence (KL) and Osteoarthritis Research Society International (OARSI) criteria preoperatively, experienced meaningful pain relief at 2 years postoperatively with no significant differences between the groups. The top and bottom of the boxes = the 25th and 75th percentile, the horizontal line within the boxes = the median, the “x” within the boxes = the mean, the circles = outliers, and the whiskers = the minimum and maximum excluding outliers.

| Patient-Reported Outcome | KL <4 | KL = 4 | Difference Between Groups (95% CI) |
|--------------------------|-------|--------|-----------------------------------|
| WOMAC pain               |       |        |                                   |
| Baseline                 | 43.0 (18.0) | 36.9 (16.4) | 6.1 (1.6, 10.6) |
| 2 yr                     | 9.2 (15.5)  | 6.2 (11.1)  | 3.0 (−0.5, 6.4) |
| Change from baseline to 2 yr | −33.8 (16.2) | −30.7 (17.1) | −3.1 (−7.6, 1.3) |
| KOOS ADL                 |       |        |                                   |
| Baseline                 | 57.8 (16.8) | 61.8 (16.2) | −4.0 (−8.4, 0.3) |
| 2 yr                     | 88.5 (17.4) | 91.6 (11.6) | −3.1 (−6.8, 0.6) |
| Change from baseline to 2 yr | 30.7 (16.9)  | 29.8 (17.9)  | 0.9 (−3.7, 5.6)  |

*Patient-reported outcome scores are reported separately for individuals with milder radiographic arthritis (KL <4) and more severe radiographic arthritis (KL = 4). Continuous variables are given as the mean, with the standard deviation in parentheses. “Difference between groups” indicates the difference in mean outcome scores between KL categories as assessed by t test or Wilcoxon rank-sum test, as appropriate. WOMAC (Western Ontario and McMaster Universities Osteoarthritis Index) pain is scaled 0 to 100, with 100 being the worst pain. KOOS ADL (Knee injury and Osteoarthritis Outcome Score activities of daily living) is scaled 0 to 100, with 100 being the best function. CI = confidence interval. Bold indicates a significant association.
with radiographically milder and more severe OA (28.6 for those with milder OA versus 31.2 for those with more severe OA; difference, −2.6 [95% CI, −7.1 to 1.9]) (Table III). Results were similar in multivariable models adjusted for age, sex, BMI, and comorbidity score.

In sensitivity analyses, we compared pain relief and functional improvement between subjects with milder and more severe radiographic findings utilizing 4 other radiographic assessments: KL grade (Fig. 2, Table IV), compartment-specific OARSI scores (Table V), compartment-specific JSN scores (Appendix Tables 1 and 2), and 4-level OARSI score category (Appendix Table 3). In each case, we did not observe clinically important differences in pain or function scores between subjects with radiographically milder and more severe OA.

Discussion

We did not observe a clinically important association between preoperative radiographic severity of OA and patient-reported outcomes 2 years following total knee replacement. Subjects with milder and more severe radiographic OA severity demonstrated meaningful and similar pain relief and functional improvement after total knee replacement. Additional sensitivity analyses supported this finding: KL grade, compartment-specific OARSI score, and compartment-specific JSN score were not associated with meaningful differences in pain relief or functional improvement.

Our study elucidates the distinct contributions of joint-space narrowing and compartment-specific tibiofemoral disease, providing a more granular analysis than previous studies. These findings may help clinicians in counseling patients that radiographic OA severity is not predictive of total knee replacement outcome. One should not infer from our findings, however, that evidence of radiographic OA alone is an indication for total knee replacement. Rather, our findings suggest that if total knee replacement is indicated for an individual patient on the basis of multiple clinical criteria, his or her outcome following total knee replacement will not be predicted by preoperative radiographic OA severity.

Our findings are consistent with those of prior literature. In an investigation that included 1,888 patients who underwent total knee replacement, Meding et al. utilized a modified KL grading system that dichotomized radiographic findings as “severe” or “mild,” with severe defined as at least 1 compartment with severe osteophytosis or bone-on-bone changes. The authors did not find clinically important differences in post-total knee replacement pain and functional improvement between individuals with severe versus mild radiographic arthritis despite identifying some small statistically significant differences.

In a study including 478 patients who underwent total knee replacement, Dowsey et al. utilized a modified KL grading system for radiographic OA assessment. The key findings of that report were that pain relief was unsatisfactory in about 30% of the cohort and functional improvement was suboptimal in about 50% of the cohort. The authors showed that, at 12 months, the likelihood of having suboptimal pain outcomes was substantially higher among those with less OA radiographic severity. The differences in findings described by Dowsey et al. and those in the current analysis could be explained mainly by the difference in outcomes. Compared with our cohort, the cohort described by Dowsey et al. had much worse outcomes. In addition, we used different pain scales (IKKS [International Knee Society score] in that study versus WOMAC in ours), which may have contributed to the difference in findings.

Other studies have reported on the association between radiographic severity of OA and outcomes of total knee replacement. Kahn et al. reported on outcomes among Osteoarthritis Initiative subjects, demonstrating a very weak correlation between radiographic OA severity and post-total knee replacement patient-reported outcomes. Keurentjes
et al. reported on the association between generic quality-of-life measures and radiographic severity of knee OA. While the authors showed that higher radiographic severity was associated with better quality of life 2 to 5 years post-total knee replacement, the response rate for the study was <50%, raising questions regarding the generalizability of the findings.

Of note, the presence of joint-space narrowing is typically required to reach higher KL grades, while the OARSI score weights joint-space narrowing, tibial osteophyte presence, and femoral osteophyte presence equally in contribution to the final score. Thus, joint-space narrowing may be given more weight in determining radiographic arthritis severity when KL-based methods are employed than when OARSI-based methods are employed. On the basis of our results and the results of others, it appears that differences in preoperative joint-space narrowing alone may be associated with small differences in patient-reported outcomes after total knee replacement. However, our data and others’ suggest that these differences do not appear to be clinically important.

Some reports did not demonstrate an association between preoperative radiographic OA severity and patient-reported outcomes following total knee replacement. Perry et al. investigated the results of 62 total knee replacements at 2 to 10-year follow-up, finding no difference in Knee Society scores between individuals with milder and more severe preoperative radiographic arthritis as determined by KL grade. Similarly, Chang et al. reported on the outcomes of 383 total knee replacement procedures at minimum of 1 year of follow-up, observing no association between radiographic OA grade and patient-reported outcomes. Importantly, the authors noted that all patients in their study had evidence of at least 75% joint-space narrowing, which implies that all patients would likely be classified as KL grade 4 in our study. In addition, the authors utilized the modified Ahlbäck radiographic scoring system, a semiquantitative radiographic assessment that considers a different complement of radiographic findings than does the KL grading system. These studies align with the overall message of our study and others that there do not appear to be clinically important differences in postoperative outcomes between those subjects with milder versus more severe preoperative radiographic arthritis.

Our study had several limitations. First, we did not evaluate all sources of intra-articular damage, especially focal cartilage defects, which may not be apparent on radiographs. Second, tibiofemoral alignment was not assessed, and this is known to be a possible contributor to differential outcome following total knee replacement. However, we assessed the role of compartment-specific OA severity, which may be a proxy for preoperative alignment. Third, this report represents total knee replacements performed at a single high-volume joint replacement center, and the surgeon and patient populations may not be representative of other surgeon and patient populations. Fourth, the decision-making process for undergoing total knee replacement was not assessed in these cases, including the utilization of advanced imaging, which may have illuminated why patients with relatively mild radiographic OA fared as well as those with severe radiographic OA after total knee replacement. Finally, we assessed the severity of baseline radiographic damage in several ways, and assessed several patient-reported outcomes, and we did not adjust for multiple testing. However, we did not find significant associations between OARSI score and patient-reported outcome in any of the primary analyses, so adjustment for multiple testing is not pertinent.

In our cohort, regardless of preoperative radiographic OA severity, subjects experienced clinically meaningful postoperative pain relief and functional improvement. We did not observe a clinically meaningful association between preoperative radiographic OA severity and patient-reported outcomes following total knee replacement. We emphasize that these findings reflect the outcomes of individuals already indicated for total knee replacement surgery, and that radiographic OA assessment should not be utilized alone to indicate a patient for total knee replacement.

Appendix

Supporting material provided by the authors is posted with the online version of this article as a data supplement at jbjs.org (http://links.lww.com/JBJSOA/A191).

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