

## Original Research

## Patients With Lymphedema are at Increased Risk of Complication After Total Knee Arthroplasty: A Population Level Study

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## ABSTRACT

**Background:** Lymphedema, a chronic disorder characterized by abnormal lymphatic fluid buildup, most commonly affects the lower extremities. Limited literature exists regarding the impact of lymphedema on outcomes following primary total knee arthroplasty (TKA). This study examined the effect of lymphedema's impact on complications and early revision-free survivorship following primary TKA using population-level data. We hypothesized that patients with preoperative lymphedema would have higher complication and revision rates.

**Methods:** Patients undergoing TKA for osteoarthritis between 2009 and 2020 were identified from a national claims database. Those with preoperative lymphedema were matched 1:1 to contemporaries without lymphedema using propensity score matching. Comparisons between matched and unmatched cohorts were performed using Chi-square and independent t-tests, while Cox proportional hazards models assessed revision risk.

**Results:** Of the 530,938 TKA patients, 1.05% ( $n = 5602$ ) had preoperative lymphedema. Matched analysis showed lymphedema had higher 90-day rates of periprosthetic joint infection (2.9% vs 1.4%,  $P < .001$ ), superficial surgical site infection (2.3% vs 1.6%,  $P = .007$ ), wound complications (2.4% vs 1.7%,  $P = .013$ ), and pulmonary embolism (6.6% vs 4.6%,  $P < .001$ ). At 2 years, lymphedema was associated with increased risk of all-cause (hazard ratio (HR) = 1.42,  $P < .001$ ) and septic revisions (HR = 1.88,  $P < .001$ ) but not aseptic revisions (HR = 0.99,  $P = .929$ ).

**Conclusions:** Preoperative lymphedema is associated with increased 90-day rates of periprosthetic joint infection and superficial surgical site infection, wound complications, and pulmonary embolism after primary TKA. Although aseptic revision risk was not increased, the association with higher all-cause and septic revisions warrants attention. These findings emphasize the need for aggressive counseling and preoperative optimization before TKA in lymphedema patients.

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## Introduction

Total knee arthroplasty (TKA) is a common procedure indicated for the definitive treatment of severe, symptomatic osteoarthritis (OA) of the knee. While complication rates have decreased from historic standards following TKA [1], periprosthetic joint infection (PJI) continues to be a devastating complication with a reported incidence rate between 0.3% and 1.9% [2]. While risk factors for PJI

such as obesity, diabetes mellitus, and renal disease have been well described, there is a relative paucity of literature regarding the association between PJI and lymphedema.

Lymphedema is a chronic disease characterized by impaired lymphatic drainage and subsequent accumulation of fluid in the interstitium [3]. Lymphedema can occur as a primary disease due to abnormal lymphatic development or as a secondary pathology associated with the damage or obstruction of previously normal lymphatic vessels caused by disease processes, recurrent infections, trauma, surgery, or radiation treatment [3]. In arthroplasty, small, single-institutional series have linked lymphedema to higher rates of complications following TKA, including increased rates of PJI and superficial wound infections.

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While informative, the existing literature regarding lymphedema and outcomes following TKA is limited by small numbers of included patients from single institutions [4,5]. Therefore, there is a need for population-level data to clarify and confirm the results of these prior investigations. Therefore, the purpose of this study was to examine the influence of lymphedema on outcomes following primary TKA using a large national database. Preoperative lymphedema was hypothesized to be associated with increased complication and revision rates.

## Material and methods

### Data source

This study utilized data from the Merative MarketScan Commercial Claims and Encounters database, which also includes the Medicare Supplemental and Coordination of Benefits database (Merative, Ann Arbor, MI). This dataset encompasses insurance claims from both commercial and Medicare health plans across the United States, covering approximately 250 million patient records. Procedural and diagnostic data were extracted using the Ninth and Tenth Revisions of the International Classification of Diseases (ICD-9 and ICD-10) as well as Current Procedural Terminology (CPT) codes (Appendix A-D). Since the dataset is fully deidentified, the study was deemed exempt from institutional review board approval.

### Study design and patient selection

Patients who underwent primary TKA for OA between January 1, 2009, and December 31, 2020, were identified using CPT code 27447 and OA-related ICD-9/10 codes as listed in Appendix A (n = 815,850). To ensure accurate reporting, patients with records indicating bilateral TKA were excluded to avoid potential complications from the contralateral procedure due to inconsistent laterality designators in the database (n = 121,134). Additionally, patients under 18 years of age (n = 233) and patients without a diagnosis of primary OA (n = 23,330) were excluded. Furthermore, patients with an additional diagnosis of traumatic arthritis (n = 2530), rheumatoid arthritis (n = 72,635), or a history of a fracture around the knee (n = 315) were excluded. Finally, individuals with less than 90 days of follow-up (n = 64,735) were excluded. Thus, the final cohort consisted of 530,938 patients (Fig. 1).

Propensity score matching was used to divide the study population into 2 cohorts. The control cohort consisted of patients without a presurgical diagnosis of lymphedema, while the lymphedema cohort included patients with a presurgical diagnosis of lymphedema. Matching was performed in a 1:1 ratio based on age, sex, Elixhauser comorbidities, Elixhauser Comorbidity Index (ECI), active smoking status, prior ischemic stroke, and prior myocardial infarction. This resulted in a final matched cohort of 5596 patients in each group.

### Baseline patient data and comorbidities

Baseline characteristics, including sex, age, active smoking, and alcohol use, were collected along with major comorbidities, such as congestive heart failure, rheumatoid arthritis, diabetes, liver disease, renal failure, obesity, and neurological disorders. The overall comorbidity burden was quantified using the ECI for each patient [6]. Patient comorbidities were determined using ICD-9 and ICD-10 codes assigned to insurance claims within 1 year of the index TKA (Appendix B).

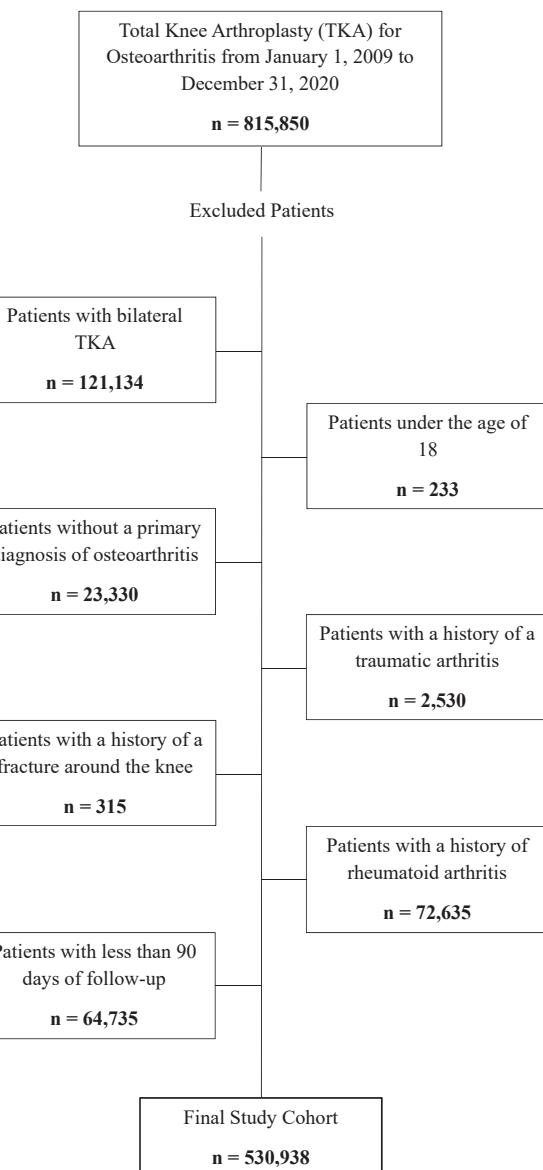


Figure 1. Study population exclusion criteria flow chart.

### Study outcomes

The outcomes evaluated were 90-day complication rates and 2-year survivorship outcomes. Complications assessed included PJI, septic revision, superficial surgical site infections (SSIs), wound complications, periprosthetic fractures, sepsis, hematoma, deep vein thrombosis, pulmonary embolism (PE), myocardial infarction, and pneumonia (Appendix C). The number of readmissions within 90 days and the number of aseptic revision surgeries were also recorded.

Extended length of stay was defined as hospital admission exceeding 2 days postindex surgery. Survivorship at 2 years free from all-cause revision, septic revision, and aseptic revision was evaluated. These outcomes were identified using ICD-9, ICD-10, and CPT codes for revision TKA, cross-referenced with primary and secondary diagnosis codes to establish the indication for revision (Appendix D). We utilized published methodology for the identification of septic and aseptic revision procedures [7].

## Data analyses

Patient characteristics and comorbidities in the unmatched and matched cohorts were compared using Chi-square tests for categorical variables and independent 2-sample t-tests for continuous variables. Ninety-day complications were also compared using Chi-square tests. Kaplan–Meier curves were constructed to evaluate 2-year survivorship free from all-cause revision, septic revision, and aseptic revision. Hazard ratios (HRs) for revision surgeries were calculated using Cox proportional hazards models, adjusting for clinically relevant covariates. The proportional hazards assumption was tested to ensure model validity. Statistical significance was set at a *P* value of  $<0.05$  for all tests.

## Results

### Patient demographic characteristics and comorbidities

Of the included 530,938 patients who were initially included, 5602 patients had a presurgical diagnosis of lymphedema (1.05%). Prior to matching, the lymphedema cohort was older (64 vs 63, *P* < .001), less often men (28.8% vs 41.1%, *P* < .001), and had higher comorbid burden (ECI 5.46 vs 3.17, *P* < .001). Specifically, the lymphedema cohort had higher rates of obesity, alcohol use disorder, congestive heart failure, diabetes mellitus, renal failure, liver disease, and neurological disorders. However, there were more smokers in the control group (6.7% vs 6.0%, *P* < .001).

After matching, there were matched cohorts consisting of 5596 patients with and without lymphedema. Matching provided well-balanced cohorts with no significant differences in age, sex, ECI, or individual comorbidities (Table 1).

### Ninety-Day complication outcomes

Within 90 days of surgery, patients in the lymphedema cohort experienced significantly higher rates of complications compared to the control group. The incidence of PJI was 2.9% in the lymphedema cohort, compared to 1.4% in the control group (*P* = .001). Similarly, the rates of SSI and wound complications were higher in the lymphedema group (2.3% vs 1.6%, *P* = .007 and 2.4% vs 1.7%, *P* = .013, respectively). Other notable differences included a higher incidence of hematoma (1.2% vs 0.6%, *P* = .004), PE (6.6% vs 4.6%, *P* < .001), and extended length of stay (9.3% vs 8.2%, *P* = .038) in the lymphedema cohort.

**Table 1**

Patient demographic characteristics and comorbidities: Unmatched and matched cohorts.

Total, n	Unmatched cohorts			Matched cohorts		
	Control (525,336)	Lymphedema (5602)	<i>P</i> value <sup>a</sup>	Control (5596)	Lymphedema (5596)	<i>P</i> value <sup>a</sup>
Age, Mean (range)	63 (19, 100)	64 (25, 94)	<b>&lt;0.001</b>	65 (19, 94)	64 (25, 94)	0.297
Sex			<b>&lt;0.001</b>			0.616
Men	215,707 (41.1)	1615 (28.8)		1591 (28.4)	1615 (28.9)	
Women	309,629 (58.9)	3987 (71.2)		4005 (71.6)	3981 (71.1)	
Elixhauser index (mean, sd)	3.17 (2.35)	5.46 (3.23)	<b>&lt;0.001</b>	5.4 (3.3)	5.4 (3.2)	0.820
Obesity (n,%)	104,398 (19.9)	2357 (42.1)	<b>&lt;0.001</b>	2392 (42.7)	2352 (42.0)	0.456
Active smoking (n,%)	35,162 (6.69)	334 (5.96)	<b>0.031</b>	293 (5.24)	332 (5.93)	0.118
Alcohol use disorder (n,%)	9307 (1.77)	153 (2.73)	<b>&lt;0.001</b>	139 (2.48)	153 (2.73)	0.441
CHF (n,%)	30,812 (5.87)	921 (16.4)	<b>&lt;0.001</b>	915 (16.4)	915 (16.4)	>0.999
Diabetes (n,%)	158,628 (30.2)	2263 (40.4)	<b>&lt;0.001</b>	2319 (41.4)	2259 (40.4)	0.257
Renal failure (n,%)	27,449 (5.23)	687 (12.3)	<b>&lt;0.001</b>	689 (12.3)	685 (12.2)	0.931
Liver disease (n,%)	31,919 (6.08)	686 (12.2)	<b>&lt;0.001</b>	674 (12.0)	683 (12.2)	0.817
Neurological disorders (n,%)	19,087 (3.63)	414 (7.39)	<b>&lt;0.001</b>	411 (7.34)	413 (7.38)	0.971

CHF, congestive heart failure.

Bolded *P* values indicate statistical significance (*P* < 0.05).

<sup>a</sup> Welch 2-Sample t-test; Pearson's Chi-squared test.

**Table 2**

Ninety-Day complication outcomes.

Complications	Control N = 6993 (%)	Lymphedema N = 6993 (%)	<i>P</i> value <sup>a</sup>
PJI	77 (1.4)	160 (2.9)	<b>&lt;0.001</b>
SSI	87 (1.6)	127 (2.3)	<b>0.007</b>
Wound complications	95 (1.7)	133 (2.4)	<b>0.013</b>
Periprosthetic Fx	7 (0.1)	8 (0.1)	1.000
Hematoma	35 (0.6)	65 (1.2)	<b>0.004</b>
DVT	28 (0.5)	36 (0.6)	0.380
PE	257 (4.6)	372 (6.6)	<b>&lt;0.001</b>
MI	117 (2.1)	122 (2.2)	0.794
PNA	41 (0.7)	25 (0.4)	0.064
Readmissions	122 (2.2)	109 (1.9)	0.425
Extended LOS	460 (8.2)	523 (9.3)	<b>0.038</b>
Aseptic revision surgery	2984 (53)	2980 (53)	0.955

Fx, Fracture; DVT, deep venous thrombosis; MI, myocardial infarction; PNA, pneumonia.

Bolded *P* values indicate statistical significance (*P* < 0.05).

<sup>a</sup> Pearson's Chi-squared test.

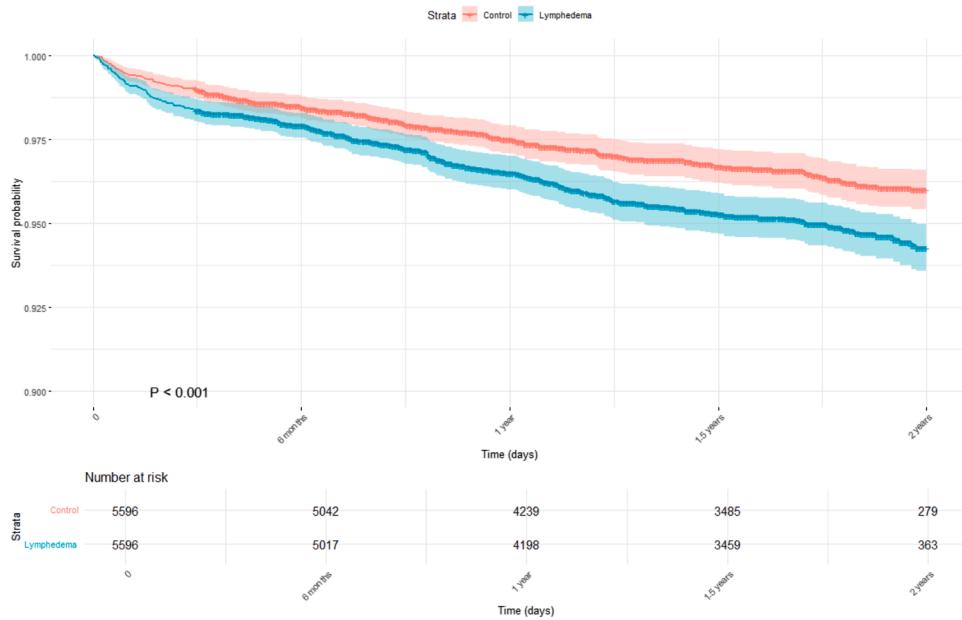
No significant differences were observed for periprosthetic fractures (0.1% vs 0.1%, *P* = 1.000, or other major complications such as deep vein thrombosis, myocardial infarction, or pneumonia. There were also no significant differences between the 2 groups for readmissions or aseptic revision surgery within 90 days of index surgery (Table 2).

### Two-year survivorship

At 2 years, Kaplan–Meier survivorship free from all-cause revision was 94.3% in the lymphedema group vs 96.0% in the control group (Fig. 2). Survivorship free from septic revision was 96.4% vs 98.2%, respectively (Fig. 3). There was no notable difference in aseptic revision (Fig. 4). In multivariable Cox proportional hazards analysis, lymphedema was associated with a significantly increased risk of all-cause revision (HR: 1.42, 95% confidence interval (CI): 1.18–1.72), and septic revision (HR: 1.88, 95% CI: 1.45–2.44). There was no significant difference in aseptic revision risk (HR: 0.99, 95% CI: 0.74–1.31) (Table 3).

## Discussion

The aim of this study was to examine 90-day complication outcomes and 2-year survivorship following primary TKA in

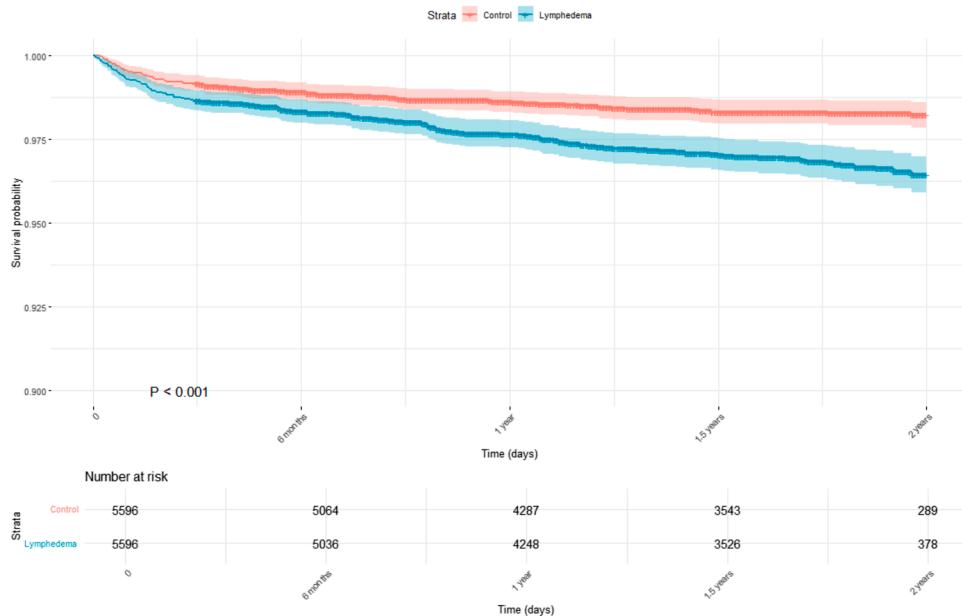


**Figure 2.** Two-year survivorship free of all-cause revision.

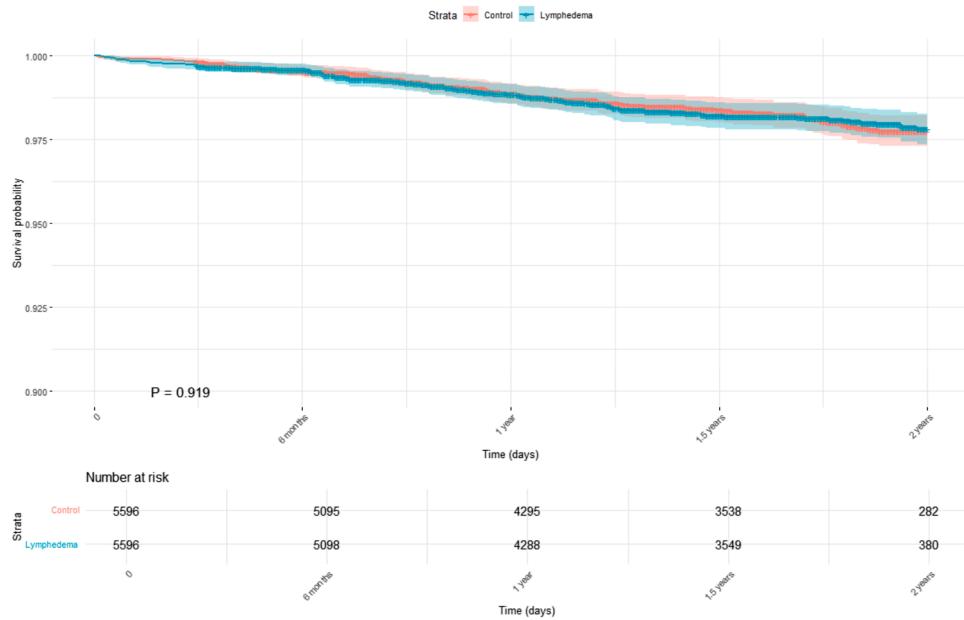
patients with lymphedema using population-level data from a national database. The results demonstrate that patients with lymphedema undergoing TKA are at significantly higher risk for postoperative complications compared to those without lymphedema. After appropriate matching, which balanced the cohorts for age, sex, and comorbidities, the lymphedema group had a higher incidence of PJI, SSI, wound complications, hematoma, and PE within 90 days of surgery. Perhaps more importantly, lymphedema was found to compromise 2-year survivorship, with significantly lower rates of survivorship free of all-cause and septic revision.

The study's results are consistent with smaller, single-institution studies of patients undergoing primary TKA. Prior studies have reported high infection rates in this population.

Shrader et al. Reported infection in 19% of patients (superficial and deep infections in 12% and 7%, respectively) among 63 individuals undergoing primary TKA [5]. Cusma et al. similarly reported a 9% infection rate and 9% revision/reoperation rate in a cohort of 67 patients with preoperative lymphedema [4]. Kolz et al., in a larger series of 432 patients, reported significantly higher rates of SSIs in patients with lymphedema compared to controls, with an infection rate of 9.7% vs 1.0%, respectively, as well as increased readmissions [8]. In contrast, our study found more modest rates of PJI in the lymphedema group (2.9%), though this remained significantly higher than in comorbidity-matched controls (1.4%). At 2 years, we also found that lymphedema was associated with an 88% increased hazard of septic revision. Similarly, and perhaps not



**Figure 3.** Two-year survivorship free of septic revision.



**Figure 4.** Two-year survivorship free of aseptic revision.

surprisingly, our results indicated that wound complications were higher in patients with lymphedema. This is consistent with findings from prior authors who also reported elevated rates of wound complications in this population.

An important finding of this study is the lower 2-year survivorship free of all-cause revision in patients with lymphedema. As noted above, this difference was largely due to an increased rate of septic revision. Patients with lymphedema had a significantly higher hazard of septic revision (HR: 1.88, 95% CI: 1.45-2.44), while the risk of aseptic failure was not significantly different between groups (HR: 0.99, 95% CI: 0.74-1.31). The overall risk of all-cause revision was also increased in the lymphedema group (HR: 1.42, 95% CI: 1.18-1.72).

Given the significant morbidity posed by PJI and the difficulty associated with infection eradication in patients with decreased local immune response due to fluid retention, these findings warrant consideration and recognition. These findings highlight the importance of thorough preoperative risk assessment and vigilant postoperative management for patients with lymphedema undergoing TKA.

The association between infection and lymphedema is likely due to a combination of systemic inflammation and the unique pathophysiology of the disease [9]. Lymphedema induces chronic inflammation, which not only disrupts normal immune function

but also fosters a proinflammatory environment that impairs wound healing and tissue repair [10]. Additionally, lymphedema leads to fibrosis of the skin and subcutaneous tissues, which compromises the integrity of the skin barrier, making it more susceptible to bacterial invasion [11].

To combat this heightened risk, a multidisciplinary approach is essential. Collaboration between orthopaedic surgeons, vascular specialists, and lymphedema therapists can optimize preoperative and postoperative care. The use of compression therapy, including compression stockings both before and after surgery, can aid in reducing fluid accumulation and improving lymphatic flow, which may help mitigate infection risk [12]. Additionally, emerging techniques such as micro lymphatic surgery may help to restore lymphatic function in severe cases and could be considered prior to TKA in high-risk patients to reduce complications and improve surgical outcomes [13,14].

The present study substantially expands upon the available data regarding TKA and lymphedema by utilizing population-level data to report on a much larger cohort of patients than has previously been studied. Despite this obvious strength, there are limitations to consider. Database studies are inherently limited by the validity and accuracy of diagnostic codes. Therefore, it is possible that some patients with lymphedema were not identified as such if not properly coded. Furthermore, the database limits the

**Table 3**  
Cox proportional hazards model for all-cause revision and septic revision.

Characteristic	All-cause revision			Septic revision			Aseptic revision		
	HR <sup>a</sup>	95% CI	P value <sup>b</sup>	HR <sup>a</sup>	95% CI	P value <sup>b</sup>	HR <sup>a</sup>	95% CI	P value <sup>b</sup>
Lymphedema	1.42	1.18, 1.72	<b>&lt;.001</b>	1.88	1.45, 2.44	<b>&lt;.001</b>	0.99	0.74, 1.31	.929
Men	1.67	1.38, 2.02	<b>&lt;.001</b>	2.31	1.80, 2.97	<b>&lt;.001</b>	1.00	0.73, 1.37	.985
BMI >35	1.43	1.18, 1.74	<b>&lt;.001</b>	1.53	1.20, 1.97	<b>.001</b>	1.29	0.96, 1.74	.097
Diabetes Mellitus	1.40	1.15, 1.71	<b>.001</b>	1.70	1.29, 2.24	<b>&lt;.001</b>	1.10	0.81, 1.48	.546
Chronic kidney disease	1.24	1.00, 1.55	.053	1.60	1.21, 2.11	<b>.001</b>	0.79	0.54, 1.16	.237
Depression	1.40	1.15, 1.71	<b>.001</b>	1.45	1.12, 1.88	<b>.006</b>	1.32	0.98, 1.78	.072

BMI, body mass index; SD, standard deviation.

Bolded P values indicate statistical significance ( $P < 0.05$ ).

<sup>a</sup> Adjusted hazard ratio.

<sup>b</sup> Wald Test.

evaluation of the severity of a patient's lymphedema. Although this may still be difficult to quantify during clinical evaluation, the degree of lymphedema and skin integrity could affect patient outcomes following TKA. The lower rate of wound complications and infections in our study, compared with smaller single-institution reports, may reflect over-diagnosis of lymphedema in the database. Similarly, the study design does not allow us to know preoperative optimization strategies. It is possible that appropriate optimization (or lack thereof) could influence our results.

## Conclusions

Patients with lymphedema undergoing primary TKA are at increased risk for early postoperative complications—including PJI, wound complications, and PE—as well as reduced revision-free survivorship at 2 years. These findings, drawn from a large, national database, reinforce lymphedema as a clinically meaningful risk factor for complications following primary TKA. Given the heightened susceptibility to infection and wound-related morbidity, lymphedema should be carefully considered during preoperative risk stratification and surgical planning. Multidisciplinary optimization and tailored postoperative care may help mitigate these risks and improve outcomes for this vulnerable patient population.

## Conflicts of interest

Ajay Premkumar is a paid consultant for Accupredict, Smith & Nephew, Stryker, Osgenics, and Naviswiss.

Jacob M. Wilson is a paid consultant for Zimmer Biomet.

The other authors declare no potential conflicts of interest.

For full disclosure statements refer to <https://doi.org/10.1016/j.artd.2025.101900>.

## CRedit authorship contribution statement

**Bryce T. Hrudka:** Writing – review & editing, Writing – original draft, Visualization, Methodology, Investigation, Formal analysis, Data curation, Conceptualization. **Evan Bailey:** Writing – review & editing, Writing – original draft. **Alyssa Woltemath:** Writing – review & editing, Writing – original draft, Conceptualization. **Grayson Nour:** Writing – review & editing, Writing – original draft. **Ajay Premkumar:** Writing – review & editing, Supervision, Methodology, Conceptualization. **Jacob M. Wilson:**

Writing – review & editing, Supervision, Project administration, Methodology, Conceptualization.

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**Appendix A**

Procedure and exclusion codes.

Procedure type	CPT codes	Definition
TKA	27447	<b>TKA</b>
Exclusion Criteria (ICD 9 and 10 Codes)		
Traumatic arthritis of knee	ICD-9	Traumatic arthropathy, unspecified knee
	ICD-10	Unilateral post-traumatic osteoarthritis of knee
Rheumatoid arthritis	ICD-9	Rheumatoid arthritis and other inflammatory polyarthropathies
	ICD-10	Felty's syndrome, knee
		Rheumatoid lung disease with rheumatoid arthritis of knee
		Rheumatoid vasculitis with rheumatoid arthritis of knee
		Rheumatoid heart disease with rheumatoid arthritis of knee
		Rheumatoid myopathy with rheumatoid arthritis of knee
		Rheumatoid polyneuropathy with rheumatoid arthritis of knee
		Other rheumatoid arthritis with rheumatoid factor of knee
		Rheumatoid arthritis without rheumatoid factor, knee
		Rheumatoid bursitis, knee
		Rheumatoid nodule, knee
		Other specified rheumatoid arthritis, knee
Fractures involving knee	ICD-9	Fracture of other and unspecified parts of femur
	ICD-10	Fracture of tibia and fibula
		Fracture of lower end of femur
		Fracture of upper end of tibia

**Appendix B**

Comorbidity ICD codes.

Comorbidity	ICD code	Description	
Active smoking	ICD-9	305.1 V15.82	
	ICD-10	F17 Z87.891	
Obesity	ICD-9	278.00 278.01 278.02 278.03	
	ICD-10	V85.30 to V85.45 E66 Z68.3 Z68.4	
Hypertension	ICD-9	401 to 405	
	ICD-10	I10 to I16	
Diabetes	ICD-9	250	
	ICD-10	E10 E11 E13	
Rheumatic disease	ICD-9	446 701.0 710.0 710.1 710.2 710.3 710.4 710.8 710.9 711.2 714 719.3 720 725 728.5 728.89 729.30	Polyarteritis nodosa and allied conditions Circumscribed scleroderma Systemic lupus erythematosus Systemic sclerosis Sicca syndrome Dermatomyositis Polymyositis Other specified diffuse diseases of connective tissue Unspecified diffuse connective tissue disease Arthropathy in Behcet's syndrome Rheumatoid arthritis and other inflammatory polyarthropathies Palindromic rheumatism Ankylosing spondylitis and other inflammatory spondylopathies Polymyalgia rheumatica Hypermobility syndrome Other disorders of muscle, ligament, and fascia Panniculitis, unspecified site
	ICD-10	L94.0 L94.1 L94.3 M05 M06 M08 M12.0 M12.3 M30 M31.0 M31.1 M31.2 M31.3 M32 M33 M34 M35 M45 M46.1 M46.8 M46.9	Localized scleroderma [morphia] Linear scleroderma Sclerodactyly Rheumatoid arthritis with rheumatoid factor Other rheumatoid arthritis Juvenile arthritis Chronic postrheumatic arthropathy [Jaccoud] Palindromic rheumatism Polyarteritis nodosa and related conditions Hypersensitivity angiitis Thrombotic microangiopathy Lethal midline granuloma Wegener's granulomatosis Systemic lupus erythematosus Dermatopolymyositis Systemic sclerosis [scleroderma] Other systemic involvement of connective tissue Ankylosing spondylitis Sacroiliitis, not elsewhere classified Other specified inflammatory spondylopathies Unspecified inflammatory spondylopathy
Atrial fibrillation	ICD-9	427.31	
Coronary artery disease	ICD-10	I489.1	
Renal disease	ICD-9	410 to 414	
	ICD-10	I20 to I25	
Renal disease	ICD-9	403	
	ICD-10	404 585 586 587 5880 V42.0 V45.11 V45.12 V56	
Renal disease	ICD-10	I12.0	
		I13.1 N18	
		Hypertensive renal disease Hypertensive heart and renal disease Chronic renal failure Renal failure, unspecified Renal sclerosis, unspecified Renal osteodystrophy Kidney replaced by transplant Renal dialysis status Noncompliance with renal dialysis Encounter for dialysis and dialysis catheter care Hypertensive chronic kidney disease with stage 5 chronic kidney disease or end stage renal disease Hypertensive heart and chronic kidney disease without heart failure Chronic kidney disease	

## Appendix B (continued)

Comorbidity	ICD code	Description
Lung disease	ICD-9	N19 N25.0 Z49 Z94.0 Z99.2 416.8 416.9 490 491 492 493 494 495 496 500 501 502 503 504 505 506.4 508.1 508.8
		Unspecified kidney failure Renal osteodystrophy Encounter for care involving renal dialysis Kidney transplant status Dependence on renal dialysis Other chronic pulmonary heart diseases Chronic pulmonary heart disease, unspecified Bronchitis, not specified as acute or chronic Chronic bronchitis Emphysema Asthma Bronchiectasis Extrinsic allergic alveolitis Chronic airways obstruction, not elsewhere classified Coalworkers' pneumoconiosis Asbestosis Pneumoconiosis due to other silica or silicates Pneumoconiosis due to other inorganic dust Pneumopathy due to inhalation of other dust Pneumoconiosis, unspecified Chronic respiratory conditions due to fumes and vapors Chronic and other pulmonary manifestations due to radiation Respiratory conditions due to other specified external agents
Liver disease	ICD-10	I27.8 I27.9 J40 J41 J42 J43 J44 J45 J47 J60 J61 J62 J63 J64 J65 J66 J67 J68.4 J70.1 J70.3
		Other specified pulmonary heart diseases Pulmonary heart disease, unspecified Bronchitis, not specified as acute or chronic Simple and mucopurulent chronic bronchitis Unspecified chronic bronchitis Emphysema Other chronic obstructive pulmonary disease Asthma Bronchiectasis Coalworker's pneumoconiosis Pneumoconiosis due to asbestos and other mineral fibers Pneumoconiosis due to dust containing silica Pneumoconiosis due to other inorganic dusts Unspecified pneumoconiosis Pneumoconiosis associated with tuberculosis Airway disease due to specific organic dust Hypersensitivity pneumonitis due to organic dust Chronic respiratory conditions due to chemicals, gases, fumes and vapors Chronic and other pulmonary manifestations due to radiation Chronic drug-induced interstitial lung disorders
		070.22 070.23 070.32 070.33 070.44 070.54 070.6 070.9 456.0 456.1 456.2 570 571 572.2 572.3 572.4 572.8 573.3 573.4 573.8 573.9 V427
		Chronic viral hepatitis B with hepatic coma without hepatitis delta Chronic viral hepatitis B with hepatic coma with hepatitis delta Chronic viral hepatitis B without mention of hepatic coma without mention of hepatitis delta Chronic viral hepatitis B without mention of hepatic coma with mention of hepatitis delta Chronic hepatitis C with hepatic coma Chronic hepatitis C without mention of hepatic coma Unspecified viral hepatitis with hepatic coma Unspecified viral hepatitis without mention of hepatic coma Esophageal varices with bleeding Esophageal varices without mention of bleeding Esophageal varices in diseases classified elsewhere, with bleeding Acute and subacute necrosis of liver Chronic liver disease and cirrhosis Hepatic encephalopathy Portal hypertension Hepatorenal syndrome Other sequelae of chronic liver disease Hepatitis, unspecified Hepatic infarction Other specified disorders of liver Unspecified disorder of liver Liver replaced by transplant
		B18 I85 I86.4 K70 K71.1 K71.3 K71.4
		Chronic viral hepatitis Esophageal varices Gastric varices Alcoholic liver disease Toxic liver disease with hepatic necrosis Toxic liver disease with chronic persistent hepatitis Toxic liver disease with chronic lobular hepatitis

(continued on next page)

## Appendix B (continued)

Comorbidity	ICD code	Description
Congestive heart failure	K71.5	Toxic liver disease with chronic active hepatitis
	K71.7	Toxic liver disease with fibrosis and cirrhosis of liver
	K72	Hepatic failure, not elsewhere classified
	K73	Chronic hepatitis, not elsewhere classified
	K74	Fibrosis and cirrhosis of liver
	K76.0	Fatty (change of) liver, not elsewhere classified
	K76.2	Central hemorrhagic necrosis of liver
	K76.3	Infarction of liver
	K76.4	Peliosis hepatis
	K76.5	Hepatic veno-occlusive disease
	K76.6	Portal hypertension
	K76.7	Hepatorenal syndrome
	K76.8	Other specified diseases of liver
	K76.9	Liver disease, unspecified
	Z94.4	Liver transplant status
	ICD-9	Rheumatic heart failure (congestive)
	398.91	Malignant hypertensive heart disease with heart failure
	402.01	Benign hypertensive heart disease with heart failure
	402.11	Unspecified hypertensive heart disease with heart failure
	402.91	Hypertensive heart and chronic kidney disease, malignant, with heart failure and with chronic kidney disease stage I through stage IV, or unspecified
	404.01	Hypertensive heart and chronic kidney disease, malignant, with heart failure and with chronic kidney disease stage V or end stage renal disease
	404.03	Hypertensive heart and chronic kidney disease, benign, with heart failure and with chronic kidney disease stage I through stage IV, or unspecified
	404.11	Hypertensive heart and chronic kidney disease, benign, with heart failure and with chronic kidney disease stage V or end stage renal disease
	404.13	Hypertensive heart and chronic kidney disease, benign, with heart failure and chronic kidney disease stage V or end stage renal disease
	404.91	Hypertensive heart and chronic kidney disease, unspecified, with heart failure and with chronic kidney disease stage I through stage IV, or unspecified
	404.93	Hypertensive heart and chronic kidney disease, unspecified, with heart failure and chronic kidney disease stage V or end stage renal disease
	425.4	Other primary cardiomyopathies
	425.5	Alcoholic cardiomyopathy
	425.8	Cardiomyopathy in other diseases classified elsewhere
	425.9	Secondary cardiomyopathy, unspecified
	428	Heart Failure
	ICD-10	Rheumatic heart failure
	I09.81	Rheumatic heart disease, unspecified
	I09.9	Hypertensive heart disease with heart failure
	I11.0	Hypertensive heart and chronic kidney disease with heart failure and stage 1 through stage 4 chronic kidney disease, or unspecified chronic kidney disease
	I13.0	Hypertensive heart and chronic kidney disease with heart failure and with stage 5 chronic kidney disease, or end stage renal disease
	I13.2	Ischemic cardiomyopathy
	I25.5	Cardiomyopathy
	I42	Cardiomyopathy in diseases classified elsewhere
	I43	Heart failure
	I50	

**Appendix C**

Complication diagnosis.

Complication ICD codes	
ICD Codes	Description
Surgical complication	
PJI	
ICD-9 codes	
711.06	Pyogenic arthritis, lower leg
996.66	Infection and inflammatory reaction due to internal joint prosthesis
996.67	Infection and inflammatory reaction due to other internal orthopaedic device, implant, and graft
ICD-10 codes	
T81.4	Infection following a procedure
T84.5	Infection and inflammatory reaction due to internal joint prosthesis
T84.7	Infection and inflammatory reaction due to other internal orthopaedic prosthetic devices, implants and grafts
SSI	
ICD-9 codes	
682.6	Cellulitis and abscess of leg, except foot
ICD-10 codes	
T81.41	Infection following a procedure, superficial incisional surgical site
T81.49	Infection following a procedure, other surgical site
Wound complications	
ICD-9 codes	
998.3	Disruption of wound
998.83	Nonhealing surgical wound
ICD-10 codes	
T81.3	Disruption of wound
Periprosthetic fracture	
ICD-9 codes	
733.1	Pathologic fracture, unspecified site
733.81	Malunion of fracture
733.82	Nonunion of Fracture
821	Fracture of other and unspecified parts of femur
822	Fracture of patella
823	Fracture of tibia and fibula
996.44	Periprosthetic fracture around prosthetic joint
ICD-10 codes	
M97	Periprosthetic fracture around internal prosthetic joint
S72	Fracture of femur
S72.4	Fracture of lower end of femur
S82.0	Fracture of patella
Hematoma	
ICD-9 codes	
998.12	Hematoma complicating a procedure
ICD-10 codes	
T84.83	Hemorrhage due to internal orthopaedic prosthetic device, implants and grafts
L76.32	Postprocedural hematoma of skin and subcutaneous tissue following other procedure
Medical complications	
Pneumonia/influenza	
ICD-9 codes	
480-488	Pneumonia and influenza
ICD-10 codes	
J09-J18	Influenza and pneumonia
Sepsis	
ICD-9 codes	
038	Septicemia
995.91	Sepsis
ICD-10 codes	
A41.9	Sepsis, unspecified organism
T81.44	Sepsis following a procedure
Cardiac	
ICD-9 codes	
393-398	Chronic rheumatic heart disease
402	Hypertensive heart disease
410-414	Ischemic heart disease
420-429	Other forms of heart disease
ICD-10 codes	
I05-I09	Chronic rheumatic heart diseases
I11	Hypertensive heart disease
I20-I25	Ischemic heart diseases
I30-I5A	Other forms of heart disease
Deep vein thrombosis	
ICD-9 codes	
453	Other venous embolism and thrombosis
ICD-10 codes	
I82	Other venous embolism and thrombosis

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**Appendix C (continued)**

Complication ICD codes	
ICD Codes	Description
Pulmonary embolism	
ICD-9 codes	
415.1	Pulmonary embolism
ICD-10 codes	
I26	Pulmonary embolism

**Appendix D**

Septic revision.

Septic revision = revision TKA CPT codes + ICD codes below		
ICD-10 codes:	Definition	Approximate ICD-9 equivalent
A40.8	Other streptococcal sepsis	038.0
A41.01	Sepsis due to Methicillin-susceptible <i>Staphylococcus aureus</i>	038.11
A41.02	Sepsis due to Methicillin-resistant <i>Staphylococcus aureus</i>	038.12
A41.1	Sepsis due to other specified <i>staphylococcus</i>	038.19
A41.51	Sepsis due to <i>Escherichia coli</i> [E. coli]	038.42
A41.81	Sepsis due to <i>Enterococcus</i>	038.8
A41.9	Sepsis, unspecified organism	038.9, 995.91
A49.01	Methicillin-susceptible <i>Staphylococcus aureus</i> infection, unspecified site	041.11
A49.02	Methicillin-resistant <i>Staphylococcus aureus</i> infection, unspecified site	041.12
B95.4	Other streptococcus as the cause of diseases classified elsewhere	041.09
B95.61	Methicillin-susceptible <i>Staphylococcus aureus</i> infection as the cause of diseases classified elsewhere	041.11
B95.62	Methicillin-resistant <i>Staphylococcus aureus</i> infection as the cause of diseases classified elsewhere	041.12
B95.8	Unspecified <i>staphylococcus</i> as the cause of diseases classified elsewhere	041.10
B96.20	Unspecified <i>Escherichia coli</i> [E. coli] as the cause of diseases classified elsewhere	041.49
L02.415	Cutaneous abscess of right lower limb	682.6
L02.416	Cutaneous abscess of left lower limb	
M00	Pyogenic arthritis	711
M86	Osteomyelitis	730
T81.4	Infection following a procedure	998.59
T83.511	Infection and inflammatory reaction due to indwelling urethral catheter	996.64
T83.518	Infection and inflammatory reaction due to other urinary catheter	
T83.598	Infection and inflammatory reaction due to other prosthetic device, implant and graft in urinary system	996.65
T84.5	Infection and inflammatory reaction due to internal joint prosthesis	996.66
T84.620	Infection and inflammatory reaction due to internal fixation device of right femur	996.67
T84.621	Infection and inflammatory reaction due to internal fixation device of left femur	
T84.69	Infection and inflammatory reaction due to internal fixation device of other site	
T84.7	Infection and inflammatory reaction due to other internal orthopaedic prosthetic devices, implants and grafts	
T85.7	Infection and inflammatory reaction due to other internal prosthetic devices, implants and grafts	996.69
Z47.3	Aftercare following explantation of joint prosthesis	V54.82
Z86.14	Personal history of Methicillin-resistant <i>Staphylococcus aureus</i> infection	V12.04