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Factors associated with persistent postsurgical pain after total knee or hip joint replacement: a systematic review and meta-analysis

Arunangshu Ghoshal^a, Shivam Bhanvadia^b, Som Singh^c, Lauren Yaeger^d, Simon Haroutounian^{e,*}

Abstract

Studies have identified demographic, clinical, psychosocial, and perioperative variables associated with persistent pain after a variety of surgeries. This study aimed to perform a systematic review and meta-analysis of factors associated with persistent pain after total knee replacement (TKR) and total hip replacement (THR) surgeries. To meet the inclusion criteria, studies were required to assess variables before or at the time of surgery, include a persistent postsurgical pain (PPSP) outcome measure at least 2 months after a TKR or THR surgery, and include a statistical analysis of the effect of the risk factor(s) on the outcome measure. Outcomes from studies implementing univariate and multivariable statistical models were analyzed separately. Where possible, data from univariate analyses on the same factors were combined in a meta-analysis. Eighty-one studies involving 171,354 patients were included in the review. Because of the heterogeneity of assessment methods, only 44% of the studies allowed meaningful meta-analysis. In meta-analyses, state anxiety (but not trait anxiety) scores and higher depression scores on the Beck Depression Inventory were associated with an increased risk of PPSP after TKR or THR. This review systematically assessed factors associated with an increased risk of PPSP after TKR or THR. This review systematically assessed factors associated with an increased risk of PPSP after TKR or THR. This review systematically assessed factors associated with an increased risk of PPSP after TKR or THR. This review systematically assessed factors associated with an increased risk of PPSP after TKR or THR. This review systematically assessed factors associated with an increased risk of PPSP after TKR or THR. This review systematically assessed factors associated with an increased risk of PPSP after TKR or THR.

Keywords: Persistent postsurgical pain, Chronic postsurgical pain, Knee replacement, Hip replacement, Systematic review, Meta-analysis

1. Introduction

Persistent postsurgical pain (PPSP) is a common sequel of surgical procedures. The prevalence varies for different surgical procedures and in different studies, ranging from a low of 3% to a high of 81%,⁹⁰ with a reported incidence of about 20% after bone and joint surgeries.³⁹ PPSP substantially impairs quality of life, has negative economic consequences,³⁵ and is a potentially preventable condition.¹⁰⁶ Patients who undergo elective procedures such as total hip or knee replacement, typically expect alleviation of the pain associated with the damaged joint. Although these procedures have a high likelihood of success in relieving pain and restoring function, PPSP is a major factor that negatively affects patient satisfaction with joint replacement

surgery outcomes.⁷² Therefore, understanding factors predisposing patients to an increased risk of persistent pain is important in attempts to prevent PPSP or minimize its occurrence. The identification of patients at high risk for PPSP will allow allocating perioperative resources for PPSP risk mitigation and may signal a need for enhanced postoperative follow-up and early referral to multidisciplinary evaluation. Several studies been have published on perioperative risk factors associated with PPSP,¹²⁰ but it is yet unclear which factors show a consistent association with PPSP.^{47,67}

Several systematic reviews have been published on psychosocial factors in general⁴³ or anxiety and catastrophizing specifically¹⁰⁷ as risk factors for PPSP, and individual studies have reported on factors associated with orthopedic surgeries

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such as total hip replacement (THR)²⁷ and total knee replacement (TKR).¹⁰⁹ An up-to-date synthesis of the existing literature can help improve perioperative planning and care, as well as guide future study designs aimed at better understanding and possible prevention of PPSP.

The objective of the current study was to perform a systematic review and meta-analysis on the reported factors associated with PPSP after total hip joint and knee joint replacement.

2. Methods

The protocol for this systematic review has been preregistered on PROSPERO (CRD42020152146).

2.1. Inclusion and exclusion criteria for studies

The studies were included if they reported PPSP occurrence after a hip joint or knee joint replacement. We included both prospective and retrospective studies that addressed PPSP as pain at the surgical or related site, present at least 2 months postoperatively. The minimum 2-month cutoff was selected as several definitions of PPSP in the past decade have been proposed,⁶⁰ with most suggesting pain duration for more than 2 or 3 months after surgery. Exclusion criteria included (1) fewer than 10 participants per arm, (2) follow-up shorter than 2 months after surgery, (3) participants younger than 18 years, (4) studies on patients with established PPSP that did not report the occurrence of persistent pain after surgery, (5) studies not reporting incidence or prevalence rate of PPSP in the cohort, and (6) abstracts.

A systematic literature search was performed in the following databases: Ovid MEDLINE, Embase, Scopus, Cochrane Central Register of Control Trials, Cumulative Index to Nursing and Allied Health Literature and PsychINFO. The full keyword search strategy can be found in Appendix 1 (available at http://links.lww.com/PR9/A184). The search was initially performed on September 17, 2019, with an update on October 14, 2020 (Appendix 1A, http://links.lww.com/PR9/A184), and subsequently expanded on July 9, 2021 (Appendix 1B, http://links.lww.com/PR9/A184), and updated on February 6, 2022 (Appendix 1C, http://links.lww.com/PR9/A184). The duplicates in the resulting abstracts were removed using EndNote.¹²

The abstracts of the resulting list of articles were screened by 2 authors for eligibility. If the studies were deemed eligible based on abstracts, then the full texts of those articles were retrieved and reviewed independently. The data from eligible articles were extracted into REDCap (Research Electronic Data Capture) and Excel spreadsheets. The data were extracted by 2 authors (A.G. and S.B./S.S.) and compared for inconsistencies. Wherever disagreements arose, a third author (S.H.) was consulted to reach the final decision. Risk of bias (ROB) assessments and data extraction were performed by the 2 reviewers (A.G. and S.B./S.S.) and compared for accuracy.

2.2. Risk of bias assessment

Most validated tools for assessing the ROB in systematic reviews⁴² refer to assessing the methodology of randomized controlled trials. Because this study aimed to analyze factors associated with PPSP (from both prospective and retrospective studies), an alternative method for the ROB systematic assessment was applied based on the methodology used in other systematic reviews focusing on the analysis of predicting factors.^{53,56,116} Potential sources of bias were evaluated and addressed whether (1) the sample adequately represented the population of interest, (2) the data represented the

sample, and (3) the outcome of interest was adequately measured to limit potential bias (Appendix 2, available at http://links.lww.com/ PR9/A184). Other potential serious sources of bias were documented—eg, selective reporting of outcomes, use of nonvalidated scales, or lack or inappropriateness of sample size calculation.

2.3. Protocol for data extraction

To address the different study designs, we developed guiding principles for data extraction across all studies (Appendix 3, available at http://links.lww.com/PR9/A184). In brief, all studies which addressed PPSP at or more than 2 months were included, and the follow-up timeframe at or closest to 6 months was obtained in case of multiple follow-ups. Although the cutoff for postsurgical duration of pain has most commonly been set at 2 or 3 months, some patients have a longer rehabilitation period after total knee or hip joint replacement surgeries.^{38,55} Thus, to remove the confounder, we decided to set the primary outcome timeframe at around 6 months after surgery in studies with multiple follow-ups. If PPSP was categorized based on a severity scale (numerical rating scale [NRS] or Likert scale), the data were dichotomized to a "no pain to mild pain" group vs a "moderatesevere pain" group, to allow 2×2 table formation. In addition, to allow systematic analysis of data, some factors were combined, eg, epidural and spinal anesthesia were merged under neuraxial anesthesia.

2.4. Data synthesis and analysis

The data analyses were performed using RStudio version 1.4.1103.⁹³ For binary factors, 2×2 tables were created, and risk ratios were calculated to be used in generating forest plots. For continuous variables (eg, age and body mass index [BMI]), weighted mean differences were used. Only raw univariate data were used for generating forest plots. Values for weighted mean differences were presented as means with standard deviation (SD). Risk Ratio (RR) or odds ratio was presented as risk or odds, with 95% confidence interval (CI). Meta-regression of multivariable data was not possible because each study accounted for different risk factors. Data on risk factors from multivariable analyses were aggregated in a tabulated form to help make qualitative inferences about common contributing factors.

3. Results

The systematic literature search resulted in a total of 10,851 potential articles. After removing 5,403 duplicate records, a total of 5,448 unique citations were added to the project library. All abstracts of identified studies were reviewed for inclusion criteria, resulting in 231 full-text articles that were considered for full review (see Appendix 1 for full details about literature search strategy and results, available at http://links.lww.com/PR9/A184). These articles were independently evaluated for inclusion by 2 reviewers (AG and SB/SS), and 81 were included in the final analysis (**Fig. 1**).

The studies analyzed included total hip and/or knee joint replacement surgeries. The overall study characteristics are outlined in **Table 1**. The minimum cutoff for determining PPSP in the included studies was 3 months after surgery.⁹⁸ The total number of patients across all total hip and/or knee joint replacement surgery studies (n = 81) was 171,354 with a mean (SD) sample size of 2,115 (14,226), 28 (34.6%) studies involving less than 100 participants. The mean PPSP rate was 16.4%.

Studies investigating PPSP after both total hip and knee joint replacement surgery (n = 11) included 10,399 participants and reported a mean PPSP incidence of 26.7%. Studies after total hip joint replacement surgery alone (n = 16) included 9,086 participants and reported a mean PPSP incidence of 16.5%, and those after total knee joint replacement surgery alone (n = 54) included 151,869 participants and reported a mean PPSP incidence of 15.6%. Overall, 69 of the studies were prospective and 12 were retrospective. A total of 36 studies have provided univariate data to be included in meta-analysis, and 64 studies have reported risk factors in multivariable analyses, from which data were used for qualitative summary (Fig. 1). The studies overall had medium to low ROB (Table 2). The main category to have significant ROB was a "follow-up rate >75% at the first follow-up" (with 14 studies (17%) with high ROB), incomplete characterization of study dropouts (with 65 studies [80%] having unclear or high ROB). Thirty five of the 81 included studies (43%) had high or unclear bias in one of the categories of pain outcome measurement—either in PPSP definition, its method of assessment or classification, or reporting—and 62 of 81 (76.5%) studies reported sources of funding. Although there is not a good way to assess publication bias related to risk prognostic studies,¹⁰⁵ funnel plots to assess asymmetry among most commonly reported risk factors did not suggest important publication bias (Appendix 4, available at http://links. lww.com/PR9/A184).

3.1. Results of meta-analysis

3.1.1. Univariate comparisons

The demographic factors with data from multiple studies that allowed testing their association with an increased risk of PPSP were age, the proportion of women in the total study population,

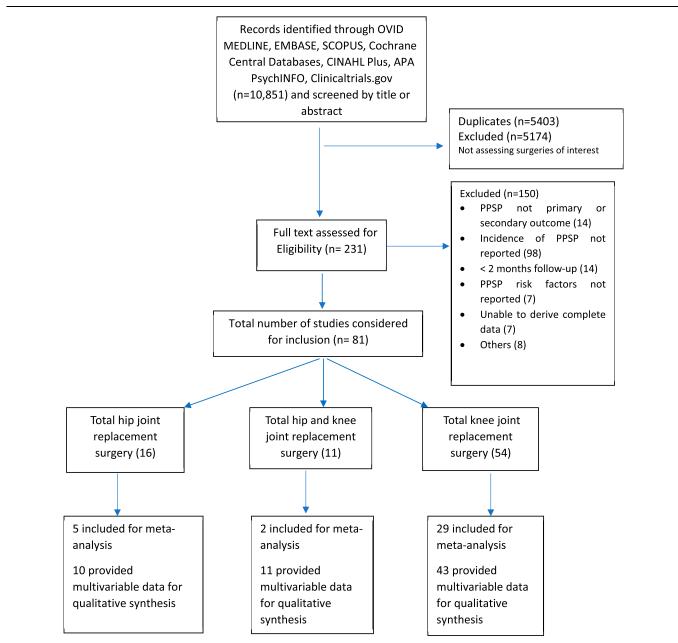


Figure 1. Preferred Reporting Items for Systematic Reviews and Meta-Analyses flow chart.

Table 1

Main characteristics of included studies (n = 81). SN Author-date Type of study Total number of participants Occurrence of PPSP percentage Follow-up Surgery (number) duration* Albayrak 2016 TKR Prospective 274 66.7% (183) 22 mo 1 2 Aso 2021 TKR Prospective 194 10.3% (20) 6 mo 3 Attal 2014 TKR 81 6 mo Prospective 39.5% (32) 4 Aveline 2014 TKR Prospective 75 17.4% (13) 12 mo Baker 2007 TKR 8010 5 Cross-sectional mail/phone 19.8% (1583) 24 mo survey Bischoff-Ferrari THR Retrospective 897 42% (377) 36 mo 6 2004 Bossmann 2017 TKR 47 7 Prospective 20% (9) 6 mo THR 94 8 Bourne 1994 Prospective 27.6% (26) 60 mo 9 Brander 2003 TKR Prospective 116 18.4% (21) 12 mo 10 Briggs 1995 TKR Prospective 65 6% (4) 24 mo 11 Bugada 2017 TKR Prospective 614 21.6% (133) 6 mo 12 Buvanendran 2010 TKR Randomized controlled 228 5.2% (placebo arm: 6 of 115) 6 mo study Buvanendran 2019 TKR 245 14% (34) 13 Prospective 6 mo Buvanendran 2020 50 TKR Prospective 34% (17) 3 mo 14 TKR 220 15 Chen 2021 Prospective 13.6% (30) 6 mo 16 Clarke 2009 THR Randomized controlled 126 (G1: placebo/placebo 26.3% (10); G2: 6 mo GPN/placebo 31.6% (12); G3: placebo/ study GPN 23.7% (9)). 17 Clarke 2010 THR Prospective 82 37.5% (31) 6 mo 18 Dong 2019 TKR Randomized controlled 122 36.06% (44) 3 mo study 19 Dowsey 2015 TKR Retrospective 689 22% (151) 12 mo-60 mo Dumenci 2019 TKR Randomized clinical trial 384 12 mo 20 18% (69) 21 Dürsteler 2021 TKR 146 Prospective 31.5% (46) 3 mo 22 Erlenwein 2017 THR 104 Prospective 17.3% (18) 6 mo 23 Fletcher 2015 THR and 145 12 mo Prospective 27.6% (40) TKR 24 Forsythe 2008 TKR 48 77.1% (37) 24 mo Prospective THR and THR 33.6% (385) 25 George 2021 2411 (1146 THR, 1265 TKR) Retrospective 6 mo TKR TKR 38.9% (492) TKR 26 Giordano 2020 Prospective 136 16.2% (22) 12 mo 27 Grosu 2016 TKR Prospective 114 10% (11) 6 mo 28 Guimaraes-Pereira THR and Prospective 19 50% (9) 3 mo TKR 2016 29 Gungor 2019 TKR Retrospective 578 31.1% (180) 3 mo Kornilov 2017 TKR 79 30 Prospective 22.8% (18) 12 mo 31 Kuchálik 2017 THR Retrospective 72 Group ITM 6% (2) 6 mo Group LIA 0% (0) 32 Kuchálik 2017 THR Randomized controlled 56 (group LIA (n = 29), group FNB Group LIA 0% (0) 6 mo Group FNB 0% (0) study (n = 27))33 Kurien 2018 TKR 50 30.4% (14) Prospective 6 mo TKR 34 Lavand'homme 112 3 mo Prospective 11% (12) 2014 Lee 2019 35 TKR Randomized controlled 24 25% (6) 6 mo study 36 Lindberg 2021 TKR Prospective 202 30.2% (61) 3 mo THR and 45.9% (473) (THR:38% (178) 37 Liu 2012 Cross-sectional mail/phone 1030 12 mo TKR:62% (295)) TKR survey THR 27.2% (200) 38 Lu 2021 Prospective 736 6 mo

71

72

Vila 2020

73 Wang 2014

Von Dincklage 2017

TKR

THR

THR

Prospective

Prospective

study

Randomized controlled

112

105

Table 1 (continued)

Main characteristics of included studies (n = 81). SN **Occurrence of PPSP percentage** Follow-up Author-date Surgery Type of study Total number of participants (number) duration* 39 Martinez 2007 TKR Prospective 20 20% (4) 4 mo 40 Masselin-Dubois TKR Prospective 89 50.6% (45) 3 mo 2013 41 Nazal 2019 THR Retrospective 10 2% (2) 24 mo 42 Neuprez 2020 THR and Prospective 626 (346 THR, 280 TKR) THR 8.99% (31) 60 mo TKR TKR 3.23% (9) 43 Nikolajsen 2006 THR 12-18 mo Cross-sectional mail/phone 1048 28.1% (294) survey 44 Noiseux 2014 TKR Prospective 215 4.6% (10) 6 mo TKR 924 45 Oh 2019 Retrospective 16% (148) 12 mo 46 Pagé 2015 TKR Prospective 108 8.3% (9) 12 mo 47 Pagé 2016 THR Prospective 111 21.6% (24) 6 mo 212 (group CFNB [n = 109] group PCIA 48 Peng 2014 TKR Randomized controlled Group CFNB 33% (36) 6 mo [n = 103]Group PCIA 52.4% (54) study THR and 49 Pereira 2016 Prospective 43 (TKR 22, THR 21) TKR 68.2% (15) 6 mo THR 23.8% (5) TKR 50 Petersen 2015 TKR Cross-sectional mail/phone 305 (215 primary surgery patients, 90 19% (41) primary surgery patients 36 mo 47% (42) revision surgery patients survey revision surgery patients) 51 Petersen 2015 TKR Prospective 78 22% (17) 12 mo 52 Petersen 2017 TKR Prospective 130 19% (25) 12 mo 53 Petersen 2020 TKR Prospective 26 35% (9) 12 mo 54 Pinedo-Villanueva TKR Retrospective 128145 15% (19222) 6 mo 2018 55 Pinto 2013 THR and 92 (THR 48, TKR 44) 41.3% (38) 4–6 mo Prospective TKR THR (13) TKR (25) TKR 56 Puolakka 2010 Cross-sectional mail/phone 562 35% (197) 4-22 mo survey 57 Rao 2020 TKR Randomized controlled 40 47.5% (19) 3-6 mo studv 58 Remérand 2009 THR Randomized controlled 142 (70 placebo, 72 ketamine) 14.8% (21) 6 mo study 15 placebo, 6 ketamine 59 Rice 2018 TKR Prospective 286 21% (60) 6 mo Sakellariou 2015 TKR 272 39.34% (107) 12-16 mo 60 Cross-sectional mail/phone survev 61 Sanders 2009 TKR Randomized controlled 56 (baclofen treatment group n = 27, 27 (baclofen treatment group n = 8, 3 mo control group n = 19) study control group n = 29) 62 Sayers 2016 THR and 560 (283 THR, 277 TKR) 5% (14) THR Randomized controlled 12 mo TKR study 12% (33) TKR 48.4% (88) 63 Sen 2020 TKR Retrospective 182 6.84 ± 4.10 mo 6 mo 64 Sideris 2021 TKR Prospective 162 9.3% (15) 65 Singh 2010 THR 5390 24 mo Cross-sectional mail/phone 8.1% (435) survey TKR 60 mo Skrejborg 2019 Cross-sectional mail/phone 604 18% (107) 66 survey 67 Sugiyama 2018 TKR Retrospective 298 6 mo 33% (97) 68 Thomazeau 2016 TKR Prospective 104 28.8% (74) 6 mo 69 Vaegter 2017 TKR Prospective 14 28.6% (4) 6 mo THR 17.0% (158) Valdes 2012 THR and 70 Prospective 1788 (THR 928, TKR 860) 38 mo TKR

Table 1 (continued)

Main characteristics of included studies (n = 81).

SN	Author-date	Surgery	Type of study	Total number of participants	Occurrence of PPSP percentage (number)	Follow-up duration*
74	W-Dahl 2014	TKR	Retrospective	2123	7.8% (165)	12 mo
75	Wylde 2009	THR and TKR	Cross-sectional mail/phone survey	2391 (1112 THR, 613 TKR)	THR 13% (144) TKR 26% (159)	60–96 mo
76	Wylde 2011	THR and TKR	Cross-sectional mail/phone survey	1294 (662 THR, 632 TKR)	THR 27% (179) TKR 44% (278)	24–48 mo
77	Wylde 2013	TKR	Prospective	51	29% (15)	13 mo
78	Wylde 2020	TKR	Retrospective	3058	11.87% (363)	3 mo
79	Yang 2020	TKR	Randomized controlled study	96	27.08% (26)	3 mo
80	Yao 2019	TKR	Retrospective	694	9.94% (69)	6 mo
81	Zachodnik 2021	THR	Prospective	62	24.19% (15)	12 mo

* Follow-up time points used for data analysis.

CFNB, continuous femoral nerve block, PCIA, patient-controlled intravenous analgesia; GPN, gabapentin; TKR, total knee replacement; THR, total hip replacement; ITM, intrathecal morphine; LIA, local infiltration analgesia; FNB, femoral nerve block.

and BMI, whereas clinical factors included duration of surgery (in minutes), preoperative factors such as presence of anxiety and depression, knee or hip joint pain, or pain elsewhere in the body.

Age (**Fig. 2**) and sex (**Fig. 3**) were not associated with an increased risk of PPSP after TKR or THR (Appendix 5 and Appendix 6 respectively, available at http://links.lww.com/PR9/A184) studies when analyzed separately. However, younger age (SMD -0.18 years [95% CI -0.30 to -0.06]) (**Fig. 4**) and female sex (RR 1.13, 95% CI 1.02-1.24) (Appendix 7, available at http://links.lww.com/PR9/A184) were associated with an increased risk in a small subset of studies reporting on the combined knee and hip replacement surgeries (n = 2).

Body mass index was not associated with a risk of PPSP in TKR studies (**Fig. 5**), but higher BMI was associated with higher PPSP occurrence in the subset of studies reporting both hip and knee replacement (SMD 0.15 kg/m² [95% Cl 0.03–0.26]) (Appendix 8, available at http://links.lww.com/PR9/A184).

Higher state anxiety (Fig. 6) (but not trait anxiety) (Appendix 9, available at http://links.lww.com/PR9/A184) scores (SMD 0.65 [95% CI 0.34–0.97]) (measured on The Spielberger State-Trait Anxiety Inventory) and higher depression scores on the Beck Depression Inventory (SMD 0.68 [95% CI 0.35–1.01]) (Fig. 7) were associated with an increased risk of PPSP in studies reporting PPSP after total knee joint replacement surgery, but not for depression (Appendix 10, available at http://links.lww.com/PR9/A184) or anxiety (Appendix 11, available at http://links.lww.com/PR9/A184) scored in the Hospital Anxiety and Depression Scale.

Higher preoperative NRS pain scores (0-10) were not associated with an increased risk of PPSP in studies reporting PPSP after total knee joint replacement surgery (SMD 0.21 [95% CI -0.01 to 0.44)] (**Fig. 8**) or total knee and hip joint replacement surgeries analyzed together (Appendix 12, available at http://links.lww.com/PR9/A184).

Other associations tested were duration of surgery (mins) (Appendix 13, available at http://links.lww.com/PR9/A184) and pre-existing pain at other sites of the body (Appendix 14, available at http://links.lww.com/PR9/A184), but these did not reach statistical significance.

3.2. Qualitative summary of multivariable analyses

Table 3 presents a qualitative summary of risk factors that were reported in multivariable analyses in the included studies. Factors

are reported as either having a significant association with an increased occurrence of PPSP (red) or no association (yellow). The variables for which the risk factors were adjusted for differ among the studies.

3.3. Demographic factors

In multivariable analysis, an increased risk of PPSP was associated with younger age in $4^{16,57,100,111}$ of 36 studies, female sex in $4^{57,85,99,111}$ of 29 studies, and with BMI in $4^{15,27,99,111}$ of 12 studies that tested these variables as independent risk factors.

3.4. Preoperative factors

Large variability in the incidence of PPSP was found by 2 studies depending on the definition¹⁹ (any pain within the last 4 weeks: NRS >0 vs current pain at the time of the interview NRS >0 vs moderate-to-severe pain at the time of the interview: NRS >3) and assessment instrument used⁷⁶ (Knee Injury and Osteoarthritis Outcome Score pain subscale vs visual analogue scale) to measure pain. Increased risk of PPSP was associated with higher pre-existing pain intensity in 17^{1,13,85,89,97,100,104,27,20,27,28,50,71,74,82,2,58} of 30 studies and with unresolved postoperative pain in the affected knee or hip joint in 12^{4,6,108,15,20,36,41,45,48,61,85,49} of 41 studies that tested these variables as independent risk factors.

3.5. Pre-existing health conditions

Higher PPSP was observed in patients with comorbidities such as hypertension, diabetes, ischemic heart disease, chronic obstructive pulmonary disease, asthma, and neurological illnesses in 7 studies,^{3,30,94,100,69,126,25} and patients with the presence of widespread pain in other parts of the body in 5 studies.^{26,70,100,111,123}

3.6. Psychological factors

Increased risk of PPSP was associated with pre-existing anxiety in $8^{13,15,25,36,65,73,89,114}$ of 20 studies, with pre-existing depression in $7^{13,15,25,36,99,111,122}$ of 20 studies, with preoperative pain catastrophizing in $2^{30,65}$ of 11 studies, and with poorer baseline performance on cognitive flexibility tests in 2 studies^{3,112} that tested these variables as independent risk factors.

Table 2

Risk of bias assessment of the included studies.

SN	Author-Date	Sample	represen	tation of	Data rej	presentati	on of the	Measurement of pain				
		the pop	ulation of	finterest	sample			outcor	nes			
		Target population adequately described	Adequate sampling frame and recruitment	Adequately described inclusion and exclusion criteria	Number of dropouts and reason for dropouts reported	Follow up rate > 75% at first follow up	Study sample not significantly different from dropout sample	PPSP clearly defined	Outcome measure and method adequately limit misclassification bias	No selective reporting of pain outcomes		
1	Albayrak 2016	•	•	•	•	•	•	•	•	•		
2	Aso 2021	•	•	•	•	•	•	•	•	•		
3	Attal 2014	•	•	•	•	•	•	•	•	•		
4	Aveline 2014	•	•	•	•	•	•	•	•	•		
5	Baker 2007	•	•	•	•	•	•	•	•	•		
6	Bischoff-Ferrari 2004	•	•	•	•	•	•	•	•	•		
7	Bossmann 2017	•	•	•	•	•	•	•	•	•		
8	Bourne 1994	•	•	•	•	•	•	•	•	•		
9	Brander 2003	•	•	•	•	•	•	•	•	•		
10	Briggs 1995	•	•	•	•	•	•	•	•	•		
11	Bugada 2017	•	•	•	•	•	•	•	•	•		
12	Buvanendran 2010	•	•	•	•	•	•	•	•	•		
13	Buvanendran 2019	•	•	•	•	•	•	•	•	•		
14	Buvanendran 2020	•	•	•	•	•	•	•	•	•		
15	Chen 2021	•	•	•	•	•	•	•	•	•		
16	Clarke 2009	•	•	•	•	•	•	•	•	•		
17	Clarke 2010	•	•	•	•	•	•	•	•	•		
18	Dong 2019	•	•	•	•	•	•	•	•	•		
19	Dowsey 2015	•	•	•	•	•	•	•	•	•		
20	Dumenci 2019	•	•	•	•	•	•	•	•	•		
21	Dürsteler 2021	•	•	•	•	•	•	•	•	•		
22	Erlenwein 2017	•	•	•	•	•	•	•	•	•		
23	Fletcher 2015	•	•	•	•	•	•	•	•	•		

Table 2 (continued)

Risk of bias assessment of the included studies.

24	Forsythe 2008	•	•	•	•	•	•	•	•	•
25	George 2021	•	•	•	•	•	•	•	•	•
26	Giordano 2020	•	•	•	•	•	•	•	٠	•
27	Grosu 2016	•	٠	•	•	•	•	•	٠	•
28	Guimaraes-Pereira 2016	•	•	•	•	•	•	•	•	•
29	Gungor 2019	•	•	•	•	•	•	•	•	•
30	Kornilov 2017	•	•	•	•	•	•	•	•	•
31	Kuchálik 2017	•	•	•	•	•	•	•	•	•
32	Kuchálik 2017	•	•	•	•	•	•	•	•	•
33	Kurien 2018	•	•	•	•	•	•	•	٠	•
34	Lavand'homme 2014	•	•	•	•	•	•	•	٠	•
35	Lee 2019	•	•	•	•	•	•	•	•	•
36	Lindberg 2021	•	٠	•	•	•	•	•	٠	•
37	Liu 2012	•	•	•	•	•	•	•	•	•
38	Lu 2021	•	•	•	•	•	•	•	٠	•
39	Martinez 2007	•	•	•	•	•	•	•	•	•
40	Masselin-Dubois 2013	•	•	•	•	•	•	•	•	•
41	Nazal 2019	•	•	•	•	•	•	•	•	•
42	Neuprez 2020	•	•	•	•	•	•	•	•	•
43	Nikolajsen 2006	•	•	•	•	•	•	•	٠	•
44	Noiseux 2014	•	•	•	•	•	•	•	•	•
45	Oh 2019	•	•	•	•	•	•	•	•	•
46	Pagé 2015	•	•	•	•	•	•	•	•	•
47	Pagé 2016	•	•	•	•	•	•	•	•	•
48	Peng 2014	•	•	•	•	•	•	•	•	•
49	Pereira 2016	•	•	•	•	•	•	•	•	•
50	Petersen 2015	•	•	•	•	•	•	•	•	•
51	Petersen 2015	•	•	•	•	•	•	•	•	•
52	Petersen 2017	•	•	•	•	•	•	•	•	•
53	Petersen 2020	•	•	•	•	•	•	•	•	•
54	Pinedo-Villanueva 2018	•	•	•	•	•	•	•	•	•
55	Pinto 2013	•	•	•	•	•	•	•	•	•
					I	I				

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Table 2 (continued)

Risk of bias assessment of the included studies.

56	Puolakka 2010	•	٠	•	•	•	•	•	•	•
57	Rao 2020	•	•	•	•	•	•	•	٠	•
58	Remérand 2009	•	٠	•	•	•	•	•	٠	•
59	Rice 2018	•	٠	•	•	•	•	•	٠	•
60	Sakellariou 2015	•	٠	•	•	•	•	•	٠	•
61	Sanders 2009	•	٠	•	•	•	•	•	•	•
62	Sayers 2016	•	٠	•	•	•	•	•	٠	•
63	Sen 2020	•	•	•	•	•	•	•	•	•
64	Sideris 2021	•	•	•	•	•	•	•	•	•
65	Singh 2010	•	•	•	•	•	•	•	•	•
66	Skrejborg 2019	•	•	•	•	•	•	•	•	•
67	Sugiyama 2018	•	•	•	•	•	•	•	•	•
68	Thomazeau 2016	•	•	•	•	•	•	•	•	•
69	Vaegter 2017	•	•	•	•	•	•	•	•	•
70	Valdes 2012	•	•	•	•	•	•	•	•	•
71	Vila 2020	•	•	•	•	•	•	•	•	•
72	Von Dincklage 2017	•	•	•	•	•	•	•	•	•
73	Wang 2014	•	•	•	•	•	•	•	•	•
74	W-Dahl 2014	•	٠	•	•	•	•	•	•	•
75	Wylde 2009	•	٠	•	•	•	•	•	•	•
76	Wylde 2011	•	٠	•	•	•	•	•	•	•
77	Wylde 2013	•	٠	•	•	•	•	•	•	•
78	Wylde 2020	•	٠	•	•	•	•	•	٠	•
79	Yang 2020	•	٠	•	•	•	•	•	•	•
80	Yao 2019	•	٠	•	•	•	•	•	•	•
81	Zachodnik 2021	•	•	•	•	•	•	•	•	•

• Low risk of bias • Unclear risk of bias • High risk of bias

Study	Total	Expo Mean			o Expo Mean		Standardized Mean Difference	SMD	95%-CI	Weight
Study	Total	wean	50	Total	wean	30	Difference	SIND	33 /0-01	weight
Albayrak I 2016	183	66.8	7.3	91	66.8	6.7	<u> </u>	0.00	[-0.25; 0.25]	6.6%
Attal N 2014	32	69.2	9.1	49	68.0	8.8		0.13	[-0.31; 0.58]	4.0%
Aveline C 2014	12	72.0	6.0	57	71.0	8.0		0.13	[-0.49; 0.75]	2.6%
Dong J 2019	58	68.2	17.1	64	69.5	13.7		-0.08	[-0.44; 0.27]	5.0%
Forsythe M 2008	37	69.6	7.9	11	69.2	7.6		0.05	[-0.62; 0.72]	2.3%
Gungor S 2019	180	64.9	9.8	398	66.6	9.0	- +	-0.18	[-0.36; -0.01]	7.8%
Kornilov 2017	18	64.0	8.0	61	63.0	8.0		0.12	[-0.40; 0.65]	3.2%
Kurien T 2018	15	65.1	8.9	35	66.9	8.0		-0.21	[-0.82; 0.39]	2.7%
Lavand'homme PM 2014	65	67.0	10.0	47	69.0	9.0		-0.21	[-0.58; 0.17]	4.8%
Lindberg 2021	139	68.9	8.6	63	66.8	10.3	÷	0.23	[-0.07; 0.53]	5.8%
Masselin–Dubois A 2013	45	69.4	7.9	44	66.5	9.9		0.32	[-0.10; 0.74]	4.3%
Oh TK 2019	90	73.6	6.1	834	71.5	6.1		0.34	[0.13; 0.56]	7.1%
Petersen 2020	9	64.0	4.0	17	70.0	2.0 <			[-3.06; -1.04]	1.2%
Rao PB 2020	25	58.5	6.0	25	58.4	5.6		0.01	[-0.54; 0.57]	3.0%
Rice DA 2018	60	67.0	8.0	231	70.0	10.0		-0.31	[-0.60; -0.03]	6.0%
Sideris 2021	15	65.6	4.1	147	67.2	8.4		-0.20	[-0.73; 0.34]	3.2%
Skrejborg P 2019	107	67.0	9.0	245	68.0	10.0		-0.10	[-0.33; 0.12]	7.0%
Sugiyama Y 2018	97	71.0	9.0	201	73.0			-0.19	[-0.43; 0.05]	6.7%
Thomazeau J 2016	30	66.5	8.0	74	69.8	9.2		-0.37	[-0.80; 0.06]	4.2%
Yang R 2020	50	65.2	7.2	47	62.7	8.1		0.33	[-0.08; 0.73]	4.5%
Yao Y 2019	266	66.9	8.5	470	66.6	8.5		0.04	[-0.12; 0.19]	8.2%
Random effects model Heterogeneity: $I^2 = 60\%$, τ^2		77, p <	0.01	3211		-1	0		[-0.15; 0.08]	100.0%

Figure 2. Forest plot for the association between age and PPSP in total knee joint replacement surgeries analyzed together. *Higher/positive SMD represents the difference in age (in years) between patients who developed PPSP vs those who did not, the horizontal box and whisker plots are the group SMDs ±95% confidence intervals, and the blue diamond is the combined SMD of groups.

Study	Exp Events	osure		posure Total	Risk Ratio	RR	95%-CI	Woigh
Study	Events	Total	Evenus	Total		nn	95 /o-Ci	weigi
Albayrak I 2016	138	183	79	91		0.87	[0.77; 0.97]	5.5%
Attal N 2014	18	26	27	43		1.10	[0.78; 1.56]	2.1%
Aveline C 2014	9	12	35	57		1.22	[0.83; 1.80]	1.8%
Buvanendran A 2019	27	34	133	211		1.26	[1.03; 1.54]	3.9%
Dong J 2019	38	58	41	64		1.02	[0.79; 1.33]	3.0%
Gungor S 2019	116	180	246	398		1.04	[0.91; 1.19]	5.2%
Kim MS 2019	18	20	45	48	-	0.96	[0.82; 1.13]	4.6%
Kornilov 2017	16	18	59	61		0.92	[0.78; 1.09]	4.5%
Kurien T 2018	10	15	20	35		1.17	[0.74; 1.85]	1.4%
Lavand'homme PM 2014	32	53	31	47		0.92	[0.68; 1.24]	2.5%
Lindberg 2021	94	139	138	202	+	0.99	[0.85; 1.15]	4.9%
Masselin-Dubois A 2013	31	45	26	44		1.17	[0.85; 1.60]	2.4%
Oh TK 2019	30	90	825	834		0.34	[0.25; 0.45]	2.6%
Petersen KK 2015	52	83	142	305		1.35	[1.10; 1.65]	3.8%
Petersen KK 2015	7	17	39	61		0.64	[0.35; 1.17]	0.9%
Petersen 2020	5	9	9	17		1.05	[0.50; 2.19]	0.6%
Pinedo-Villanueva 2018	10343	18522	57771	107542		1.04	[1.03; 1.05]	7.0%
Puolakka P 2010	105	130	268	296		0.89	[0.81; 0.98]	6.0%
Rao PB 2020	17	25	16	25		1.06	[0.71; 1.58]	1.7%
Rice DA 2018	22	45	128	243		0.93	[0.67; 1.28]	2.3%
Sideris 2021	8	15	83	147		0.94	[0.58; 1.55]	1.2%
Skrejborg P 2019	71	107	143	245	+	1.14	[0.96; 1.35]	4.4%
Skrejborg P 2020	54	94	49	91	<u> </u>	1.07	[0.82; 1.38]	3.0%
Sugiyama Y 2018	78	97	153	201		1.06	[0.93; 1.20]	5.3%
Thomazeau J 2016	25	30	50	74		1.23	[0.99; 1.54]	3.5%
Vila M 2020	33	47	33	65		1.38	[1.02; 1.87]	2.5%
Wylde V 2011	53	93	104	183		1.00	[0.81; 1.25]	3.6%
Yang R 2020	38	50	41	47		0.87	[0.72; 1.05]	4.1%
Yao Y 2019	195	266	328	470	<u>+</u>	1.05	[0.96; 1.15]	5.9%
Random effects model		20503		112147	•	1.00	[0.95; 1.07]	100.0%
Heterogeneity: $I^2 = 73\%$, τ^2	= 0.0132	p < 0.0	1	1				

Figure 3. Forest plot for the association between the proportion of women in the study population and PPSP in total knee joint replacement surgeries. *Risk ratio (RR) >1 represents higher, RR <1 represents lesser, while RR = 1 represents a similar occurrence of PPSP in women compared with the study population, the horizontal box and whisker plots are the group RRs \pm 95% confidence intervals, and the blue diamond is the combined RR of groups.

Study	Total	Expo Mean		o Expo Mean		Standardized Mean Difference in age (Years)	SMD	95%-CI	Weight
Liu SS 2012 Pinto PR 2013	473 38	65.0 63.3	 	67.0 64.5				0.30; –0.06] •0.57; 0.26]	92.0% 8.0%
Random effects model Heterogeneity: $I^2 = 0\%$, τ^2		= 0.89	611		ا _	1 0	1 -0.18 [-(0.30; –0.06]	100.0%

Figure 4. Forest plot for the association between age and PPSP in total knee and hip joint replacement surgeries analyzed together. *Higher/positive SMD represents the difference in age (in years) between patients who developed PPSP vs those who did not, the horizontal box and whisker plots are the group SMDs \pm 95% confidence intervals, and the blue diamond is the combined SMD of groups.

3.7. Psychophysical assessments

Low preoperative pressure pain detection threshold was associated with PPSP in 1 study,⁷⁷ whereas temporal summation of pain (TSP) or conditioned pain modulation (CPM) alone showed no associations, but the association with CPM was seen in male subgroup patients.¹⁰ Increased odds of PPSP were also observed with heat hyperalgesia,⁶² preoperatively facilitated TSP,⁴⁵ expected pain and TSP,⁸⁹ and with preoperative widespread pain sensitization measured using pressure algometry,¹²⁵ whereas hypoalgesia after cold pressor stimulation and aerobic exercise assessed preoperatively by cuff algometry was associated with less PPSP.¹¹⁰

3.8. Perioperative interventions

There were no sufficient data to combine perioperative interventions in the meta-analyses. Data from 1 or 2 articles were available on each intervention; therefore, only a qualitative summary is provided below.

In the 4 studies which reported on regional anesthesia in TKR, a reduced incidence of PPSP was reported with continuous femoral nerve block analgesia with standardized rehabilitation therapy in 1 study,⁷⁶ whereas analgesic benefit was noticed only for up to 1 month after surgery in 1 study¹⁵ or even for a shorter immediate postoperative period but not 3 or 6 months postsurgery in the other 2.^{51,128} There was reduced pain at 3 months after TKR with intrathecal baclofen used as an adjuvant to spinal anaesthesia,⁹⁵ with sevoflurane anaesthesia,¹²⁷ and with preventive epidural analgesia.⁸⁶ Perioperative pregabalin

administration reduced the incidence of chronic neuropathic pain in a single study.¹⁷

After THR, a single 600-mg dose of gabapentin given preoperatively or postoperatively neither reduced morphine consumption nor pain scores in the hospital or after 6 months within the context of spinal anesthesia and a robust multimodal analgesia regimen.²¹ There was no difference between local infiltration analgesia and intrathecal morphine or femoral nerve block,⁵¹ and no effect of a single spinal dose of ketorolac¹¹⁵ or perioperative ketamine⁸⁸ on PPSP prevention. There was no effect on PPSP from preoperative analgesia and neuraxial anesthesia in a study analyzing THR and TKR together.³⁶

4. Discussion

This systematic review and meta-analysis focused on identifying reported factors associated with PPSP risk across a group of hip joint and knee joint replacement surgeries. Overall, 81 studies met the eligibility criteria and were included for further analyses. The studies selected based on inclusion criteria had mostly low or unclear ROB for appropriate representation of the surgical population tested, study design, likelihood of publication bias, and the measurement of pain outcomes. However, based on ROB assessment, important sources of bias and heterogeneity were low follow-up rate and inconsistent characterization of participants who dropped out of studies, as well as inconsistent definition of PPSP, its assessment methods and reporting. The overall mean PPSP rate was 16.4% (16.5% after THR and 15.6% after TKR), which was similar to the previous literature review (9% or more after THR and about 20% after TKR).

Study	Total	Exposure Mean SD		Exposure Mean SD	Standardized Mean Difference in BMI units	SMD	95%-CI	Weigh
Albayrak I 2016	183	32.1 4.6	91	32.6 4.8		-0.11	[-0.36; 0.14]	11.29
Aveline C 2014	12	28.7 5.6	57	26.3 4.7		→ 0.49	[-0.14; 1.12]	1.99
Lavand'homme PM 2014	65	29.0 5.0	47	28.0 5.0		0.20	[-0.18; 0.57]	5.29
Lindberg 2021	139	29.2 4.9	63	29.3 4.6		-0.02	[-0.32; 0.28]	8.19
Oh TK 2019	90	26.9 3.6	834	27.0 3.3		-0.03	[-0.25; 0.19]	14.69
Petersen 2020 51	9	29.0 2.0	17	30.0 1.0 ←		-0.68	[-1.52; 0.15]	1.19
Sideris 2021	15	33.1 5.2	147	31.2 6.9		0.28	[-0.25; 0.81]	2.69
Skrejborg P 2019	107	30.7 5.2	245	29.9 5.0		0.16	[-0.07; 0.39]	13.59
Thomazeau J 2016	30	31.7 5.5	74	30.3 5.4		0.26	[-0.17; 0.68]	4.19
Vila M 2020	47	33.0 7.0	65	32.0 5.0		0.17	[-0.21; 0.54]	5.2%
Yang R 2020	50	27.0 5.4	47	25.8 3.5		0.27	[-0.13; 0.67]	4.69
Yao Y 2019	266	25.7 3.9	470	25.5 4.2		0.05	[-0.10; 0.20]	27.99
Random effects model	1013		2157			0.07	[-0.02; 0.15]	100.09

Figure 5. Forest plot for the association between body mass index (BMI) in kg/m² and PPSP in total knee joint replacement surgeries analyzed together. *Higher/ positive SMD represents the difference in BMI units between patients who developed PPSP vs those who did not, the horizontal box and whisker plots are the group SMDs ±95% confidence intervals, and the blue diamond is the combined SMD of groups.

SMD 95%–CI	95%–Cl We
0.51 [0.06; 0.96]	[0.06; 0.96] 47
0.78 [0.35; 1.21]	[0.35; 1.21] 52
0.65 [0.34; 0.97]	[0.34; 0.97] 100
0.00	

Figure 6. Forest plot for the association between state anxiety and PPSP in total knee joint replacement surgeries analyzed together. *Higher/positive SMD represents the difference in the state anxiety score between patients who developed PPSP vs those who did not, the horizontal box and whisker plots are the group SMDs \pm 95% confidence intervals, and the blue diamond is the combined SMD of groups.

4.1. Knee joint replacement surgeries

The results of the meta-analyses in knee joint replacement surgeries from 54 studies demonstrated an increased risk of PPSP in individuals with higher state anxiety (but not trait anxiety) scores and higher depression scores (in Beck Depression Inventory). Literature suggests that a bidirectional relationship exists between pain and anxiety or depression, and observations from functional imaging studies suggest that this bidirectional relationship is due in part to shared neural circuitry, particularly related to mechanisms of emotional regulation of pain.¹³⁰

The qualitative summary of multivariable analysis from 43 studies pointed out several possible factors associated with PPSP but with limited data and evidence. Presurgical pain intensity was the only factor where more studies reported an independent risk with PPSP compared with studies that did not find such an association. Although factors such as younger age, female sex, higher BMI, high catastrophizing, more severe acute postoperative pain, pre-existing depression, or anxiety were independently associated with PPSP in some studies, there were generally more studies that did not find such independent association. Certain factors were demonstrated in single studies to increase the risk of PPSP. Among these were being a widower; being a housewife; not having higher education; working at a job that requires physical effort;¹ presurgical restricted walking distance; pre-existing knee pain; neuropathic pain;⁸¹ pre-existing chronic pain states;¹¹⁹ poorer cognitive flexibility; comorbidities such as diabetes mellitus, cancer, and fibromyalgia;³³ revision surgery compared with primary TKA;79 mechanical complication of prosthesis;48 and painful postoperative period.³⁴ Given the paucity of data, future research will be needed to test these associations further.

Interventions such as preoperative exercise and education were suggested to have low-quality to moderate-quality evidence in a recent review,²⁴ but we did not find any evidence in the included studies to quantify these measures. Perioperative pregabalin administration reduced the incidence of chronic neuropathic pain

in 1 study¹⁷ but has been refuted in a meta-analysis.⁶³ Continuous regional anesthesia provided analgesic benefit for up to 1 month after surgery but did not influence PPSP outcomes at 6 months,¹⁵ and single-injection femoral nerve block did not affect PPSP.¹²⁸ Because these findings are based on a limited number of studies, it is impossible to draw robust conclusions on the effectiveness of these perioperative interventions to prevent PPSP.

4.2. Hip joint replacement surgeries

The results of the meta-analyses in hip joint replacement surgeries only from 5 studies could not demonstrate any statistically significant results for factors consistently associated with an increased risk of PPSP. The qualitative summary from multivariable analysis results from 10 studies pointed out several possible factors associated with PPSP; however, there is insufficient evidence to conclude these contributing factors because of a small number of studies reporting them. Young age,⁹¹ female sex,¹⁰³ higher time spent with pain in the perioperative period, presurgical anxiety or depression, higher BMI, poor Harris Hip Score,¹¹ presence of loose beads from the prosthesis, pain at other sites of the body, preoperative pain disability, higher cumulative opioid consumption,⁵² and high intraoperative nociception were the factors reported to be associated with an increased risk of PPSP in some, but not other studies. The current data on preoperative education of patients undergoing THR are unclear, 32,46,68 and we did not find any conclusive evidence on PPSP prevention with this approach. Perioperative ketamine was associated with a reduction in the risk of PPSP in 1 study⁸⁷ but has been refuted in a meta-analysis.⁴⁹ A single perioperative dose of gabapentin,²¹ spinal ketorolac, local infiltration analgesia, and femoral nerve block did not show a protective effect on the presence of PPSP at 6 months after THR. These findings are based on limited data; therefore, no conclusion can be drawn of effective strategies for preventing PPSP after THR.

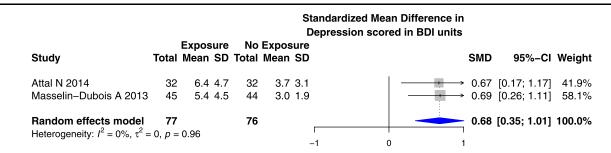


Figure 7. Forest plot for the association between depression scored in the Beck Depression Inventory (0–39) and PPSP in total knee joint replacement surgeries was analyzed together. *Higher/positive SMD represents the difference in depression scored in the Beck Depression Inventory (score range 0–39) units between patients who developed PPSP vs those who did not, the horizontal box and whisker plots are the group SMDs ±95% confidence intervals, and the blue diamond is the combined SMD of groups.

		F	Na	р	Standardized Mean Difference reoperative NRS pain score ur			
Study		Exposure Mean SD		Exposure Mean SD		SMD	95%–Cl	Weight
Albayrak I 2016	183	8.1 0.5	91	7.9 1.2	<u> </u>	0.25	[0.00; 0.50]	9.1%
Attal 2014	32	5.3 1.9	49	4.3 2.3		0.46	[0.01; 0.91]	7.3%
Forsythe M 2008	37	7.0 1.9	11	6.1 2.0		0.46	[-0.22; 1.14]	5.3%
Giordano 2020	22	6.3 1.8	114	6.6 1.8		-0.17	[-0.62; 0.29]	7.2%
Gungor 2019	180	7.9 1.9	398	6.8 2.2		0.52	[0.34; 0.70]	9.6%
Gungor S 2019	180	7.9 1.9	398	6.8 2.2		0.52	[0.34; 0.70]	9.6%
Kurien 2018	15	6.5 2.1	35	4.7 2.3	+ >	0.79	[0.16; 1.42]	5.7%
Lindberg 2021	139	5.3 2.1	63	6.1 1.9		-0.39	[-0.69; -0.09]	8.6%
Masselin–Dubois A 2013	45	5.0 2.1	44	4.4 2.2		0.28	[-0.14; 0.69]	7.6%
Petersen 2017	25	6.5 0.3	105	6.7 0.2	< ·	-0.89	[-1.34; -0.44]	7.3%
Rice DA 2018	60	5.5 1.8	231	4.4 2.0		0.56	[0.27; 0.85]	8.8%
Sideris 2021	15	2.5 3.4	147	2.4 2.5		0.04	[-0.49; 0.57]	6.5%
Thomazeau 2016	30	4.8 3.3	74	3.7 2.8		0.37	[-0.06; 0.80]	7.5%
Random effects model Heterogeneity: $I^2 = 83\%$, τ^2	963 = 0.130	07, <i>p</i> < 0.01	1760	-	1 0 '	0.21	[-0.01; 0.44]	100.0%

Figure 8. Forest plot for the association between preoperative numerical rating scale pain score (0–10) and PPSP in total knee joint replacement surgeries analyzed together. *Higher/positive SMD represents the difference in preoperative numerical rating scale pain score (0–10) units between patients who developed PPSP vs those who did not, the horizontal box and whisker plots are the group SMDs ±95% confidence intervals, and the blue diamond is the combined SMD of groups.

4.3. Hip and knee joint replacement surgeries

The results of the meta-analyses in studies that combined patients undergoing hip and knee joint replacement surgeries demonstrated an increased risk of PPSP in women and patients with higher BMI, but the number of studies is likely too small to draw any meaningful conclusions. The qualitative summary of multivariable analyses from 11 studies pointed out several possible factors associated with PPSP. Functional severity of joint pathology, higher time spent with pain in the preoperative period, presurgical anxiety or depression, and presence of pain in other areas of the body were associated with higher instances of PPSP, whereas preoperative analgesia and neuraxial anesthesia were associated with less PPSP in 1 study. It has been reported that younger patients are more likely to develop PPSP after orthopedic surgery,⁸ but this finding was not consistent across studies.⁷⁵ Mechanisms explaining the increased incidence of chronic pain in younger patients are unknown but may relate to a reduction in peripheral nociceptive function and density with increased age.¹²⁹ It has been reported that women experience higher pain intensity at a lower level of inflammation after knee surgery compared with men, which has been attributed to central mechanisms of pain perception.¹⁰¹ Data from the American College of Surgeons National Surgical Quality Improvement Program database has shown that the degree of obesity (high BMI) relates to the of risk of postoperative complications in hip and knee arthroplasty,⁹² which is consistent with our findings from the qualitative summary. Several factors, such as employment status, illiteracy, history of hypertension or diabetes, and previous knee or hip replacement surgeries have been reported to be associated with higher PPSP in the literature but did not show association in this combined analysis.83,84

4.4. Assessment of chronic pain

Chronic pain as a subjective lived, and sensory experience presents particular challenges to assessment.⁷ This review has identified the following tools used for assessing chronic pain: Oxford Knee Score (joint-specific questionnaire assessing pain severity and interference), Western Ontario and McMaster

Universities Osteoarthritis Index, Knee Injury and Osteoarthritis Outcome Score (disease-specific questionnaire assessing pain severity), the Brief Pain Inventory (disease-agnostic questionnaire with a pain severity subscale [4 items] and a pain interference subscale [7 items]), Short-Form McGill Pain questionnaire (disease-agnostic questionnaire assessing sensory and affective qualities of pain), Pain visual analogue scale (unidimensional assessment of pain severity), PainDETECT (a neuropathic pain guestionnaire), and Douleur Neuropathique 4 (DN4, a neuropathic pain questionnaire). Because these tools are selected based on study objectives and each measure a somewhat different dimension of pain severity, characteristics, and interference, direct comparison among studies is challenging. It is increasingly recognized that the accurate assessment of patient-reported outcome measures (PROMs) in orthopedic surgery trials and registries is important and ideally should include metrics of quality of life, functioning, disability, and patient satisfaction.9,66 In both clinical and research settings, the approach to assessing chronic pain after THR or TKR needs to be in-depth and multidimensional to understand the characteristics and impact of this pain. Unfortunately, the heterogeneity in PROM assessment in the included studies did not allow to robustly test the associations between PPSP and PROMs. We could not identify risk factors that are associated with diminished pain-related quality of life after knee and hip joint replacement surgeries. Only 1 study showed diminished quality of life in patients with PPSP compared with those without,³⁶ while 2 other studies focused on quality of life measures in the context of perioperative anesthesia selection¹²⁷ and general longitudinal outcomes of joint replacement surgery.⁶⁹

4.5. Quantitative sensory testing modalities

Quantitative sensory testing (QST) is a psychophysical approach used to quantify somatosensory function in response to controlled stimuli.⁵ The idea of using QST in the preoperative setting to determine an individual's response to painful stimuli is appealing as a possible predictor of a response to a painful surgical procedure. In this review, heat pain threshold such as hyperalgesia was associated with perioperative pain after TKR⁶²

Table 3

Qualitative summary of risk factors for PPSP reported in multivariable analyses in included studies.

	Author, year	Type of surgery	Younger age	Female sex	Pre-surgical pain	Higher BMI	Preop depression	Preop anxiety	Pain catastrophizing	Higher postop pain scores	Other risk factors, including quantitative sensory testing (QST)
1	Albayrak 2016	TKR									Increased risk in widowers, having a primary school education level or under, housewives, people with jobs that require physical effort, those suffering from pre-surgical restricted walking distance
2	Aso 2021	TKR									Preoperative severe pain and postoperative coronal malalignment were independent risk factors of CPSP after TKA
3	Attal 2014	TKR									Increased risk with premorbid limited cognitive flexibility and memory capacities
4	Aveline 2014	TKR									
5	Baker 2007	TKR									Oxford knee score was used
6	Bischoff-Ferrari 2004	THR									Pain in the operated hip was strongly associated with poor functional status 3 years after THR
7	Bossmann 2017	TKR									Dysregulation of the nociceptive and autonomic nervous systems may be predictive of pain severity 6 months after primary TKR
8	Bourne 1994	THR									Increased risk with poor Harris Hip Score, presence of loose beads which were shed from the prosthesis
9	Brander 2003	TKR									
10	Briggs 1995	TKR									Use of the Tricon hybrid system has resulted in 94% of all patients having a good or excellent result an average of 24 months postoperatively
11	Bugada 2017	TKR									Continuous regional anesthesia provided analgesic benefit for up to 1 month after surgery but did not influence PPSP at 6 months. Better pain control at 1 month was associated with reduced PPSP. Patients with higher expectations from surgery, enhanced basal inflammation, lower ASA physical status and a pessimistic outlook are more prone to develop PPSP
12	Buvanendran 2010	TKR									Perioperative pregabalin administration reduced the incidence of chronic neuropathic pain
13	Buvanendran 2019	TKR									
14	Buvanendran 2020	TKR									Cognitive-behavioral therapy interventions delivered prior to surgery in person or via telehealth can reduced pain catastrophizing scores; however, this reduction did not lead to improved 3-month pain outcomes

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Table 3 (continued)

Qualitative summary of risk factors for PPSP reported in multivariable analyses in included studies.

15	Chen 2021	TKR			Preoperative angiotensin II type 2 receptor is a predictor for developing chronic post-surgical pain after total knee arthroplasty surgery
16	Clarke 2009	THR			A single 600mg dose of gabapentin given pre- operatively or post-operatively did not reduce morphine consumption or pain scores in hospital or at 6 months after THR within the context of spinal anesthesia and multimodal analgesia
17	Clarke 2010	THR			Maximum movement-evoked acute pain did not predict the presence of CPSP at 6 months
18	Dong 2019	TKR			Non-tourniquet combined with controlled hypotension technique can alleviate chronic pain
19	Dowsey 2015	TKR			
20	Dumenci 2019	TKR			
21	Dürsteler 2021	TKR			Preoperative sensory testing predicts poor pain outcome after TKR
22	Erlenwein 2017	THR			
23	Fletcher 2015	THR and TKR			Functional impairment was associated with CPSP severity, percentage of time in severe pain on postoperative day one
24	Forsythe 2008	TKR			Number of comorbidities per patient predicted the presence of pain postoperatively
25	George 2021	THR and TKR			Non-white race and TKR were the only two variables associated with chronic pain outcomes
26	Giordano 2020	TKR			Increased risk with dysregulation of circulating microRNAs hsa-miR-146a-5p, -145-5p, and -130 b-3p
27	Grosu 2016	TKR			The trajectories of the different types of pain after TKR show their non-linear evolution.
28	Guimaraes- Pereira 2016	THR and TKR			
29	Gungor 2019	TKR			Increased risk in African Americans (vs whites), adductor canal saphenous nerve (vs femoral nerve) blocks
30	Kornilov 2017	TKR			
31	Kuchálik 2017 (Retrospective)	THR			No difference between local infiltration analgesia and intrathecal morphine
32	Kuchálik 2017 (Prospective)	THR			No difference between local infiltration analgesia and femoral nerve block
33	Kurien 2018	TKR			Patients with knee osteoarthritis with neuropathic pain-like symptoms identified using the PainDETECT questionnaire are most at risk
34	Lavand'homme 2014	TKR			
35	Lee 2019	TKR			Decreased risk with prerecorded hypnotic intervention
36	Lindberg 2021	TKR			
37	Liu 2012	THR and TKR			Increased risk with knee versus hip replacement, presence of pain in other areas of the body
38	Lu 2021	THR			

Table 3 (continued)

Qualitative summary of risk factors for PPSP reported in multivariable analyses in included studies.

			 r				
39	Martinez 2007	TKR					Heat hyperalgesia was the predominant QST symptom associated with perioperative pain after total knee arthroplasty, and was predictive of postoperative morphine consumption
40	Masselin- Dubois 2013	TKR					
41	Nazal 2019	THR					Orthopedic surgeons performing THA should consider minimizing disruption and/or ensuring repair of the anterior capsule to decrease this potential cause of iliopsoas tendinopathy
42	Neuprez 2020	THR and TKR					The determinants of "good outcome" for THR were a higher preoperative disability score, lack of comorbidities, and the absence of postoperative complications. In patients undergoing TKR, the degree of preoperative stiffness, level of education, degree of radiological severity, lack of comorbidities, place of discharge, and length of hospital stay were identified as predictors for good outcome
43	Nikolajsen 2006	THR					Increased risk with presence of pain at other sites of the body
44	Noiseux 2014	TKR					
45	Oh 2019	TKR					Decreased risk with perioperative magnesium sulfate administration
46	Pagé 2015	TKR					
47	Pagé 2016	THR					Increased risk with preoperative pain disability and anxiety as well as cumulative morphine consumption
48	Peng 2014	TKR					Decreased risk with continuous femoral nerve block analgesia 24 h following surgery
49	Pereira 2016	THR and TKR					Decreased risk with preoperative analgesia and neuraxial anesthesia
50	Petersen 2015	TKR					Increased risk with revision surgery compared with patients after primary TKA
51	Petersen 2015	TKR					Low preoperative pressure pain detection threshold (PDT) was associated with a less postoperative pain relief, whereas temporal summation of pain (TSP) or conditioned pain modulation (CPM) alone showed no associations with postoperative pain relief
52	Petersen 2017	TKR					Increased risk with preoperatively facilitated TSP
53	Petersen 2020	TKR					CE-MRI and DCE-MRI synovitis was associated with higher pain intensity 12 months after TKA
54	Pinedo- Villanueva 2018	TKR					Decreased risk with IT baclofen used as an adjuvant to spinal anesthesia
55	Pinto 2013	THR and TKR					Increased risk with chronic timeline perception of the disease, postsurgical anxiety
56	Puolakka 2010	TKR					
57	Rao 2020	TKR					Decreased risk with preventive epidural analgesia
58	Remérand 2009	THR					Decreased risk with peri-operative Ketamine

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Table 3 (continued)

Qualitative summary of risk factors for PPSP reported in multivariable analyses in included studies.

59	Rice 2018	TKR				Increased risk with expected pain, TSP
	Sakellariou					Increased risk with length of the operative procedure, medical history of diabetes, presence
60	2015	TKR				of preoperative flexion contracture, and patellofemoral joint overstuffing
61	Sanders 2009	TKR				Decreased risk with IT baclofen used as an adjuvant to spinal anesthesia
62	Sayers 2016	THR and TKR				
63	Sen 2020	TKR				Increased risk with neuropathic mechanisms and central sensitization
64	Sideris 2021	TKR				Patient-specific biologic response to surgery may influence longer-term clinical outcomes after TKA.
65	Singh 2010	THR				
66	Skrejborg 2019	TKR				Increased risk in fibromyalgia, chronic pain in body parts other than the knee, previous diagnosis of cancer, knee instability
67	Sugiyama 2018	TKR				
68	Thomazeau 2016	TKR				Increased risk with lower education and a lack of physical activity in adulthood. An association of borderline statistical significance was found with the A allele of the COMT gene
69	Vaegter 2017	TKR				Hypoalgesia after cold pressor stimulation and aerobic exercise assessed preoperatively by cuff algometry was associated with less pain
70	Valdes 2012	THR and TKR				Increased risk with presence of chronic widespread pain
71	Vila 2020	TKR				Increased risk with poorer baseline performance on the colored word matching Stroop test
72	Von Dincklage 2017	THR				Increased risk with higher intraoperative nociception
73	Wang 2014	THR				
74	W-Dahl 2014	TKR				Risk affected by the type of instrument used to measure pain
75	Wylde 2009	THR and TKR				Increased risk with TKR (vs THR)
76	Wylde 2011	THR and TKR				Increased risk with presence of pain elsewhere
77	Wylde 2013	TKR				Increased risk with preop widespread pain sensitization, measured using pressure algometry
78	Wylde 2020	TKR				More severe pain at 13-weeks postop was associated with poorer general health and physical health, more pain worry, and lower satisfaction with surgery outcome. More severe functional limitation was associated with higher levels of depression, more pain worry, lower satisfaction with surgery outcome and higher pain acceptance

Table 3 (continued)

Qualitative summary of risk factors for PPSP reported in multivariable analyses in included studies.

79	Yang 2020	TKR					Decreased risk with Sevoflurane anesthesia
80	Yao 2019	TKR					
81	Zachodnik 2021	THR					No association between older adults with moderate to severe levels of pain during the first 5 days postoperatively and 1 year after surgery

Results are color coded for the type of associations:

Red – increased odds of PPSP; Yellow – no association with PPSP; White – not reported.

TKR = total knee replacement; THR = total hip replacement; Preop = preoperative; postop = postoperative

Results are color coded for the type of associations.

Red, increased odds of PPSP; Green, decreased odds of PPSP; Yellow, no association with PPSP; White, not reported.

and preoperatively facilitated TSP and CPM assessed using mechanical stimuli or manual cuff stimuli with knee osteoarthritis.^{78,80,125} Hypoalgesia after cold pressor stimulation and aerobic exercise assessed preoperatively by cuff algometry was associated with pain relief 6 months after TKR.¹¹⁰

4.6. Potential modifiers

Certain factors such as anxiety, depression, catastrophizing, and BMI are potentially modifiable, and future studies should investigate whether preoperative modification of these factors can mitigate PPSP risk. Cognitive behavior therapy and other holistic approaches may be relevant in this setting.¹¹⁸ Other risk factors such as age, and sex are not modifiable; however, they may be important to developing risk stratification tools. Intraoperative and perioperative factors such as surgical and anesthesia techniques as well as perioperative care and analgesia plans can be modified to mitigate the risk for PPSP in these patients.

4.7. Limitations of our study

It is important to acknowledge the limitations of this review when interpreting the results. First, although we performed a systematic literature search, gray literature as well as abstracts and dissertations were not evaluated, potentially missing some studies. Some full-text articles described data in a format which could not be retrieved for meta-analyses; these data were only included in a qualitative summary. We contacted authors in such situations but were only successful to obtain additional data in a few cases. We included a mix of both retrospective and prospective studies, as well as observational and interventional studies. There are no guidelines how to weigh risk factors based on differences in study design,23 and this may have led to heterogeneity in the study population and recall bias in retrospective studies.⁴¹ Second, with the main outcome of PPSP occurrence as a categorical variable, it is also possible that we missed certain constructs of PPSP as captured by multidimensional or descriptive tools, such as PROMs⁶⁶ after hip and knee arthroplasty which has been well described elsewhere. 40,44,61,117,124,130 Third, this review included data both from studies on THR and TKR, although the literature has shown that pain and patient satisfaction outcomes from THR are generally better in comparison with TKR.⁶⁴ However, we analyzed the data separately, where possible. Fourth, we could not account for the effect of pharmacological and nonpharmacological interventions on the reporting of pain from the available

data. For example, there were several predictor variables reported as important in the individual studies, but for which not enough data were available to conduct a meaningful metaanalysis. These include coping skills, negative beliefs about opioids having a higher PPSP risk,²² biomechanical factors, and QST measures. These factors were qualitatively summarized based on multivariate analyses in some studies but require further and more rigorous evaluation, ideally in the setting of multicenter studies. Several of the included studies described perioperative that were interventions available onlv from 1 study.^{14,18,115,128,19,21,36,51,76,86,88,96} We did not exclude them and summarized their findings qualitatively; however, the ability to draw generalizable conclusions from these studies remains limited. Fifth, there was also marked heterogeneity in the study designs, interventions (eg, spinal anesthesia may offer shorter duration of analgesia compared with continuous epidural analgesia³¹), statistical analyses undertaken, and ROB scores. THA or TKA may be performed under general, regional, or neuraxial anesthesia (or a combination of these); the choice may be based on institutional practices, patient comorbidities, and patient preference.²⁸ In this review, 4 articles focused on regional anesthesia^{14,51,76,128} and 2 on intrathecal analgesia.^{96,115}; however, with insufficient information to draw conclusions on differences in anesthetic approaches in the context of PPSP prevention. Finally, the follow-up periods were grouped to a mean of 6 months postoperatively (as per inclusion criteria) so some precision may be lost in the meta-analyses. As no widely available risk of the bias assessment system is available for a group of studies with mixed designs, we used previously published tools to adapt to the needs of this review; it has been used in reviews with similar methodology⁵⁶; however, the entire battery of questions has not been validated as a single tool.

5. Conclusions

Our review systematically addressed factors associated with risk for PPSP and identified that higher preoperative state anxiety and depression increase the risk of PPSP after TKR. No consistent risk factors for PPSP after THR were identified in the metaanalysis. In the qualitative summary of multivariable analyses performed in individual studies, higher preoperative pain scores and more severe acute postoperative pain were associated with a higher risk of PPSP after TKR or THR. There was a wide variation in the methods of pain assessment and in the tools used to collect perioperative variables. Standardization of tools and methods for perioperative data collection and PPSP assessment, improvement in follow-up rates, proper characterization of patients who dropped off from the studies, and ascertaining the reproducibility of PPSP prevention strategies by conducting multicenter studies can all improve the understanding and mitigation strategies of PPSP after joint replacement surgeries.

Disclosures

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Appendix A. Supplemental digital content

Supplemental digital content associated with this article can be found online at http://links.lww.com/PR9/A184.

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