

## Review Article

# Preoperative Optimisation in Enhanced Recovery after Surgery (ERAS) Programs for Total Hip and Knee Arthroplasty: A Systematic Review

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**Abstract**

**Background:** Enhanced recovery after surgery (ERAS) programs aim to reduce peri- and post-operative complications by targeting modifiable risk factors through pre-operative and peri-operative interventions. The evidence for ERAS in reducing complications following total hip (THA) and knee arthroplasty (TKA) is mixed. This systematic review aimed to describe pre-operative optimisation protocols for THA and TKA and report their effects on post-operative morbidity and patient-reported outcome measures (PROMs).

**Methods:** This systematic review was performed according to the Preferred Reporting Items for Systematic reviews and Meta-Analyses guidelines. Databases were searched for studies with pre-operative optimisation components in ERAS protocols, and data on interventions, frequency, duration, and outcomes (readmissions, complications, PROMs) were extracted. Meta-analysis was not performed due to study heterogeneity.

**Results:** Thirty-eight studies were included. Common interventions included patient education and physiotherapy. There are nine studies offering medical optimisation such as diabetes or weight loss management. Most studies only involved a single clinic visit and lacked clear targets. ERAS protocols led to a decrease in hospital readmissions in six studies (median -1.8%) and an increase in five (median 0.3%) compared to pre-ERAS protocols. Nine studies reported a decrease in post-operative complications (median 5.0%), while four showed an increase (median 1.9%).

**Conclusion:** The review found that pre-operative optimisation protocols within ERAS are poorly designed, with low-quality interventions and limited attention to intervention frequency and duration. Consequently, it remains unclear whether pre-operative optimisation affects readmissions, complications, or PROMs. Future research should focus on prospective studies addressing risk factors like obesity and diabetes to improve patient outcomes.

**Trial registration:** The systematic review protocol was submitted to the International Prospective Register of Systematic Reviews (CRD42021274156).

**INTRODUCTION**

The prevalence of hip and knee osteoarthritis is increasing worldwide [1]. Globally, hip and knee osteoarthritis was ranked as the 11th highest contributor to global disability and 38th highest in disability-adjusted life years [2]. Osteoarthritis affects approximately 9.3% of the Australian population, causing pain and disability, restricting employment and hobbies, and reducing quality of life [3]. Increasing rates of obesity and an

ageing Australian population are driving an increase in prevalence from an estimated 2.1 million in 2015 to an estimated 3.1 million (12% of the population) by 2030 [3]. Total hip arthroplasty (THA) and total knee arthroplasty (TKA) are recommended for end-stage osteoarthritis when all appropriate conservative options, delivered for a reasonable period of time, have failed [4]. Arthroplasty is the most cost-effective and clinically effective treatment for end-stage osteoarthritis in appropriately selected individuals [5,6].

THA and TKA are major surgical procedures associated with post-operative medical and surgical complications. These include, but are not limited to, peri-prosthetic joint infection, peri-prosthetic fracture, osteolysis with implant loosening, and venous thromboembolism (VTE). These complications are associated with significant morbidity and mortality, necessitating hospital readmission, medical treatment and/or revision surgery [7-9]. There are known modifiable risk factors which increase the risk of post-operative complications, including body mass index (BMI)  $\geq 30$ , malnutrition, poor glycaemic control, anaemia, smoking, use of opioids and vitamin D deficiency [10-13]. In addition, poor patient-reported outcomes are also associated with modifiable physiological and psychological factors such as catastrophising, anxiety, depression, and poor self-efficacy [14,15].

In an effort to improve peri-operative outcomes and reduce morbidity, Enhanced Recovery After Surgery (ERAS), a peri-operative care pathway initially developed in the field of colorectal surgery, was adapted for patients undergoing hip and knee arthroplasty [16]. According to ERAS principles, recovery is facilitated by a multimodal approach directed towards global modulation of the surgical stress response [17]. This is achieved through pre-operative patient preparation, standardised anaesthetic regimens intraoperatively, and early mobilisation post-operatively [18]. ERAS protocols are designed and implemented by a multidisciplinary team of surgeons, anaesthetists, nursing, and allied health, and have consistently been associated with faster recovery, decreased morbidity and reduced hospital length of stay (LOS) in multiple surgical subspecialties [19]. However, evidence for the effect of ERAS implementation in orthopaedic surgery on surgical outcomes is conflicting. A recent systematic review concluded that ERAS protocols reduce the length of hospital stay after THA and TKA, but have minimal to no impact on perioperative morbidity or readmission [20]. A meta-analysis concluded that ERAS reduced post-operative complications and 30-day mortality after arthroplasty, with no effect on hospital readmissions [21].

One explanation for the inconsistency in the literature regarding the benefits of ERAS protocols on morbidity and mortality may be the heterogeneity in the pre-operative management of patients, with no clear consensus on how patients should be optimised before TKA or THA. Emerging evidence supports the optimisation of diabetes control and weight loss, however how best it is achieved is very much in its infancy [22]. Only one randomised trial has shown that pre-operative weight loss via bariatric surgery resulted in fewer complications in people with

obesity undergoing TKA [23]. There are no definitive trials proving diet or drug-based weight loss improves outcomes pre-arthroplasty surgery [24]. There is also no high-level evidence demonstrating the efficacy of opioid tapering prior to arthroplasty surgery in chronic opioid users [25].

In light of the current knowledge gaps, it remains unclear what constitutes effective pre-operative optimisation as part of an ERAS protocol for THA and TKA for the purposes of reducing complications and improving patient-reported recovery. This systematic review aimed to: (1) describe the pre-operative optimisation protocols for THA and TKA that are being used worldwide, including information on the type, duration, frequency and goals of interventions; and (2) report the effects of pre-operative optimisation protocols for THA and TKA on post-operative morbidity (including hospital readmission, complications and mortality) and patient reported outcomes.

## METHODS

This systematic review was prepared according to the Preferred Reporting Items for Systematic Reviews and Meta-Analysis (PRISMA) Guidelines [26,27]. The PRISMA checklist is provided in Supplementary Table S1. The systematic review was registered in the International Prospective Register of Systematic Reviews (PROSPERO) (CRD42021274156) and uploaded to medRxiv September 2021 (doi: <https://doi.org/10.1101/2021.09.27.21264189>).

### Eligibility Criteria

**Types of Participants:** Studies including adults aged 18 or older receiving primary elective THA or TKA surgeries were included. The surgeries could be either unilateral or bilateral and no restriction was placed on sex or race. Studies including participants receiving partial arthroplasties (i.e., uni-compartmental or hemiarthroplasties) or arthroplasties indicated for fracture, were not included.

**Types of Intervention:** Eligible studies investigated pre-operative optimisation protocols administered prior to participants receiving a primary elective THA or TKA, with/without a comparative group that received a standard care protocol. No restriction was placed on the duration of pre-operative optimisation protocol. Only studies that had a comparative group were used for aim 2 of this review. To provide a comprehensive review on pre-operative optimisation, all studies investigating ERAS with a pre-operative component were included in the review.

**Type of Outcome Measures:** The primary outcomes

of this review for assessing aim 2 were: 1) hospital readmissions within 90 days; and 2) any complication including surgical site or other infection, cardiovascular event, VTE, or death. The secondary outcomes included length of hospital stay, and patient-reported pain and function outcomes. Studies were only included if investigating post-operative outcomes. No restriction was placed on when the outcomes were measured in the studies.

**Types of Studies:** For Aim 1, we included randomised controlled trials (RCTs), non-randomised clinical trials, prospective observational studies, and retrospective studies. For Aim 2, only RCTs or non-randomised clinical trials were to be included in meta-analyses if one was possible. Systematic or literature reviews, case reports or series, or conference abstracts were excluded.

### Search Strategy

To identify eligible, published studies, we searched the following electronic databases:

- MEDLINE
- EMBASE
- Cumulative Index to Nursing and Allied Health Literature (CINAHL)
- Cochrane Central Register of Controlled Trials (CENTRAL)

Search strategies were established using medical subject headings (MeSH) and related text words and tailored to each database. A combination of different keywords for THA, TKA and pre-operative optimisation protocols were used to identify relevant studies. The full search terms and search strategy are included in Supplementary Table S2. No restriction was placed on the publication period, but only studies in the English language were included. We searched the reference lists of eligible studies and relevant reviews to include any missed but relevant published studies. Citation searching for forward citation of recent studies and citation alerts (i.e., Google Scholar) on included studies were also be used to identify new studies as they appeared during the review progress. The search strategy was run in October 2021 and again in March 2023, several weeks prior to manuscript drafting, to retrieve and include any relevant studies.

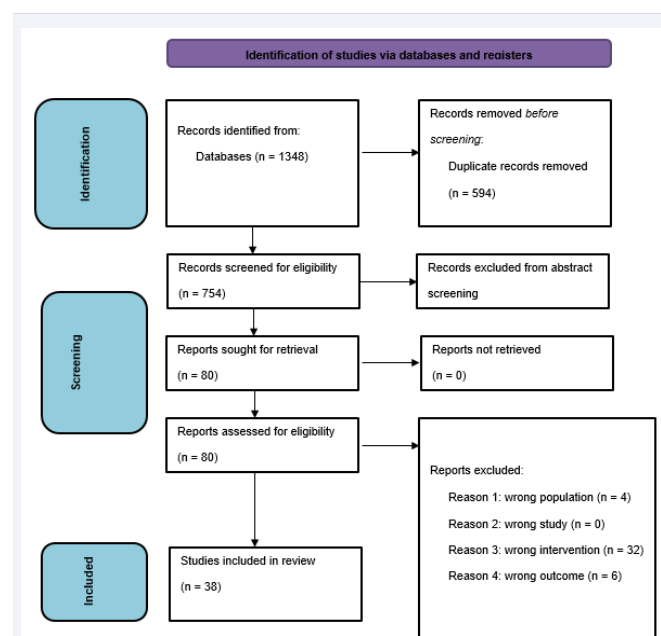
### Study Selection

EndNote X9 software (Clarivate Analytics) was used to store, organise, and manage all search results and to remove duplicate records. Two reviewers independently

evaluated the title and abstract of all studies identified through the search against the eligibility criteria. The full text of the selected studies was then retrieved. For studies retrieved from trial registrations, the full text was defined as all associated files and information. For any studies with uncertainty about the eligibility, the full text was obtained for further information. Any disagreement over study eligibility was resolved by consensus, and an additional reviewer was consulted if required. The Preferred Reporting Items for Systematic Review and Meta-Analysis Flow Chart with reasons for exclusion is presented in Figure 1.

### Data Extraction

A customized data extraction spreadsheet was developed and piloted on two studies relevant to this review before it was used to extract data from all the included studies. Two reviewers independently performed data extraction from the final list of selected studies.



**Figure 1** PRISMA flow diagram outlining the systematic review process. PRISMA, Preferred Reporting Items for Systematic Reviews and Meta-Analyses

Reason 1: Is the population adults aged > 18 and received selective THA or TKA (either primary or revision; unilateral or bilateral)? Excluded if: partial or hemi-replacement or due to acute fracture

Reason 2: Was the study a RCT, non-RCT, prospective observational study or retrospective study? Excluded if: systematic or literature reviews, case reports or series or conference abstracts

Reason 3: Did the study investigate an ERAS program prior to participants receiving THA/TKA (with or without a comparative group)? Excluded if: ERAS program commenced after THA/TKA.

Reason 4: Is the study primary outcome: hospital readmission within 90 days or complications after THA/TKA? Excluded if: primary outcome is not readmission/complications after THA/TKA

Any disagreements in the extracted data were resolved through discussion with a third reviewer. We extracted the following information from the included studies:

- Study characteristics: the first author, year of publication, study design, country, and study setting.
- Participants: age, sex, type of surgery (i.e., THA or TKA, unilateral or bilateral, left or right), duration of knee/hip osteoarthritis, co-morbidities, use of opioids, and the number of participants allocated in each intervention group.
- Interventions: details of the pre-operative optimisation protocol (i.e., type, duration, number of interventions, frequency, intervention providers, goals of intervention), details of pre-, peri- or post-operative care protocols as a part of a continuum optimisation program if available.
- Outcome measures: the type of measure used to assess primary and secondary outcomes, at any peri- or post-operative timepoints.
- Results: data on the primary and secondary outcomes measured at any timepoints.
- If data were missing, authors of the studies were emailed a maximum of three times, after which the data was considered irretrievable.

### Study Quality and Risk of Bias

Study quality and risk of bias was assessed by two independent reviewers using the Cochrane Risk of Bias version 1 (RoB 1) tool for RCTs [28], and the Cochrane Risk of Bias in Non-Randomised Studies - of Interventions (ROBINS-I) tool for non-randomised studies [29]. Disagreement between reviewers was resolved through discussion and a third reviewer was consulted if consensus was not achieved. The risk of bias was evaluated on the following domains: random sequence generation, allocation concealment, blinding of participants and personnel, blinding of outcome assessment, incomplete outcome data, selective reporting, and other potential sources of bias. The ROBINS-I tool was used to assess the risk of bias for studies that have not used randomisation for intervention allocations (i.e., cross-sectional or cohort study designs). The ROBINS-I tool covers seven domains: confounding and participants' selection (pre-intervention), intervention classification (during intervention), and deviations, missing data, measurements and selection of reported results (post-intervention) [29]. Both RoB 1 and ROBINS-I used signalling questions to guide judgments for each domain and an overall risk of bias assessment.

### Data Synthesis

A narrative summary was conducted to provide an overview of pre-operative optimisation protocols for THA and TKA respectively. The details of pre-operative optimisation protocols from all selected studies were summarised in Table 1. The risk factors targeted by pre-operative optimisation protocols and the corresponding intervention provided were listed. For aim 2, we planned a meta-analysis, however it was not feasible given the high heterogeneity and high risk of bias of included studies. Thus, a narrative approach was used to summarise the results from non-RCTs and retrospective studies based on study quality and type of surgery (THA/TKA). Study findings including data reduction, display and comparison, conclusion and the classification of evidence from individual studies were then summarised in tabular form [30,31]. From each study, the extracted data on post-operative morbidity (including hospital readmission, complications and mortality) and PROMs were descriptively analysed and summarised in charts.

### RESULTS

The search strategy generated a total of 754 unique articles, of which 80 full-text articles were assessed for eligibility. Following full text screening, 38 studies were eligible for analysis (Figure 1).

### Study Characteristics

Ten studies were prospective cohort studies [32-85], Seventeen were retrospective studies [34-90], seven studies utilised the before and after design [38-81], and there were four case control studies [33-91]. No RCTs were included in the review. Twenty studies investigated patients undergoing THA and TKA, ten studies investigated TKA only and four studies THA only. Unilateral and bilateral TKA, unilateral and bilateral THA, bilateral TKA and bilateral THA were each investigated by one study. In all studies, there was a pre-operative, intra-operative and post-operative component to the ERAS protocol. Two studies investigated pre-operative optimisation as a stand-alone intervention in reducing readmissions or complications, with standardised intra-operative and post-operative protocols between groups [32,33]. All remaining studies had different peri-operative and post-operative protocols between groups and the effect size of the ERAS intervention was attributed to the implementation of all three components.

### Type of Interventions

Thirty-two studies incorporated patient education and thirteen studies included physiotherapy as part of their



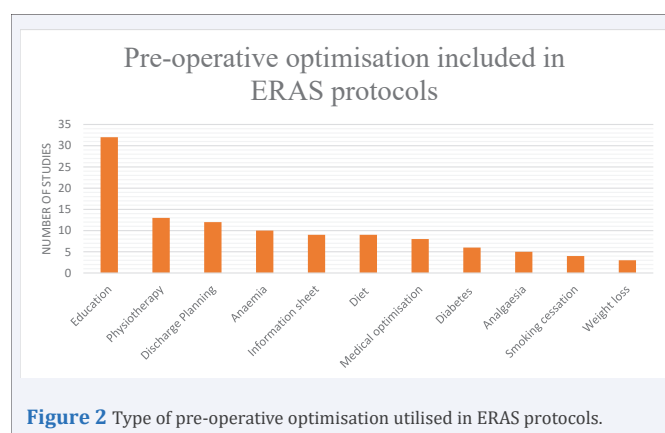
pre-operative optimisation (Figure 2). In fourteen studies this was the only form of optimisation offered to patients. Nine studies offered medical optimisation before surgery, with five specifically focusing on diabetes (Figure 3). Three studies included weight loss for obesity management, but none specified the type of weight loss intervention offered. The details of pre-operative optimisation offered are summarised in Table 1.

### Duration of Intervention

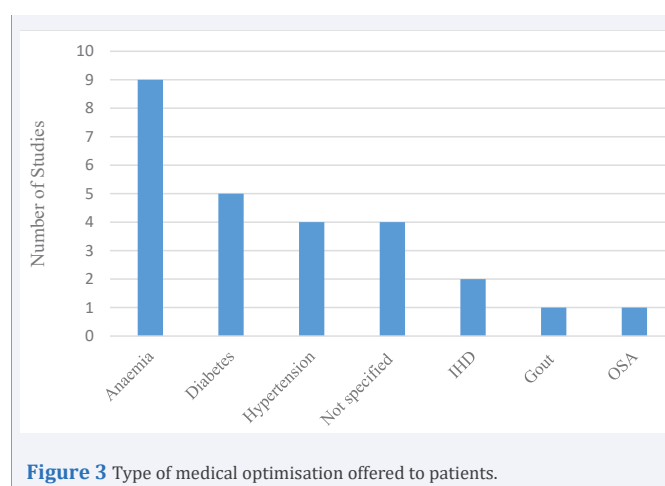
In twenty-one studies the duration of optimisation was assumed to be one day [33-90]. This consisted of a single preoperative clinic visit which incorporated education, physiotherapy or discharge planning without further optimisation before surgery (Figure 4). Four studies optimised patients for 4 weeks [32- 85], and two studies optimised patients for approximately 2 months [34, 35]. In one study surgery was delayed if necessary, allowing for sufficient duration for medical comorbidities to be managed [36]. In ten studies, the duration of optimisation was not stated.

### Goals of Optimisation

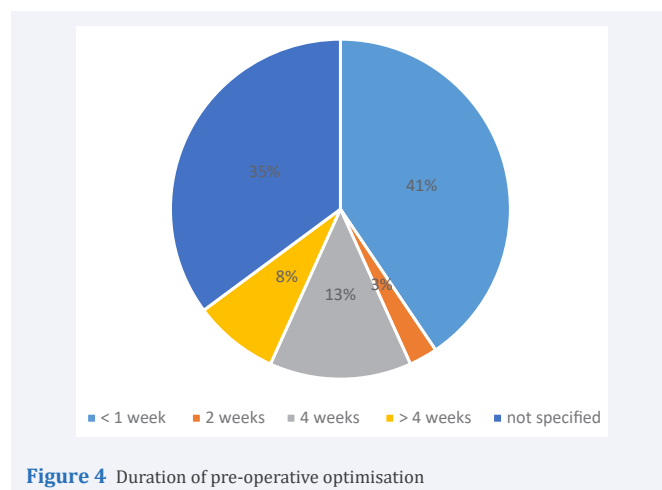
In the five studies incorporating diabetes management,



**Figure 2** Type of pre-operative optimisation utilised in ERAS protocols.



**Figure 3** Type of medical optimisation offered to patients.



**Figure 4** Duration of pre-operative optimisation

only one study set a target HbA1c for optimisation. None of the three studies utilising weight loss as an optimisation set a target weight or weight loss percentage. Two of the nine studies that included anaemia optimisation set a target haemoglobin. No studies followed up patients after the implementation of an intervention. In other words, the specific effect of the intervention on the organ or target of interest (e.g. effect of dietary intervention of weight) was not evaluated in any study.

### Clinical Efficacy of Pre-Operative Optimisation Protocols on Post-Surgical Outcomes

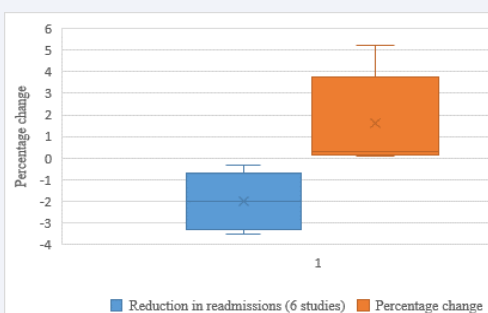
**Primary Outcomes:** The effects of pre-operative optimisation protocols alone on hospital readmissions and complications (both medical and surgical) following THA and TKA were unclear as most studies included differing pre-, peri-, and post-operative ERAS interventions, and did not investigate the stand-alone effect of pre-operative optimisation. The implementation of ERAS protocols showed a decrease in hospital readmissions within 90 days in six studies (median -1.8%, interquartile range (IQR) -2.3%) [33-41], and an increase in five studies (median 0.3%, IQR 3.6%) [36-45], compared to pre-ERAS protocols (Figure 5). For post-operative complications, nine studies [32-48], showed a decrease in incidence upon implementation of ERAS (median -5.0%, IQR -5.1%), with four studies [36-49], recording an increase (median 1.9%, IQR -5.3%) (Figure 6).

**Secondary Outcomes:** No study investigated the isolated impact of pre-operative optimisation (controlling the intra-operative and post-operative variables) on the length of stay or patient-reported outcomes. Therefore, the effects on in secondary outcomes were attributed to the implementation of ERAS as a whole, with no indication of the weighting of the pre-operative component to these differences. The implementation of ERAS resulted in a

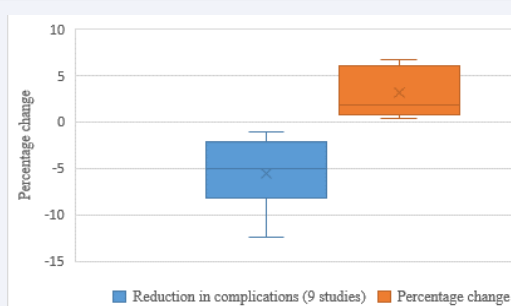
**Table 1:** Characteristics of pre-operative optimisation protocols from included studies

Study Author	Year	Study Design	Summary of pre-operative optimisation
Ascione et al[51]	2020	Prospective cohort	Preoperative education and physiotherapy
Auyong et al[37]	2015	Retrospective cohort	Preoperative education and discharge planning
Azam et al[42]	2021	Prospective cohort	Preoperative education, physiotherapy, diabetes optimisation (HbA1c < 6.5%), hypertension and anaemia optimisation, dietary supplementation and smoking cessation advice
Blum et al[46]	2019	Retrospective cohort	Preoperative education
Christellis et al[38]	2015	Before-and-after intervention	Preoperative education, physiotherapy and dietician counselling
de Carvalho et al[49]	2021	Prospective cohort	Preoperative education, physiotherapy and discharge planning
Frassantio et al[75]	2020	Prospective cohort	Preoperative education and physiotherapy
Garriga et al[76]	2019	Retrospective cohort	Medical assessment of haemoglobin levels and comorbidities including cardiovascular/respiratory disease
Garriga et al[77]	2019	Before and after intervention	Preoperative education, physiotherapy and discharge planning
Glassou et al[78]	2014	Retrospective cohort	Preoperative education
Gleicher et al[79]	2021	Before-and-after intervention	Preoperative education and discharge planning
Gwynne-Jones et al[43]	2017	Before-and-after intervention	Weekly preoperative education and physiotherapy
Hansen et al[32]	2012	Prospective cohort	Identification and optimisation of risk factors: malnutrition, obesity, general health (hypertension, diabetes, infection), physiotherapy, cessation of smoking and alcohol consumption.
Husted et al[80]	2006	Prospective cohort	Preoperative education and discharge planning
Jiang et al[47]	2019	Prospective cohort	Preoperative physiotherapy
Kelmer et al[33]	2021	Case control	Preoperative education, physiotherapy and discharge planning
Maempel et al[81]	2015	Before-and-after intervention	Preoperative education and physiotherapy
Maempel et al[52]	2016	Retrospective cohort	Preoperative education and physiotherapy
Malviya et al[48]	2011	Retrospective cohort	Preoperative education
Otte et al[82]	2011	Case control	Preoperative education
Pamilo et al[44]	2018	Before-and-after intervention	Preoperative education and discharge planning
Pamilo et al[45]	2018	Before-and-after intervention	Preoperative education and discharge planning
Petersen et al[83]	2020	Prospective cohort	Preoperative education
Picart et al[39]	2021	Case-control	Preoperative education and iron deficiency anaemia treated
Plenge et al[84]	2020	Prospective cohort	Preoperative education, assessed in optimisation clinic, pre-op analgesia rationalised, treatment of anaemia
Qiu et al[36]	2016	Retrospective cohort	Preoperative education, medical optimisation (diabetes, hypertension, sleep apnoea, gout) in primary care and specialist setting, lifestyle modification (smoking, obesity)
Ripolloes-Melchor et al[85]	2020	Prospective cohort	Preoperative education, smoking and alcohol cessation 4 weeks before surgery, anaemia treated
Savaridas et al[86]	2013	Retrospective cohort	Preoperative education
Stambough et al[87]	2015	Retrospective cohort	Preoperative education and analgesia optimisation
Starks et al[40]	2014	Retrospective cohort	Preoperative education, discharge planning, and pre-existing medical conditions optimised (anaemia, hypertension, ischaemic heart disease)
Stowers et al[88]	2016	Retrospective cohort	Weekly education classes
Van Horne et al[34]	2019	Retrospective cohort	Preoperative education, physiotherapy, medical optimisation of modifiable risk factors, weight loss and analgesia optimisation
Van Horne et al[35]	2020	Retrospective cohort	Preoperative education, physiotherapy, medical optimisation of modifiable risk factors, weight loss and analgesia optimisation
Vendittoli et al[89]	2019	Retrospective cohort	Preoperative education, physiotherapy, optimisation of anaemia, diabetes and other medical comorbidities
Wilches et al[50]	2017	Retrospective cohort	Preoperative education, physiotherapy and treatment of anaemia
Winther et al[90]	2015	Retrospective cohort	Preoperative education and physiotherapy
Yanik et al[41]	2018	Retrospective cohort	Preoperative education
Zhang et al[91]	2018	Case control	Preoperative education, physiotherapy, diet supplementation and analgesia optimisation

\*\*When an underlying medical condition such as anaemia was identified in the preoperative period, surgery was rescheduled



**Figure 5** Box and whisker plot demonstrating change in readmission rate after implementation of ERAS. Separated into studies showing increase/decrease in readmissions.



**Figure 6** Box and whisker plot demonstrating change in complication rate after implementation of ERAS. Separated into studies showing increase/decrease in complications.

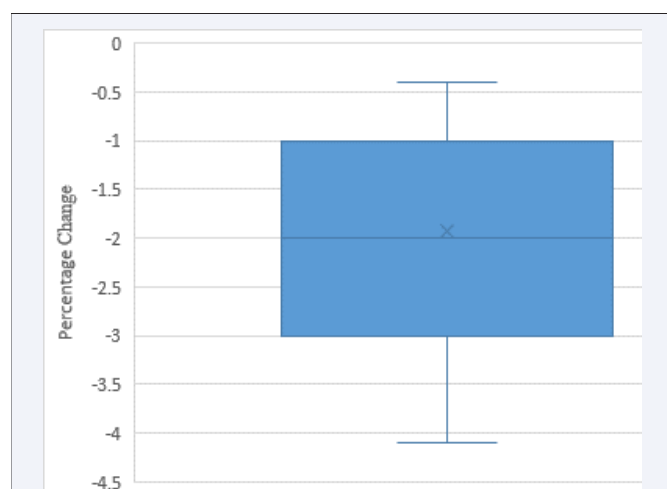
reduction in length of stay in 23 studies (median -2 days, IQR 2) (Figure 7), with one study reporting an increased length of stay (3.3 days) [50]. Patient-reported outcome measures were improved in seven studies [37-52]. Objectively measured physical function was improved after ERAS implementation in three studies [37-42]. No studies reported a deterioration in PROMs following implementation of ERAS.

### Risk of Bias

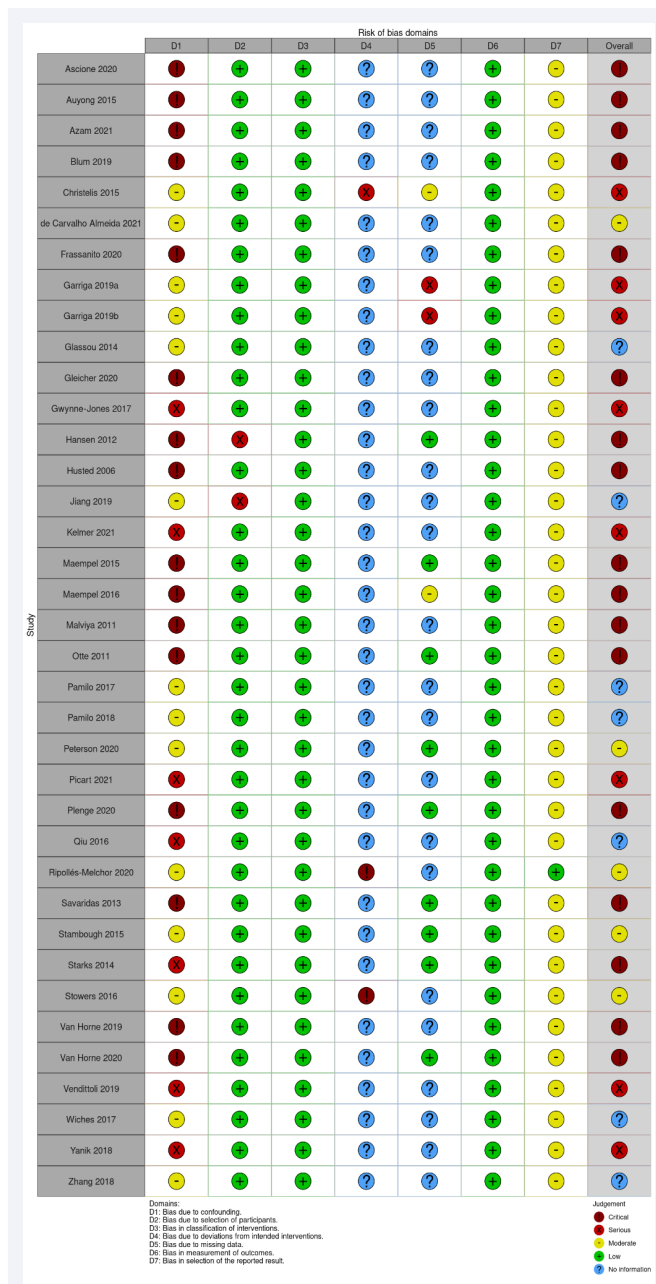
All studies included in the systematic review were observational studies, and a moderate to critical risk of bias was present in all studies (Figure 8). A moderate to critical degree of bias due to confounding affected all studies. There was low risk of selection and measurement bias, and an unclear risk of attrition bias.

### DISCUSSION

This systematic review found that pre-operative optimisation protocols before THA and TKA focused heavily on patient education and physiotherapy, with a minority of studies optimising medical comorbidities and obesity. The duration of optimisation is likely to be inadequate in most studies for improvement of physiological processes or reversal of pathophysiology, and typically no follow-up has been undertaken to ensure patients are adequately optimised prior to surgery. There is conflicting evidence whether ERAS reduces complications or readmissions (Figure 5,6), however it appears that the length of hospital stay is reduced, and PROMs are improved. However, the evidence for the efficacy of pre-operative optimisation cannot be determined as very few studies investigated pre-operative optimisation alone in reducing complications or readmissions.



**Figure 7** Box and whisker plot demonstrating change in length of stay after ERAS implementation



**Figure 8** Risk Of Bias In Non-randomised Studies - of Interventions (ROBINS-I) tool

The application of ERAS protocols to the peri-operative practice of multiple surgical sub-specialties has produced significant clinical and economic benefits. It was initially implemented in the field of colorectal surgery and there is a large body of evidence which demonstrates significant reductions in morbidity, faster recovery, and reduced hospital length of stay [53-55]. A significant amount of research has confirmed the efficacy of intra-operative components of ERAS. Standardised intra-operative anaesthetic protocols utilise neuraxial and regional anaesthesia, goal-directed fluid therapy, multimodal analgesia and prevention of post-operative nausea and

vomiting [56], and these intra-operative practices are associated with a lower incidence of post-operative pulmonary complications, acute kidney injury and opioid analgesia requirements [17-59]. Furthermore, surgical practices including the use of tranexamic acid to reduce blood loss, abolition of tourniquet use and surgical drains and early indwelling catheter removal are all evidence-based recommendations with robust evidence to support their use or not [60].

In comparison, the current pre-operative interventions for THA and TKA do not have the evidence to support their routine inclusion in ERAS protocols. Our review has shown that 23 of the 38 studies (60.5%) only offered patient education and/or physiotherapy with no other interventions. Although patient education is a low-cost intervention and unlikely to cause harm, two Cochrane reviews found little evidence that education influences pain, recovery, or reduce length of hospital stay and post-operative morbidity [61,62]. Pre-operative physiotherapy, in isolation, also does not confer any clinically important benefits and does not shorten length of hospital stay [63]. Therefore three-fifths of all current pre-operative protocols utilise ineffective strategies to optimise patients' outcomes following THA and TKA.

The length of pre-operative optimisation was found to be grossly inadequate in most studies. Sixty three percent of the protocols interrogated had allocated one day (a single pre-admission clinic appointment) as the duration of the pre-operative optimisation. This is especially concerning in medical optimisation as the topic was scarcely addressed in the included studies. Medical optimisation is a key aspect of the ERAS protocol, aim to address numerous conditions that may influence surgical outcomes. By optimising these conditions prior to surgery, ERAS protocols aim to reduce complications such as infections, delayed wound healing, and prolonged hospital stays, ultimately improving overall recovery [64].

Despite the established benefits of preoperative medical optimisation, its application in practice appears to be limited. Among the studies that did offer some form of medical optimisation the duration varied significantly for patients- ranging from 2 weeks to over 1 month. Only one study which offered medical optimisation of chronic conditions specified a variable duration of optimisation to ensure sufficient time for patients' conditions to be assessed and treated. This highlights a gap in the literature, as many studies did not detail the timing or adequacy of the preoperative optimisation period. No studies allocated follow-up to assess the efficacy of the interventions on the specific chronic condition, and no studies indicated that

surgery was postponed if patients were not adequately optimised before surgery. The lack of consistency in the application and duration of medical optimisation in these studies points to a significant area for improvement within ERAS programs, where a more standardised approach to the management of comorbidities could help improve patient outcomes and safety.

Diabetes and obesity are two chronic medical conditions that have strong associations with post-operative complications including surgical site infection, periprosthetic joint infection, VTE and myocardial infarction [65-70]. It is postulated that hyperglycaemia adversely affects wound healing by delaying the synthesis of collagen and suppresses the immune response to infection by impairing phagocytosis [71,72], and obesity may influence the risk of a complication via a number of pathways including the influence of visceral fat on dysmetabolism and chronic inflammation [73]. These conditions are rarely diagnosed in isolation, and are strongly associated with other co-morbidities that may independently increase risk of complications such as coronary artery disease, hyperlipidaemia, hypertension, and sleep apnoea [74].

Our review has demonstrated that neither the length of optimisation nor the efficacy of interventions was sufficient to address these multifaceted issues. The three studies incorporating weight loss did not specify a modality or target weight, and only one study specified the duration of the weight loss program (4 weeks). Only one of the five studies optimising diabetes set a target HbA1c (6.5%), and similarly no duration was specified. Only one study referred patients to a specialist physician for management of diabetes. Obesity and diabetes are complex medical issues which have afflicted patients for many years prior to their arthroplasty surgery, and it is essential that these conditions are addressed by a multidisciplinary team of physicians, dieticians and allied health professionals [74]. Goals should be established, and sufficient time must be allocated prior to their surgery for interventions to be implemented for patients to have these conditions adequately optimised. The Australian Diabetes Society has set a target HbA1c of 8.0% for patients undergoing surgery and recommends postponing surgery if elevated. The Royal Australian College of General Practitioners has recommended a minimum weight loss target of 5-7.5% of body weight for patients with osteoarthritis [4]. There is no clear consensus on the duration of optimisation, however patients who are on the public hospital waiting list for THA (median 179 days) and TKA (median 308 days) [75], have a flexible time interval of several months which can be exploited to optimise patient comorbidities



in a multidisciplinary setting. Patients undertaking arthroplasty in a private hospital setting have a much smaller window for optimisation due to a significantly shorter waiting period (usually less than a month), and therefore the potential for optimisation of risk factors is considerably reduced.

The strength of this review is comprehensive analysis of all currently available pre-operative ERAS protocols. This is the first review presenting a thorough analysis of pre-operative optimisation components offered in all ERAS protocols. Other strengths of this review include the a priori protocol which was pre-registered and the strict adherence to the PRISMA guidelines for performing and reporting the review. The main weakness of this review is the moderate to high risk of bias in most included studies, which were observational and retrospective in nature. Further, intra-operative and post-operative components were not standardised between groups. The intervention being assessed in most studies was ERAS as a whole (encompassing pre-, peri-, and post-operative protocols), with only two studies evaluating the impact of pre-operative interventions on patient complications and readmissions. Meta-analysis of complications and readmissions was unable to be performed on the differences between ERAS and pre-ERAS groups given the degree of confounding in the results. Although a meta-analysis was not performed, it was clear that there was little evidence to inform the efficacy of pre-operative optimisation prior to THA or TKA. The effects of the pre-operative interventions studies (focusing heavily on education and physiotherapy) did not significantly change complication or readmission rates. Future studies investigating the utility of pre-operative optimisation must address the risk factors associated with post-operative adverse events, including diabetes, obesity and opioid use, with specified duration and targets for optimisation. Intra-operative and post-operative treatments between intervention groups must be standardised to reduce risk of bias.

## CONCLUSION

Within the ERAS paradigm, current pre-operative optimisation protocols for THA and TKA are poorly designed, providing limited insight into the duration, frequency and number of interventions offered, with a heavy focus on low-quality interventions and little attention towards the duration, frequency and number of interventions offered. There is a paucity of evidence to guide the pre-operative management of chronic conditions known to be associated with post-operative morbidity and mortality. Given these limitations, it is unclear whether pre-operative optimisation affects readmission rates, complications or

PROMs. Future work should be prospective and controlled, addressing risk factors associated with post-operative morbidity, particularly obesity and diabetes, in order to optimise patient outcomes.

## DECLARATIONS

### Availability of data and materials

The datasets used and/or analysed during the current study are available from the corresponding author on reasonable request.

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### Authors' contributions

VL and JL drafted the manuscript. WJC, JN and SA conceptualised the study and developed the methodology of the review. All the authors reviewed and approved the final version of the manuscript.

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