



# Effectiveness of eHealth Tools for Hip and Knee Arthroplasty: A Systematic Review

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**Objective:** This study aimed to compare the effectiveness and costs of eHealth tools with usual care in delivering health-related education to patients' undergoing total hip or knee arthroplasty due to osteoarthritis.

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**Data Sources:** Six electronic databases were searched to identify randomized controlled trials and experimental designs (randomized or not) examining the effect of eHealth tools on pre- or post-operative care. Only manuscripts written in English were included. In the current study, no specific primary or secondary outcomes were selected. Any study that investigated the impacts of eHealth tools on hip or knee arthroplasty outcomes were included.

**Review Methods:** Two researchers reviewed all titles and abstracts independently and in duplicate. Two researchers also conducted full-text screening and data extraction from the 26 selected articles.

**Results:** The data were descriptively reported, and themes could emerge from each outcome. Two researchers separately assessed the Risk of Bias for each paper using the Cochrane risk of bias assessment tool. The majority of studies evaluated the impact of eHealth tools on physical (n = 23) and psychosocial outcomes (n = 19). Cost-related outcomes were measured in 7 studies. eHealth tools were found to be equivocal to usual care, with few studies reporting statistically significant differences in physical or psychosocial outcome measures. However, cost-related outcomes showed that using eHealth tools is more cost-effective than usual care.

**Conclusions:** This review demonstrated that eHealth tools might be as effective as usual care, and possibly more cost-effective, a crucial implication for many overly burdened health care systems.

Keywords: eHealth, rehabilitation, joint replacement surgery, arthroplasty, prehabilitation

#### EHealth for Hip/Knee Arthroplasty

### SUMMARY

#### Strengths and limitations of this study

- Including any type of eHealth interventions.
- Investigating the impact of eHealth interventions on physical, psychological, and cost-related outcomes.
- Not conducting meta-analysis due to diversity in eHealth interventions.

### INTRODUCTION

Osteoarthritis (OA) is a leading contributor to global mobility impairment, driving the rapidly-increasing demand for total hip and knee arthroplasty (THR, TKR) surgery in the US and Canada (1, 2). THR and TKR surgeries, in turn, result in substantial health-care costs (3). Health promotion interventions delivered mainly by physiotherapists and occupational therapists, such as preoperative (prehab) and postoperative (rehab) education which may focus on different topics such as exercise, pain management, nutrition and weight management, surgery and precautions and recovery after surgery, are vital in optimizing surgical outcomes and reducing hospitalization costs (4-6). Several studies have shown that providing prehab and rehab education, which increases health literacy (7), is effective in reducing preoperative anxiety in patients undergoing joint arthroplasty (8, 9) and in patients with other types of surgeries (10-12). Both prehab and rehab education can reduce direct and indirect costs by up to 30% in patients who undergo joint arthroplasty (13), improve patient care, and recovery (14, 15), reduces hospitalization stays by half (13, 16), and improves physical functioning and quality of life in elective surgical procedures (17-19).

Currently, most prehab and rehab education is provided either through in-person and group sessions or educational booklets (9). These types of educational delivery methods which consist the majority of the current "usual care" may be inaccessible for many patients due to various reasons, including mobility issues due to OA, not being able to take time off work to attend inperson and group sessions, and not being able to travel great distances if living in remote and rural areas (20). Reduced access to education can result in lower health literacy (e.g., insufficient knowledge about surgery and precautions) which is the single best predictor of poor health outcomes (21–23).

EHealth tools offer an attractive alternative mode of delivery for health-related education. In this study, eHealth is used to refer to any type of intervention or treatment that is delivered with information and communication technology (e.g., videoconferencing, telemonitoring, and phone calls). Health education using eHealth approaches has been lauded for being interactive and enabling learners to re-engage over sustained periods. eHealth also has the potential to improve the quality of care for older adults (24–26), enhance communication between patients and health care providers [e.g., medical professionals] (27), reduce care costs, and increase access to health care and evidence-based health information. As older adults are increasingly using computers to seek health information (28), the feasibility of harnessing eHealth tools for patients undergoing THR and TKR due to OA is also increasing, providing an opportunity for older adults to benefit from the advantages of eHealth tools. Finally, considering the restrictions caused by the COVID-19 pandemic in accessing in-person education, eHealth education can be considered as a reliable and sustainable way of delivering education (29).

The development of eHealth tools and their evaluation for delivering education for patients undergoing THR and TKR is gaining prominence (30). However, there is no systematic evidence evaluating whether eHealth tools are effective in this space. The present study aimed to assess the effectiveness of eHealth tools (any tools that use information and communication technologies) on the outcomes (any outcomes including physical, psychological, and cost-related outcomes) of patients undergoing THR and TKR due to OA.

## METHODS

#### Patient and Public Involvement

"No patient involved."

### **Eligibility Criteria**

The current study was not limited to only one type of eHealth tools. Specifically, articles were included for analysis if they met the following inclusion criteria: (1) included patients with hip/knee osteoarthritis who either will have or had a hip/knee arthroplasty; (2) studied (randomized or not randomized experimental study designs) eHealth tools (mentioned telehealth, mHealth, eHealth, phone calls tools in the title and/or abstract); (3) investigated the economic, psychosocial or physical impact of eHealth tools (no primary or secondary outcomes selected); (4) published manuscripts and advance access publications in peerreviewed journals; (5) mentioned hip/knee replacement surgery or arthroplasty in the title and/or abstract, and (6) were published in English. Studies were excluded if they were (1) on other joint arthroplasties (e.g., shoulder, ankle); (2) were on the management of OA; (3) were not related to eHealth for patients undergoing THR/TKR; (4) were related solely to the cost of using eHealth tools on health care system; (5) were cohort studies and nonrandomized controlled trials; and (6) were without complete data (e.g., protocols of RCTs and conference abstracts).

### **Study Sources and Searches**

This research applied a systematic search approach to investigate the impacts of eHealth tools on patients with hip and knee arthroplasty. The following databases were searched without being limited to any date: Ovid MEDLINE, Embase, Cochrane Controlled Register of Trials (Central), CINAHL, Web of Science, and Google Scholar (Please see **Supplementary Material A** for an example of the search strategy used in this study). The first search strategy was conducted in June 2018. The search has been updated in July 2019 and June 2020. **Supplementary Material A** presents the search strategy used for Medline. To search Google Scholar, four separate but simplified search strategies were created and the first 100 results for each search strategy were examined at search time points.

## **Study Selection**

To select the potential articles for full-text screening, two researchers (SM and a research assistant who received training from a librarian prior to screening) reviewed all titles and abstracts independently and in duplicate. Full-text screening and data extraction from the selected articles were also conducted by two researchers independently (SM and a research assistant who received training from a librarian prior to screening) and in duplicate. In all the phases, if there were any disagreements, the researchers discussed the issue, and a third researcher (JMR) was consulted to resolve the final conflicts when consensus could not be achieved.

## **Data Extraction and Quality Assessment**

All selected articles for full-text review were imported to Covidence (Covidence.org, Melbourne, Australia) to facilitate data extraction. The data on the author-corresponding information, method (e.g., duration of the study), population (e.g., inclusion and exclusion criteria, demographic information of the sample), intervention (e.g., description, duration, timing), and outcomes were extracted.

To assess the risk of bias of the sample, the standard Cochrane Risk of Bias assessment form (31) which was integrated into Covidence.org was used. The Cochrane Risk of Bias assessment form assesses sequence generation, allocation concealment, blinding of participants and personnel, blinding of outcome assessors, incomplete outcome data, selective outcome reporting, and other sources of bias (e.g., funding). Quality assessment of the sample completed by two researchers separately, and then all the assessments were compared. Disagreements were first discussed among the two assessors. If consensus could not be achieved, a third researcher was consulted.

The Cochrane risk of bias assessment tool defines a study with a "low risk of bias" as a study that has low risk of bias for all domains, a study with unclear risk of bias has been defined as a study with "unclear risk of bias for one or more key domains." Finally, a study with high risk of bias has been defined as a study "High risk of bias for one or more key domains" (32).

# **Data Synthesis and Analysis**

Meta-analyses were not conducted in the sample due to the substantial heterogeneity of the outcomes and the measurements. Hence, the outcomes were compiled into themes, and a descriptive and narrative synthesis of the data was used to provide information regarding the findings of the included studies and assess the impact of prehab and rehab education on patients' outcomes after total hip or knee arthroplasty due to OA.

# RESULTS

## **Search Results**

The detailed search in six main databases resulted in 12,032 references. After removing the duplicates, 6,312 abstracts remained for the title and abstract screening. Independent title and abstract screening resulted in the exclusion of 6,209 articles. In total, 103 articles were selected for full-text screening. Of these, 21 articles were excluded because participants did not have OA

of hip or knee; 20 were excluded due to study design (e.g., cohort studies); 15 articles were excluded as they were RCT protocols; 10 were excluded because they were conference abstracts; five had wrong patients population; two were not in English, and one was excluded because of outcome type (i.e., clinicians' outcomes). A total of 29 articles (33–61) were included in this study. **Figure 1** shows the PRISMA flow diagram of search returns that were retrieved and included.

# **Study Characteristics**

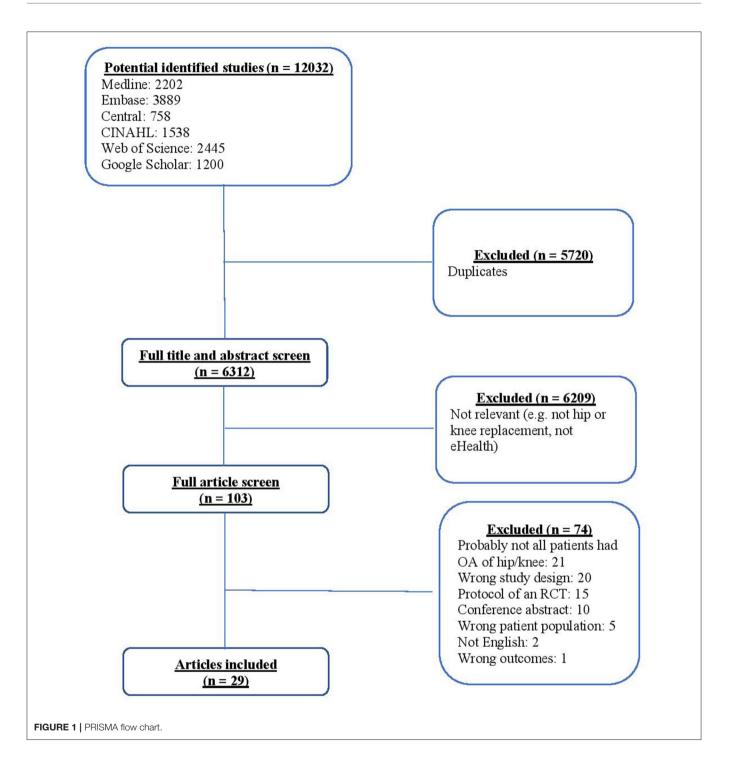
Of the 29 studies that formed our final sample (33-61), most of the studies had one intervention arm and a control arm. However, two studies had two interventions arms (50, 53). Duration of the studies ranged from 7 days (54) up to 52 weeks [e.g., (33)]. Studies were performed in 10 different countries: Canada, USA, Denmark, Germany, Portugal, Spain, UK, South Korea, China, and Australia. Except for one study (53) all interventions were conducted postoperatively. The number of participants ranged from 5 to 209. Some examples of interventions included six 45-min telerehabilitation sessions vs. six 45-min in-person rehabilitations for 6 weeks (34); a web-based follow-up with a surgeon vs. an in-person appointment with the surgeon (40); and 16 sessions of 45-60 min videoconferences vs. 16 45-60 min in-person physiotherapy sessions for 16 weeks. Table 1 lists all descriptive characteristics of the studies included in the final sample.

# Participants

In our sample, 25 studies targeted patients undergoing total knee arthroplasty due to the OA. The remaining studies recruited patients undergoing total hip arthroplasty or a combination of patients undergoing hip or knee arthroplasty.

# Type of Interventions

Delivering rehabilitation sessions via video conference (i.e., telerehabilitation) by an expert (e.g., physiotherapist) was one of the main eHealth approaches used in our sample (n = 9)(34, 35, 39, 42–44, 53, 56, 57). The telerehabilitation sessions were focused on assessment, treatment techniques and exercises. Of the selected studies, six studies (33, 37, 46-50) used telephone as the format of the intervention. The focus of phone calls was on various topics, including wound care, pain management, and health assessment. In addition, phone calls were used to remind patients about their exercises and reinforce care behaviors. In three studies (38, 49, 52) patients were asked to watch educational videos related to their surgery, postoperative care and other topics. In two studies (40, 41) web-based interactions (e.g., remote viewing of x-ray images) were used to perform a follow-up meeting with the surgeon. Eisermann et al. (36) used computer-supported training. Three studies used mobile applications to deliver the training and education to patients (54, 59, 60). Finally, other studies used virtual reality training (55, 61) and the Vivofit 2 device (provides feedback on daily steps) (51, 58).



### **Risk of Bias**

We used the Cochrane risk of bias assessment tool (32) to assess the risk of bias of included studies. The main unmet criterion was not blinding the study personnel. Specifically, eight studies either fail to blind the personnel (35, 38, 40, 42, 49–51, 56, 57) or 11 did not provide sufficient information on blinding the personnel (33, 34, 36, 37, 45, 47, 48, 52–55, 58). Incomplete outcome data for all outcomes and other sources of bias (e.g., not reporting funding sources) were the other two unmet criteria. **Figure 2** provides the information related to the risk of bias assessment.

### Main Outcomes of the Included Studies Physical Outcomes

In 23 studies (33–37, 39, 42, 43, 46–55, 57–61) physical outcomes were measured. We categorized the outcome types into three main categories: (1) pain, (2) physical health and functioning,

TABLE 1 | Characteristics of the included studies.

References	Country	Sample size	Population	Time of intervention	Duration/length of follow up	Intervention
Russell et al. (34)	Australia	Intervention: 10 Control: 11	TKR	Post- operative	6 weeks	Intervention group: attended six 45-min telerehabilitation sessions. Control group: attended six 45-min in-person rehabilitation session. The focus of the sessions was on assessment, exercises an treatment techniques.
Kramer et al. (33)	Canada	Intervention: 80 Control: 80	TKR	Post- operative	52 weeks	Intervention group: received at least one phone call from the clinician between weeks 2 and 6 and another call from the clinician between weeks 7 and 12. The goal was to reinforce the critical role of doing exercise, wound care, scar treatmen and pain control. Control group: attended two physiotherapy sessions per week between weeks 2 and 12 after surgery.
Eisermann et al. (36)	Germany	Intervention: 79 Control: 70	TKR	Post- operative	24 weeks	Intervention group: used a computer-aided training, 3–5 time (30 min) for 2–4 weeks without any supervision. Control group: received self-training for 3–4 weeks.
Hørdam et al. (37)	Denmark	Intervention: 82 Control: 93	THR	Post- operative	36 weeks	Intervention group: 2 and 10 weeks after their surgery, received telephone support and counseling to understand patients' health status and their additional needs. Moreover, they received standard care. Control group: received the standard care (i.e., discharging from hospital 5–7 days after surgery and a clinical control 3 months after surgery).
Russell et al. (35)	Australia	Intervention: 31 Control: 34	TKR	Post- operative	6 weeks	Intervention group: attended six 45-min tele-rehabilitation sessions. Control group: attended six 45-min in-person rehabilitation session. The focus of the sessions was on assessment, exercises and treatment techniques.
Leal-Blanquet et al. (38)	Spain	Intervention: 42 Control: 50	TKR	Post- operative	48 weeks	Intervention group: watched a video related to surgery procedure, recovery, post-operative care, outpatients care at 2, 6, and 12 months after surgery. Control group: received verbal information about surgery, potential complications and habilitation.
Piqueras et al. (39)	Spain	Intervention: 90 Control: 91	TKR	Post- operative	12 weeks	Intervention group: received 10 1-h interactive virtual tele-rehabilitation sessions. The focus of the sessions was or exercises. Control group: received 10 1-h in-person rehabilitation sessions after surgery.
Marsh et al. (40)	Canada	Intervention: 66 Control: 61	THR	Post- operative	48 weeks	Intervention group: completed a web-based follow up with their surgeon. Control group: completed an in-person follow up with their surgeon.
Marsh et al. (41)	Canada	Intervention: 61 Control: 61	THR	Post- operative	48 weeks	Intervention group: completed a web-based follow up with their surgeon. Control group: completed an in-person follow up with their surgeon.
Mobolaji and Lynne (42)	UK	Intervention: 69 Control: 71	TKR	Post- operative	6 weeks	Intervention group: used a rehabilitation Visualization System for 6 weeks after surgery when exercising. In addition, patients received a video call 3 weeks after their surgery from a clinician and had a clinical checkup 6 weeks after surgery. Control group: received exercise handbook and DVD. In addition, patients received a checkup call 2 weeks after surgery.
Tousignant et al. (45)	Canada	Intervention: 97 Control: 100	TKR	Post- operative	16 weeks	Intervention group: received 16 telerehabilitation sessions via videoconferences led by a physiotherapist. Control group: received 16 in-person rehabilitation sessions led by a physiotherapist. The rehabilitation was focused on assessment and functional rehabilitation.

(Continued)

#### TABLE 1 | Continued

References	Country	Sample size	Population	Time of intervention	Duration/length of follow up	Intervention
Moffet et al. (43)	Canada	Intervention: 104 Control: 101	TKR	Post- operative	16 weeks	Intervention group: received 16 sessions of 45–60 min physiotherapy sessions <i>via</i> videoconference. Control group: received 16 45–60 min in-person physiotherapy sessions. The content of the sessions was on the assessment before and after exercise, supervised exercise training, and prescription of home exercises.
Szots et al. (48)	Denmark	Intervention: 54 Control: 54	TKR	Post- operative	12 weeks	Intervention group: received Telephone follow ups after their discharge (4 and 14 days) as well as standard care. The focus of Telephone Follow ups was on the patients' wound, pain management and exercises. Control group: received standard care.
Szots et al. (47)	Denmark	Intervention: 59 Control: 58	TKR	Post- operative	48 weeks	Intervention group: received Telephone follow ups after their discharge (4 and 14 days) as well as standard care. The focus of Telephone follow ups was on the patients' wound, pain management, and exercises. Control group: received standard care.
Chen et al. (46)	China	Intervention: 101 Control: 101	TKR	Post- operative	48 weeks	Intervention group: received a structured telephone call aiming to reinforce care and the standard care. Control group: received only the standard care with no telephone follow up. Standard care included home exercises, rehabilitation manua and a video.
Moffet et al. (44)	Canada	Intervention: 84 Control: 98	TKR	Post- operative	8 weeks	Intervention group: received 16 45–60 min telerehabilitation session (i.e., videoconference) Control group: received 16 45–60 min in-person physiotherapy session.
Bini and Mahajan (49)	USA	Intervention: 14 Control: 15	TKR	Post- operative	12 weeks	Intervention group: received instructional exercise videos. Patients received additional videos after clinicians reviewed patients uploaded videos of their exercise performance. The clinicians and the patients determined the endpoint. Control group: received the standard care and joining other patients in the rehabilitation clinic.
Park and Song (50)	South Korea	Telephone Intervention: 21 Text message group: 19	TKR	Post- operative	12 weeks	Intervention group: received telephone counseling 1, 3, 5, 7, 9, and 11 weeks after discharge. The focus of the calls was on general condition, activity of daily living, and affected joint dysfunction. Control group: patients in the text message group received telephone counseling 1, 3, 5, 7, 9, and 11 weeks after discharge. The focus of the text messages was on general condition, activity of daily living, and affected joint dysfunction
Culliton et al. (52)	Canada	Intervention: 209 Control: 207	TKR	Post- operative	48 weeks	Intervention group: watched 32 brief educational videos. The videos provided information on pain, functional outcomes, limitations, and restrictions. Control group: received an educational booklet before their surgery without any reminder.
Doiron-Cadrin et al. (53)	Canada	Interventions (2): 22 Control: 11	THA/TKA	Pre-operative	12 weeks	Intervention group 1: received a home-based prehabilitation program at home <i>via</i> telecommunication application. Intervention group 2: received a home-based prehabilitation program at the orthopedic clinic. Control group: received usual care without prehabilitation.
Hardt et al. (54)	Germany	Intervention: 26 Control: 27	ТКА	Post- operative	Days	Intervention group: received the GenuSport application. The application provided training for active knee extension. Patients also received the standard care. Control group: received the standard care. The standard care consisted of "knee mobilization, gait training, assisted walking with crutches, strength exercises, stair climbing, manual lymphatic drainage, and cryotherapy" three times a day.

(Continued)

TABLE 1 | Continued

References	Country	Sample size	Population	Time of intervention	Duration/length of follow up	Intervention
Jin et al. (55)	China	Intervention: 33 Control: 33	ТКА	Post- operative	24 weeks	Intervention group: received virtual reality training. In the VR training, patients rowed a boat three times a day (30 minutes/per session). Control group: performed knee flexion exercises using their arms. Exercises should perform 30 times a day (~30 min).
Van der Walt et al. (51)	Australia	Intervention: 81 Control: 82	TKA/THA	Post- operative	24 weeks	Intervention group: used the Vívofit 2 device. Patients were able to see their daily steps count. Control group: used the Vívofit 2 device. Patients were not able to see their daily steps count.
Timmers et al. (59)	Netherlands	Intervention group: 114 Control group: 99	TKR	Post- Operative	4 weeks	Intervention group: used the Patient Journey App to receive day-to-day information about recovery Control group: Received information related to recovery two times per week
Pronk et al. (60)	Netherlands	PainCoach: 38 Control group: 33	TKR	Post- Operative	4 weeks	Intervention group: downloaded the PainCoach app, that provides information on using pain medicine, exercises, rests, when to call the clinic and the usual care. Control group: only received the usual care in which they received the advice similar to the PainCoach advice.
Gianola et al. (61)	Italy	Intervention group: 44 Control group: 41	TKR	Post- Operative	6-7 days	Intervention group: received VR Based rehabilitation. Control group: received traditional rehabilitation.
Christiansen et al. (58)	USA	Intervention: 20 Control: 19	TKR	Post- operative	12 months	Intervention group: received a Fitbit Zip, received a recommended number of steps per week/day from a physiotherapist and received monthly follow up calls. Control group: received an exercise program and an exercise log. The physical therapist updated the record weekly.
Nelson et al. (56, 57)	Australia	Intervention: 35 Control: 35	THR	Post- Operative	6 months	Intervention group: received a telerehabilitation program and an exercise program using an iPad. Control group: received in-person physiotherapy and home exercise program.

and (3) functioning of the operated hip/knee. A total of 42 different types of measures and scales were used in the sample to assess physical outcomes. As an example, pain was measured using the Visual Analog Scale (62), pain subscale of the Knee Injury and Osteoarthritis Outcome Score (KOOS) (63), and the pain subscale of the Western Ontario and McMaster Universities Osteoarthritis Index (WOMAC) (64), Outcomes related to physical health and functioning were assessed by measures such as Knee Society Score (KSS) (65), Patient Specific Functional Scale (PSFS) (66), and Timed Up and Go Test (TUG) (67) and outcomes related to the functioning of the operated hip/knee were evaluated by measures including but not limited to Oxford Knee Score (OKS) (68), Biodex System-3 dynamometers, Nicholas Manual Muscle tester (NMMT) dynamometer (Kg), Angle degree, and Limb girth measurement. In total, 195 separate comparisons related to physical outcome types were reported. Of these 195 separate comparisons, in 169 comparisons, no statistically significant difference between the intervention and control groups was not observed. Only 28 comparisons [in 12 studies (34, 35, 46, 51, 52, 54, 55, 58-61)] showed a statistically significant difference between the intervention and control groups (i.e., patients who received the usual care) in the measured outcome at the study endpoint, in 25 cases, the intervention groups had significantly better outcomes. Specifically, in the pain category, in 36 comparisons there was only six significant differences between the intervention groups and the control groups [the intervention groups reported less pain in two studies (54, 55, 59, 60)].

In the physical health and functioning category, in 90 comparisons (out of 106), no significant differences were observed between the results of the intervention and the control groups. However, in 16 cases, the patients in the intervention groups showed statistically significant differences from the patients in the control group. For example, patients in the intervention group showed higher scores in overall functioning intervention types: Educational videos; VR intervention; six 45min tele-rehabilitation sessions; GenuSport application for the active knee extension; journey App (35, 54, 55, 59), exercises and physical activity, and mobility [intervention type: journey App; FitBit Zip; wearing the Vívofit 2 device; Virtual rehabilitation (39, 51, 58, 59)]. Only in three comparisons [intervention type: educational videos (52)], the control groups were found to have a higher score in functional activity (p = 0.02), less symptoms, and higher change in rigidity [Intervention type: VR based rehabilitation (61)], compared to the intervention group at the end of the study period.

In the functioning of the operated knee/hip category, in 47 comparisons (out of 53), no significant differences were found between the intervention and the control groups. However, in six comparisons related to overall functioning of the operated



joint [intervention types: GenuSport application for the active knee extension; VR intervention (54, 55)], muscle strength [intervention type: interactive virtual tele-rehabilitation sessions (39)], range of motion [intervention type: GenuSport application for the active knee extension; VR intervention (54, 55)], the patients in the intervention group scored better than the patients in the control groups. **Table 2** lists the details of the statistically significant analyses related to physical outcomes.

#### **Psychological and Social Outcomes**

In 19 studies (35, 37, 38, 42-44, 46-52, 54, 57, 59, 61, 71, 72) psychological and social outcomes were assessed. In total, 67 separate comparisons were conducted between the intervention and the control groups, using 16 distinct measures including Medical Outcomes Study Short Form (73), Short-Form questionnaire (74), Beck Depression Inventory (BDI) (75), Hospital Anxiety/Depression Scale (76), Pain Catastrophizing Scale (77), and Health Care Satisfaction Questionnaire (78). We categorized these outcomes in eight main psychological outcome types: (1) mental health, (2) intrinsic motivation, (3) self-efficacy, (4) emotional role, (5) quality of life, (6) patient satisfaction, (7) expectations, and (8) social function. Mental health was the most studied psychological outcome (n = 9 studies) (37, 42, 46–49, 51– 53). In 54 cases (out of 67), there were no statistically significant differences between the psychological or social outcomes of the eHealth intervention group and the control group, or both the intervention group and the control group had improved at the end of the study compared to baseline. However, for 12 discrete comparisons [in five studies (38, 46, 52, 57, 59)], there was a statistically significant difference between the intervention and the control group. For example, Culliton and colleagues [intervention type: educational videos (52)] reported that 12 months after surgery, patients in the eHealth intervention group had higher levels of anxiety (p = 0.02), pain-related magnification (p = 0.02), pain-related rumination (p = 0.02), and pain-related helplessness (p = 0.02) than the control group. However, Leal-Blanquet and colleagues [intervention type: educational video (38)] found that 4 months after surgery, the control group had statistically significant increased expectations of knee range of motion than the intervention group (p = 0.04), and the intervention group had significantly increased expectations for going up (p = 0.03) and down (p = 0.03) stairs than the control group participants. Table 3 lists the details of the statistically significant analyses related to psychosocial outcomes.

#### **Cost-Related Outcomes**

In seven studies (40–42, 45, 54, 56, 59) usability and costrelated outcomes were measured using 27 comparisons. In total, four main outcome types were assessed: (1) adherence to the rehab program, (2) travel distance, (3) time, and (4) cost. In 15 comparisons [in 5 studies (40, 41, 45, 56, 59)] the intervention group reported lower expenses and costs than the control groups. For example, Marsh and colleagues [intervention type: a webbased follow-up with the surgeon (40)] in their study found that travel distance to the medical facility (p < 0.01), travel TABLE 2 | Summary of the statistically significant physical outcomes assessed in the included studies.

Outcome type	References	Measure/scale	Subscale	Finding(s)
Pain	(54)	Numeric Rating Scale (NRS)	Pain in motion	Significant differences between the intervention group (mean = 4) and the control group (mean = 5) about 7 days after surgery ( $p = 0.006$ ; the intervention group reported lower pain).
Pain	(55)	Visual Analog Scale (62)	NA	Significant differences between the intervention group (mean = 3.87) and the control group (mean = 4.42) about 1 week after surgery ( $p = 0.002$ ; the intervention group expressed less pain).
Pain	(60)	Opiate use	Oxycodone usage	Significant differences after 2 weeks between the intervention group and the control group ( $p = 0.02$ ; Intervention group used 32.2% less opiate).
Pain	(59)	NRS	Pain during night	Significant difference between the intervention group (mean = 4.18) and the control group (mean = 5.21) 4 weeks after discharge ( $p = 0.003$ ; the intervention group reported less pain).
Pain	(59)	NRS	Pain at rest	Significant difference between the intervention group (mean = 3.45) and the control group (mean = 4.59) 4 weeks after discharge ( $\rho = 0.001$ ; the intervention group reported less pain).
Pain	(59)	NRS	Pain during activity	Significant difference between the intervention group (mean = $3.99$ ) and the control group (mean = $5.08$ ) 4 weeks after discharge ( $p < 0.001$ ; the intervention group reported less pain).
Physical health and functioning	(61)	Stabilometric platform of the Virtual Reality Rehabilitation System	Global proprioception	Significant difference in mean change (i.e., week 1 and week 4 after discharge) between the intervention (mean change = $73.46$ ) and the control group (mean change = $59.86$ ; $p = 0.002$ ; the intervention group had more improvement).
Physical health and functioning	(34)	Timed Up and Go Test (TUG) (67)	NA	There was a significant difference between the intervention group and the control group $p < 0.001$ ; the intervention group reported higher mean change after the end of the study). Means have not been reported.
Physical health and functioning	(46)	Days	Home exercise	Significant difference after 12 months between the intervention (mean = 78.35) and the control group (mean = 70.21; $p < 0.01$ ; the intervention group had a higher mean).
Physical health and functioning	(46)	Minutes	Home exercise	Significant difference after 12 months between the intervention (mean = 54.12) and the control group (mean = 48.95; $\rho < 0.01$ ; the intervention group had a higher mean).
Physical health and functioning	(52)	Knee Society Score (KSS) (65)	Symptoms	Significant difference between the intervention group (mean = 18.90) and the control group (mean = 19.84) 12 months after surgery ( $p = 0.04$ ; better outcome for the control group).
Physical health and functioning	(52)	KSS (65)	Functional activities	Significant difference between the intervention group (mean = 64.75) and the control group (mean = 68.18) 12 months after surgery ( $p = 0.04$ ; higher score in the functional activity in the control group).
Physical health and functioning	(35)	Patient Specific Functional Scale (PSFS) (66)	NA	Significant difference in mean change (baseline to 1.5 months after surgery) between the intervention group (mean = 5.05) and the control group (mean = 3.97; $\rho$ = 0.04; intervention group showed higher score in functioning).
Physical health and functioning	(54)	KSS (65)	Function	Significant differences between the intervention group (mean = 42) and the control group (mean = 37) about 7 days after surgery ( $p = 0.01$ ; the intervention group reported higher score in functioning).
Physical health and functioning	(51)	Daily Step Count	Mobility	Significant differences between the intervention group (mean = 137) and the control group (mean = 117) 6 months after surgery ( $p = 0.030$ ; the intervention had more daily steps).
Physical health and functioning	(55)	WOMAC (64)	NA	Significant differences between the intervention group (mean = 21.58) and the control group (mean = 26.33) about 6 months after surgery ( $p = 0.000$ ; the intervention group had less functional limitations).
Physical health and functioning	(59)	OA KOOS-Physical Function Shortform (KOOS-PS) (69)	Functioning limitation	Significant difference at week 4 between the intervention (mean = 37.61) and the control group (mean = 43.08; $\rho < 0.001$ ; the intervention group reported less functioning limitation).
Physical health and functioning	(59)	EuroQol-5 Dimensions	NA	Significant difference at week 4 between the intervention group (mean = 0.76) and the control group (mean = 0.67; $\rho < 0.001$ ; the intervention group reported more improvement).
Physical health and functioning	(59)	NRS	Physiotherapy exercises	Significant difference in mean change between the intervention (mean = 7.50) and the control group (mean = 6.88) 4 weeks after discharge ( $p = 0.03$ ; the intervention reported higher ability to perform physical exercises).

(Continued)

TABLE 2 | Continued

Outcome type	References	Measure/scale	Subscale	Finding(s)
Physical health and functioning	(61)	WOMAC (64)	Rigidity	Significant difference in mean change (i.e., week 1 and week 4 after discharge) between the intervention (mean change = $-45.43$ ) and the control group (mean change = $-67.05$ ; $\rho = 0.046$ ; the intervention reported less change in rigidity).
Physical health and functioning	(58)	Minutes	Physical activity	Significant difference at 12 months between the intervention (mean = 133.8) and the control group (mean = $57.7$ ) 12 months after surgery (95% CI: 10.5, 141.5; the intervention group had a higher mean in physical activity).
Physical health and functioning	(58)	Daily steps	Physical activity	Significant difference between the intervention (mean $=$ 6,144) and the control group (mean $=$ 4,169) 12 months after surgery (95% CI: 466, 3,422; the intervention group had a reported more daily stems).
Functioning of the Operated Knee/Hip	(55)	Hospital for Special Surgery (HSS hip) (70)	NA	Significant differences between the intervention group (mean = 87.55) and the control group (mean = 80.39) about 6 months after surgery ( $p = 0.000$ ; the intervention group had better score in functioning).
Functioning of the Operated Knee/Hip	(46)	Angle degree	Active range of motion	Significant difference between the intervention group and the control group ( $p = 0.01$ ; the intervention group had a higher mean change after 12 months).
Functioning of the Operated Knee/Hip	(39)	Nicholas Manual Muscle tester (NMMT) dynamometer (Kg)	Quadriceps muscle strength	Significantly increased in the intervention group (mean change = 8.48) compared to the control group (mean change = 5.89) 3 months after surgery ( $\rho = 0.018$ ).
Functioning of the Operated Knee/Hip	(54)	KSS (65)	Knee	Significant differences between the intervention group (mean = 71) and the control group (mean = 59) about 7 days after surgery ( $p = 0.0002$ ; intervention group had better functioning).
Functioning of the Operated Knee/Hip	(54)	Angle degrees	Active range of motion	Significant differences between the intervention group (mean = 76) and the control group (mean = 67) about 7 days after surgery ( $p = 0.034$ ; intervention group had better active range of motion).
Functioning of the Operated Knee/Hip	(55)	Angle degree	Knee range of motion	Significant differences the intervention group (mean = 93.73) and the control group (mean = 86.36) about 2 weeks after surgery between ( $\rho = 0.000$ ; intervention group had higher range of motion).

costs (p < 0.01), time spent completing the follow-up assessment (p < 0.01) and time spent by caregivers (p < 0.01) were lower in the eHealth intervention group than the control group. Similarly, Nelson et al. (56) reported a statistically significant difference in patient/carer travel costs between the intervention group (mean = \$0.77) and control group (mean = \$77.69). In another study, Marsh and colleagues [intervention type: a web-based follow-up with the surgeon (41)] showed that the cost of the follow-up assessment based on both the societal (p < 0.01) and the health-care payer perspectives (p < 0.01) was lower in the intervention group than the control group.

**Table 4** lists the details of the statistically significant analyses

 related to usability and cost-related outcomes.

### DISCUSSION

This review investigated the effectiveness of eHealth tools in providing pre- and post-operative education for patients undergoing total hip and knee arthroplasty. Based on the type of analyses that have been conducted, the findings showed that often both the intervention group and the control groups show statistically significant improvements at the end of the study period compared to baseline. Overall, there were no major differences between the intervention and the control groups at the end of the study period indicating that eHealth tools are as effective as usual care.

Findings from this review uncovered a few significant benefits of eHealth tools when physical or psychological and social outcomes were considered. These results are consistent with other studies in the field of surgery that have generally found no statistically significant differences in the outcomes of eHealth tools in postoperative care vs. traditional or usual care interventions (80-83). A systematic review focused on telemedicine conducted by Grunter et al. (84) demonstrated that complication rates after surgery do not differ between eHealth intervention groups and control groups in various patient populations. However, our findings are at odds with other work that has found eHealth tools to be more effective than usual care in other contexts, for example, in improving physical activities in older patients [e.g., (85)]. In addition, in one study (52), researchers reported slightly higher levels of anxiety, pain-related rumination, magnification, and helplessness in the intervention group compared to the control group. While these may be considered minor and non-clinically relevant findings, it is important to conduct more research on the potential benefits and the harms of eHealth tools.

While costs and expenses have been measured only in few studies (40–42, 45), in most cases, eHealth tools were found to be more cost-effective than usual care, making lower-cost a key advantage of eHealth tools over usual care. In line with this finding, Hwa et al. (86) found that using telephone follow-ups can lead to 110 additional opening spots in their clinics. Besides

Outcome type	References	Measure/scale	Sub-scale	Finding(s)
Mental health	(46)	Medical Outcomes Study Short Form (SF-36) (73)	Mental component summary	Significant difference between the intervention (mean T0 = 45.6; mean T3 = 52.5) and the control group (mean T0 = 44.7; mean T3 = 50.5; $\rho$ = 0.03 the intervention group had a bigger change).
Mental health	(52)	Hospital Anxiety and Depression Scale (76)	Anxiety	Significantly more anxiety in the intervention group (mean = 3.40) compared to the control group (mean = 2.81) 12 months after surgery ( $\rho$ = 0.02).
Mental health	(52)	Pain Catastrophizing Scale (PCS) (77)	Rumination	Significantly more rumination in the intervention group (mean = 2.16) compared to the control group (mean = 1.51) 12 months after surgery ( $p = 0.02$ ).
Mental health	(52)	PCS (77)	Magnification	Significantly more magnification in the intervention group (mean = 1.03) compared to the control group (mean = 0.69) 12 months after surgery ( $p = 0.02$ ).
Mental health	(52)	PCS (77)	Helplessness	Significantly more helplessness in the intervention group (mean = 2.32) compared to the control group (mean = 1.76) 12 months after surgery ( $p = 0.02$ ).
Quality of life	(59)	Numerical Rating Scale (NRS) (0–10)	Daily self-care	Significant difference between the intervention (mean = $8.32$ ) and the control group (7.64) 4 weeks after discharge ( $p = 0.004$ ; the intervention group reporte higher daily self-care).
Quality of life	(59)	NRS (0–10)	Satisfaction with information	Significant difference between the intervention (mean = 7.61) and the control group (mean = 5.32) 4 weeks after discharge ( $p < 0.001$ ; the intervention group were more satisfied).
Quality of life	(59)	NRS (0–10)	Patient-perceived involvement by the hospital	Significant difference between the intervention (mean = 7.24) and the control group (mean = 4.90) 4 weeks after discharge ( $p < 0.001$ ; the intervention group perceived more involvement).
Satisfaction	(57)	Questionnaire	Home exercise compliance	The intervention group (overall compliance=86%) reported higher compliant that the control group (overall compliance=74; $p = 0.048$ ) after surgery.
Expectations	(38)	The Hospital for Special Surgery Knee Replacement Expectations Surgery (KRES) (79)	Knee range of motion	The intervention group (mean of the change = 0.0) had significantly decreased expectations of knee ROM in comparison to the control group (mean of the change = 0.1) 4 months after surgery ( $p = 0.04$ ).
Expectations	(38)	KRES (79)	Going up stairs	The intervention group (mean of the change = 0.1) had significantly increased expectations for going up stairs than the control group (mean of the change = $-0.04$ ) 4 months after surgery ( $\rho = 0.03$ ).
Expectations	(38)	KRES (79)	Going down the stairs	The intervention group (mean of the change = 0.2) had significantly increased expectations for going down the stairs than the control group (mean of the change = $-0.02$ ) 4 months after surgery ( $p = 0.03$ ).

TABLE 3 | Summary of the psychological and social outcomes assessed in the included studies.

the health care systems' benefits, patients who use eHealth tools and their family members can also benefit financially by traveling fewer kilometers, dedicating less time for travel, reducing the amount of time they take off from their work and decreasing money spent on transit (82, 87).

Despite broad inclusion criteria that encompassed prehab interventions, our search strategy only uncovered one study where eHealth tools were used before surgery. More research is needed to understand better the effect of eHealth tools that aim to deliver prehab education. In addition, most of the research included in this review did not meet all the criteria for high-quality studies. While character limitations and journal requirements may play a role in the quality of the papers, still the findings should be considered with caution and may not represent the actual impact of eHealth tools. The overrepresentation of studies focused on using eHealth tools on patients with knee arthroplasties over hip arthroplasties also limits the generalizability of the impacts uncovered. Similarly, our sample contained a majority of studies assessing the impact of eHealth tools designed to deliver exercises and physiotherapy and as such, physical outcomes were the main focus. This narrow focus highlights the need to consider the potential of eHealth tools in promoting a holistic view of both prehab and rehab, which includes attention to psychosocial factors. Furthermore, our sample mainly contained studies that did not fully embrace contemporary eHealth approaches (e.g., interactive designs). Therefore, it is possible that using interactive eHealth tools can improve eHealth benefits from usual care. While we did not restrict our search to any language, but we had to exclude studies that were not in English (two studies) due to our limited resources. Qualitative studies were excluded from this study. Therefore, potential benefits and harms that were uncovered through qualitative work have not been captured in this work. Furthermore, because of the lack of evidence about the most common outcomes measured in the studies related to eHealth tools and hip and knee arthroplasties, our team decided not to select any primary or secondary outcome measures. Finally, due to the heterogeneity of the outcome and the measurements, we did not perform a meta-analysis. The absence of meta-analyses will limit our ability to estimate TABLE 4 | Summary of the statistically significant cost-related outcomes assessed in the included studies.

Outcome type	References	Measure/scale	Subscale	Finding(s)
Travel distance	(40)	Kilometers	Travel distance	Travel distance to the radiology appointment was significantly lower for the intervention group (mean = 28.2) compared to the travel distance to the radiographs and clinic visits (mean = 103.7) for the control group about 12 months after surgery ( $p < 0.01$ ).
Time	(40)	Minutes	Time to complete	Completion time for follow-up assessment was significantly less for the intervention group (mean = 121.7) than the control group (mean = 228.8) about 12 months after surgery ( $\rho < 0.01$ ).
Time	(40)	Minutes	Caregiver time	Caregiver time for follow-up assessment was significantly less for the intervention group (mean = 44.1) than the control group (mean = 127.2) about 12 months after surgery ( $p < 0.01$ ).
Time	(56)	Minutes	Patient time	Significant difference in patient time between the intervention group (mean $=$ 79) and the control group (mean $=$ 331) ( $\rho$ -value not reported); intervention group spent less patient time.
Time	(56)	Minutes	Carer time	Significant difference in carer time between the intervention group (mean $=$ 38) and the control group (mean $=$ 302) ( $p$ -value not reported); intervention group spent less carer time.
Costs and expenses	(40)	Dollar	Travel cost	Travel costs were significantly lower for the intervention group (mean = 10.45) compared to the control group (21.36) about 12 months after surgery ( $p < 0.01$ ).
Costs and expenses	(41)	Dollar	Follow up appointment cost based on societal perspective	Mean cost of follow-up per patient was significantly lower for the intervention group (mean = CDN\$98) compared to the control group (mean = CDN\$162) about 12 months after surgery ( $p < 0.01$ ).
Costs and expenses	(41)	Dollar	Follow up appointment cost based on health-care payer perspective	Mean cost of follow-up per patient was significantly lower for the intervention group (mean = CDN\$45) compared to the control group (mean = CDN\$71) about 12 months after surgery ( $p < 0.01$ ).
Costs and expenses	(41)	Dollar	Software Cost (i.e., Licensing Fee)	The intervention group would be significantly cost-saving when a surgeon follow-up 20 or more patient per year using this method compared to the control group from 12 to 24 months after surgery ( $\rho < 0.01$ ).
Costs and expenses	(41)	Dollar	Value of unpaid time for (\$0/h, \$10.25/h, \$26.19/h)	The intervention group was significantly more cost-saving if the value of the unpaid time was \$0/h ( $\rho < 0.01$ ) or \$10.25/h ( $\rho < 0.01$ ) or even \$26.19/h ( $\rho < 0.01$ ) compared to the control group.
Costs and expenses	(45)	Dollar	Total cost for treatment (includes received and canceled treatments)	Total cost was significantly lower for the patients in the intervention group (mean = \$1,224) than the patients in the control group (mean = \$1,487; $\rho < 0.001$ ).
Costs and expenses	(45)	Dollar	Cost for one treatment	Cost per treatment was significantly lower for the patients in the intervention group (mean = \$80.99) than the patients in the control group (mean = \$93.08; $\rho = 0.008$ ).
Costs and expenses	(59)	Researcher made question	Point of contact with hospital	Significant difference in overall point of contact between the intervention (mean = 1.22) and the control group (mean = 1.62; $p = 0.014$ ; the intervention group had fewer contacts with the hospital).
Costs and expenses	(56)	Number	Intervention physiotherapy sessions	Significant difference between intervention group (mean = 2.5) and control group (mean = 2.9; $p$ -value not reported); intervention group had less sessions.
Costs and expenses	(56)	Dollar	Patient/Carer travel costs	Significant difference in patient/carer travel costs between the intervention group (mean = $0.77$ ) and control group (mean = $77.69$ ; $\rho$ -value not reported); intervention group had less patient/carer travel costs.

the effect size of eHealth interventions. It will also limit our ability to generalize our findings (88). In summary, findings from this review revealed that eHealth tools are as effective as usual care interventions and may be more cost-effective in their implementation. The use of eHealth intervention requires attention to several factors. Patient preferences and computer literacy levels are critical to the success of remote interventions, especially when using interactive designs. While for some eHealth modalities such as telephone follow-ups, high computer or literacy skills are not critical, other interventions delivered through apps and computer programs require not only access to these devices but also comfort in using them. Another important issue raised by researchers in this field (84, 89) is patients' privacy when using eHealth tools. For example, Watzlaf et al. (90) found that most voices over internet protocol (VOIP) videoconferencing software that is used for videoconferencing have serious security vulnerabilities. Moving forward, attention to ethical issues such as privacy, confidentiality and quality are crucial in ensuring eHealth tools are both adopted and beneficial (91).

#### Conclusion

The overarching goal of the current study was to provide an understanding of the effectiveness of eHealth tools on the outcomes of patients undergoing total hip and knee arthroplasty. While the included studies used heterogeneous group of interventions, in conclusion, the overall findings showed that regardless of the type of the eHealth intervention that was used in each study, in the majority of the cases, eHealth tools were as effective as usual care interventions, but more cost-effective which can be a good argument in supporting their development and application in the health care system. However, only a smaller set of studies investigated the cost-related outcomes, and more investigations, especially longitudinal investigations, are needed to assess the short- and long-term impacts of eHealth tools on cost-related outcomes. The findings of this study do not indicate that using eHealth tools will totally remove the costs of prehabilitation and rehabilitation intervention. However, it shows that eHealth tools can significantly reduce some of the expenses (e.g., travel time). Considering the evidence around the procedures for tool development that suggests eHealth tool should contain personalized advice, have features that enable communications between patients and their health care providers, and include patients' health profile (92) to increase the effectiveness of the eHealth tools, more robust approaches in developing these tools should be taken into account in future. Furthermore, more research should compare the effect of eHealth education with standard care. Especially usability and feasibility of different aspects and features of eHealth education (e.g., videos, text, quizzes) should be assessed. In addition, the effectiveness of the different types of eHealth education (e.g., webinars, online applications with or without professional

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support) should be compared with in-person education. While the COVID-19 pandemic restricted many from accessing inperson education, it also mitigated the transformation of inperson education to online education, an opportunity that should not be missed (93).

### **Clinical Outcomes**

- Using eHealth tools in providing health-related education for patients undergoing hip/knee arthroplasty can be as effective as usual care.
- eHealth tools are more cost-effective than usual care.

## DATA AVAILABILITY STATEMENT

The original contributions generated for the study are included in the article/**Supplementary Materials**, further inquiries can be directed to the corresponding author.

## **AUTHOR CONTRIBUTIONS**

SM: study design, databases search, study selection, quality assessment, and manuscript preparation. WCM: supervision, study design, and manuscript preparation. JW: data extraction, manuscript preparation, and quality assessment. CP: databases search, quality assessment, and manuscript preparation. JMR: supervision, study design, study selection, quality assessment, and manuscript preparation. All authors read and approved the final manuscript.

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### SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fresc. 2021.696019/full#supplementary-material

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