



Outpatient Total Hip Arthroplasty: A Meta-Analysis

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Abstract: Introduction: Outpatient total hip arthroplasty (THA) is increasingly popular. This meta-analysis investigated the potential advantages of outpatient regimes for THA. Methods: This study followed the PRISMA guidelines. PubMed, Web of Science, Google Scholar, Embase, and Scopus databases were accessed in June 2021. All clinical studies investigating outpatient THA were considered. The outcomes of interest were pain, infection, mortality, revision, dislocation, readmission rates, and deep vein thrombosis (DVT). Results: Data from 102,839 patients were included. A total of 52% (153,168 of 102,839 patients) were women. The mean age of patients was 62.6 \pm 4.6 years, the mean BMI was 29.1 \pm 1.8 kg/m². Good comparability was found in age, BMI, and gender (p > 0.1). No difference was found in pain (p = 0.4), infections (p = 0.9), mortality (p = 0.9), rate of revision (p = 0.1), dislocation (p = 0.9), and readmission (p = 0.8). The outpatient group demonstrated a greater rate of DVT (OR 3.57; 95% CI 2.47 to 5.18; *p* < 0.0001). **Conclusions:** In selected patients, outpatient THA can be performed safely with optimal outcomes comparable with inpatient THA. Clear and comprehensive pre-operative planning should involve a multi-disciplinary group composed of orthopaedic surgeons, anaesthesia and rehabilitation specialists, and physiotherapists. Each centre performing outpatient THA should implement continuous homecoming welfare activity, to supervise physiotherapy and monitor anticoagulant therapy.

Keywords: total hip arthroplasty; inpatients; outpatients

1. Introduction

Total hip arthroplasty (THA) is the treatment choice for advanced hip osteoarthritis. THA has typically been considered as an inpatient procedure [1,2]. Postoperative care after THA has evolved during the past 40 years [3]. Initially, patients undergoing THA could have been hospitalized even longer than two weeks and spend further weeks in a rehabilitation centre [4]. Weightbearing was limited and mobilisation delayed [4]. Postoperative protocols have evolved to decrease costs and allow faster discharge and return to previous daily living activities [5–9]. Improvement in pain management and bleeding control, regional anaesthesia, direct postoperative full weightbearing, and fast track concepts in hospitalization and rehabilitation have been introduced [1,5,10–12]. Long hospitalisation has been associated with a higher rate of complications [13,14]. However, outpatient THA remains reserved to a few selected patients: less than 1% of THAs in the United States are performed as outpatient procedures [15–17]. A patient's selection is mandatory. The current exclusion criteria for outpatient THA includes heart and vascular



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Copyright: © 2021 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (https:// creativecommons.org/licenses/by/ 4.0/). diseases, a history of transient ischemic attack, chronic obstructive pulmonary disease, untreated obstructive sleep apnoea, obesity, haemoglobin < 130 g/L, diabetes mellitus, end-stage hepatic and/or renal disease, history of delirium or dementia, solid organ transplant, and advanced neoplasm [18,19].

Several clinical studies comparing outpatient versus inpatient THA (>2 days) are available [1,11,13,14,17,20–30]. However, most of them have been recently published, and, therefore, have not yet been considered in previous systematic reviews and meta-analyses. This meta-analysis compared outpatient versus inpatient THA (>2 days) in terms of symptoms, rates of infection, mortality, revision, dislocation, readmission, and deep vein thrombosis (DVT). We hypothesised that, for selected patients, outpatient THA provides comparable results to inpatient procedures.

2. Material and Methods

2.1. Search Strategy

This systematic review was conducted according to the Preferred Reporting Items for Systematic Reviews and Meta-Analyses: the 2020 PRISMA statement [31]. The PICO algorithm guided the initial search:

- P (Population): end stage hip osteoarthritis;
- I (Intervention): outpatient THA;
- C (Comparison): inpatient (>2 days) THA;
- O (Outcomes): pain, infections, mortality, revisions, dislocations, readmissions, DVT.

2.2. Data Source and Extraction

The literature search was performed by two authors (F.M. and L.C.) independently. In June 2021, the main databases were accessed: PubMed, EMBASE, Google Scholar, and Scopus. The following keywords were used in combination: *one day, day surgery, outpatient, inpatient, fast track, arthroplasty, hip, prosthesis, replacement, readmission, revision, stay, hospitalization, deep vein thrombosis, complications, failure.* The resulting articles were examined and, if of interest, the full text version was accessed. The bibliographies were also screened for inclusion of further articles. Any disagreements were discussed and settled by consensus.

2.3. Eligibility Criteria

All clinical trials comparing one outpatient versus inpatient (>2 days) THA were considered. Only study comparative investigations published in peer reviewed journals were considered. According to the authors' language capabilities, articles in English, German, Italian, French, and Spanish were eligible. According to Oxford Centre of Evidence-Based Medicine [32], levels I to IV of evidence were considered. Reviews, opinions, letters, and editorials were not considered. Animals, biomechanics, computational, and cadaveric studies were not eligible. Only articles which reported quantitative data with regard to symptoms and complications were eligible. Studies which enhanced THA with cell therapies or with experimental surgical or rehabilitative protocols were not suitable, nor where those experimenting new implants designs.

2.4. Selection and Data Collection

Two authors (F.M. and L.C.) independently performed the database search. All the resulting titles were screened and, if suitable, the abstract was accessed. The full text of abstracts matching the topic were accessed. A cross reference of the bibliography of full-text articles were also screened for inclusion. Disagreements were debated and the final decision was decided by a third author (NM).

2.5. Data Items

Two authors (F.M. and L.C.) independently performed data extraction. The following data were extracted: generalities (author, year, journal, type of study) and patient baseline

demographic information (number of samples, sex, mean age and BMI, length of the hospital stay) were collected. The outcomes of interest were: pain, rates of infection, mortality, revision, dislocation, readmission, and deep vein thrombosis (DVT).

2.6. Methodology Quality Assessment

The Coleman Methodology Score (CMS) was used to evaluate the quality of the methodology of each investigation. The CMS is widely used to evaluate the methodological quality of scientific articles for systematic reviews and meta-analyses [33]. The CMS evaluates the study size, length of the follow-up, type of study, diagnosis description, surgical technique, and rehabilitation. Additionally, outcome criteria assessment, procedures for assessing outcomes, and the subject selection process were also evaluated. The CMS rated articles with values between 0 (poor) and 100 (excellent). A mean value greater than 60 points was considered satisfactory.

2.7. Statistical Analysis

Statistical analyses were performed by the main author (F.M.). These meta-analyses were performed using the Review Manager Software 5.3 (The Nordic Cochrane Collaboration, Copenhagen). Binary data were evaluated through a Mantel–Haenszel analysis, with odd ratio (OR) effect measure. The comparisons were performed with a fixed model effect as set up. Heterogeneity was assessed through the χ^2 and Higgins-I² test. If $\chi^2 < 0.05$ and If I² test > 50%, high heterogeneity was detected. In cases of heterogeneity, a random model effect was used. The confidence intervals (CI) were set at 95% in all comparisons. The overall effect was considered statistically significant if p < 0.05. The funnel plot of the most reported outcome was performed to assess the risk of publication bias. Egger's linear regression was performed through the STATA/MP Software version 16 (StataCorp, College Station, TX, USA) to assess funnel plot asymmetry, with values of p < 0.05 indicating statistically significant asymmetry.

3. Results

3.1. Search Result

A total of 147 articles were identified during the initial search. Of these, 35 duplicates were excluded. An additional 72 articles were excluded because of the following reasons: not comparative study (n = 31), type of study (n = 35), uncertain methodology or results (n = 5), and language limitation (n = 1). An additional 24 articles were excluded as they lacked quantitative data under the outcomes of interest. This left 16 studies for inclusion (Figure 1).

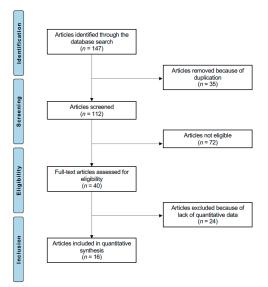


Figure 1. Literature search flow chart.

3.2. Methodological Quality Assessment

The study size and the length of the follow-up were reliable in most of considered articles. The surgical technique, diagnosis, and rehabilitation protocols were generally well described. The retrospective design of 50% (8/16) of the included studies represented an important limitation highlighted by the CMS. Outcome measures and the assessment timing were often well defined, providing moderate reliability. Concluding, CMS resulted in 72.4 \pm 5.5, attesting to the present study a good quality of the methodological assessment (Table 1).

Table 1. Methodological quality assessment.

Endpoint	Mean	SD	Range			
Part A: Only one score to be given for each of the 7 sections						
Study size: number of patients	9.43	1.63	4 to 10			
Mean follow-up	0.25	1.0	0 to 4			
Surgical approach	8.25	2.64	0 to 10			
Type of study	5.31	5.61	0 to 15			
Description of diagnosis	4.37	1.70	0 to 5			
Descriptions of surgical technique	6.94	3.48	0 to 10			
Description of postoperative rehabilitation	3.12	2.5	0 to 5			
Part B: Scores may be given for each option in each o	of the 3 sections					
Outcome criteria	2.5	0.5	2 to 3			
Procedure of assessing outcomes	3.74	0.84	3 to 5			
Description of subject selection process	4.83	0.88	0 to 5			

3.3. Risk of Publication Bias

The funnel plot of most reported outcome was performed to assess the risk of publication bias (Figure 2). The graph evidenced minimal asymmetry. Most of effects points were located in the pyramidal shape. Moreover, the Egger's test evidenced no statistically significant asymmetry (p = 0.08). Concluding, the funnel plot indicated a low to moderate risk of publication bias.

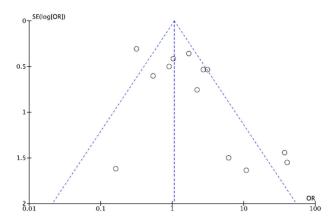


Figure 2. Funnel plot.

3.4. Patient Demographic

Data from 102,839 patients were included. A total of 52% (53,168 of 102,839 patients) were women. The mean age of patients was 62.6 ± 4.6 years, and the mean BMI was 29.1 ± 1.8 kg/m². A good between-groups comparability was found in age, BMI, and the female:male ratio (p > 0.1). An overview of the included studies is shown in Table 2.

3.5. Outcomes of Interest

No difference was found in pain (p = 0.4), infections (p = 0.9), mortality (p = 0.9), revision (p = 0.1), dislocation (p = 0.9), and readmission (p = 0.8) between the two groups. The outpatient group demonstrated a greater rate of DVT (OR 3.57; 95% CI 2.47 to 5.18; p < 0.0001). These results are shown in greater detail in Figure 3.

Author et al., Year	Journal	Study Design	CMS	Length of Stay	Patients (n)	Age	Female (%)	BMI (kg/m ²
Aynardi et al., 2014 [11]	HSS J.	Prospective	80	1 Day >2 Days	119 78	59.0 61.5	40 67	28.1 33.2
Bertin et al., 2005 [1]	Clin. Orthop. Relat. Res.	Retrospective	69	1 Day >2 Days	10 10	62.0 63.0	40 50	30.0 29.6
Carey et al., 2019 [20]	J. Arthroplast.	Retrospective	70	1 Day >2 Days	623 1869	61.0 61.0		
Crampet et al., 2019 [21]	Orthop. Traumatol. Surg. Res.	Retrospective	72	1 Day >2 Days	50 77	62.7 70.5	44 81	25.6 26.5
Goyal et al., 2017 [22]	Clin. Orthop. Relat. Res.	RCT	85	1 Day >2 Days	112 108	59.8 60.2	47 46	27.6 28.3
Greenky et al., 2019 [23]	J. Arthroplast.	Retrospective	75	1 Day >2 Days	310 28,408	71.4 73.8		29.6 29.2
Gromov et al., 2019 [17]	Acta Orthop.	Prospective	75	1 Day >2 Days	70 339	61.0 62.0	41 43	28.0 28.0
Kelly et al., 2018 [24]	J. Arthroplast.	Prospective	74	1 Day >2 Days	23 28	59.2 64.1	58 28	30.4 32.7
Krenk et al., 2014 [25]	Anesth. Analg.	Prospective	67	1 Day >2 Days	220 220	72.0 68.0	9 91	
Lovecchio et al., 2016 [13]	J. Arthroplast.	Prospective	77	1 Day >2 Days	183 585	64.0 65.0	56 55	30.0 29.0
Nelson 2016 et al., 2016 [26]	J. Arthroplast.	Retrospective	70	1 Day >2 Days	420 63,424	62.0 65.0	47 55	28.0 29.0
Otero et al., 2016 [14]	J. Arthroplast.	Prospective	70	1 Day >2 Days	249 1940	62.6 59.9	50 38	29.4 29.4
Richards et al., 2018 [27]	J. Arthroplast.	Prospective	68	1 Day >2 Days	136 396	53.2 55.4	71 54	27.5 29.6
Tedder et al., 2018 [28]	J. Foot Ankle Surg.	Retrospective	75	1 Day >2 Days	66 535	58.9 64.0	58 49	29.0 30.0
Weiser et al., 2018 [29]	J. Arthroplast.	Retrospective	70	1 Day >2 Days	164 1315	56.8 58.0	52 57	26.9 28.2
Ziemba-Davis et al., 2019 [30]	J. Arthroplast.	Retrospective	62	1 Day >2 Days	164 588	63.4 63.0	61 60	32.6 30.1

 Table 2. Studies generalities and patient demographics (CMS: Coleman Methodology Score).

Study or Subgroup W		Odds Ratio H, Fixed, 95% CI	Odds Ratio M-H, Fixed, 95% Cl				
, , ,		0.60 [0.11, 3.22]					
		5 [0.24, 105.71]					
Kielik et al., 2014	11.0/0 5.0	5 [0.24, 105.71]					
Total (95% CI) 10	00.0%	L.11 [0.30, 4.10]					
Total events							
Heterogeneity: Chi ² = 1.4	7 df = 1 (0)	$2 = 0.22 + 1^2 = 22\%$					
Test for overall effect: Z =			0.01 0.1 1 10 100				
rest for overall effect. z =	0.16 (P =	0.87)	Favours 1 Day Favours >2 Days				
Forest plot of the outcome: dislocation							
10	lest pi						
		Odds Ratio	Odds Ratio				
Study or Subgroup	Weight	M-H, Fixed, 95% CI					
Avnardi et al., 2014	2.5%	5.21 [0.28, 98,13]					
Crampet et al., 2019	4.8%	0.50 [0.02, 12.64]					
Gromov et al., 2019	32.8%	5.62 [3.17, 9.96]					
Krenk et al., 2014	8.1%	1.00 [0.14, 7.16]					
Lovecchio et al., 2016	24.6%	4.45 [2.17, 9.14]					
Nelson 2016., 2016	17.1%	0.95 [0.23, 3.81]					
Otero et al., 2016	9.2%	1.56 [0.34, 7.17]					
Ziemba-Davis et al., 2019	0.9%	10.80 [0.44, 266.31]					
Total (95% CI)	100.0%	3.57 [2.47, 5.18]					
Total (95% CI) Total events	100.0%	3.57 [2.47, 5.18]					
Total (95% CI) Total events Heterogeneity: Chi ² = 10.9	100.0%	3.57 [2.47, 5.18] P = 0.14); I ² = 36%	•				
Total (95% CI) Total events	100.0%	3.57 [2.47, 5.18] P = 0.14); I ² = 36%					
Total (95% CI) Total events Heterogeneity: Chi ² = 10.9 Test for overall effect: Z =	100.0% 94, df = 7 (F 6.75 (P < 0	3.57 [2.47, 5.18] P = 0.14); I ² = 36%	0.01 0.1 1 10 100 Favours 1 Day Favours >2 Days				
Total (95% CI) Total events Heterogeneity: Chi ² = 10.9 Test for overall effect: Z =	100.0% 94, df = 7 (F 6.75 (P < 0	3.57 [2.47, 5.18] P = 0.14); I ² = 36%					
Total (95% CI) Total events Heterogeneity: Chi ² = 10.9 Test for overall effect: Z =	100.0% 94, df = 7 (F 6.75 (P < 0	3.57 [2.47, 5.18] P = 0.14); I ² = 36%	0.01 0.1 1 10 100 Favours 1 Day Favours > 2 Days				
Total (95% CI) Total events Heterogeneity: Chi ² = 10.9 Test for overall effect: Z =	100.0% 94, df = 7 (F 6.75 (P < 0	3.57 [2.47, 5.18] P = 0.14); I ² = 36%	0.01 0.1 1 10 100 Favours 1 Day Favours > 2 Days				
Total (95% CI) Total events Heterogeneity: Chi ² = 10.9 Test for overall effect: Z =	100.0% 94, df = 7 (F 6.75 (P < 0	3.57 [2.47, 5.18] P = 0.14); I ² = 36%	0.01 0.1 1 10 100 Favours 1 Day Favours > 2 Days				
Total (95% CI) Total events Heterogeneity: Chi ² = 10.9 Test for overall effect: Z =	100.0% 94, df = 7 (F 6.75 (P < 0	3.57 [2.47, 5.18] P = 0.14); I ² = 36%	0.01 0.1 1 10 100 Favours 1 Day Favours > 2 Days				
Total (95% CI) Total events Heterogeneity: Chi ² = 10.9 Test for overall effect: Z = Forest p	100.0% 94, df = 7 (F 6.75 (P < 0	3.57 [2.47, 5.18] 9 = 0.14); 1 ² = 36% 1.00001) he outcome:	0.01 0.1 10 Favours 1 Day Favours >2 Days deep vein thrombosis Odds Ratio				
Total (95% CI) Total events Heterogeneity: Chi ² = 10.9 Test for overall effect: Z =	100.0% 04, df = 7 (F 6.75 (P < 0 DIOT OF t	3.57 [2.47, 5.18] P = 0.14); I ² = 36% .00001) he outcome: Odds Ratio	0.01 0.1 1 10 10 Favours 1 Day Favours >2 Days deep vein thrombosis Odds Ratio M-H, Fixed, 95% Cl				

Study or Subgroup	Weight	M-H, Fixed, 95% CI	M-H, Fixed, 95% CI
Greenky et al., 2019	44.7%	0.25 [0.04, 1.81]	
Gromov et al., 2019	5.9%	1.62 [0.17, 15.84]	
Nelson 2016., 2016	20.3%	0.56 [0.08, 4.03]	
Richards et al., 2018	20.2%	1.68 [0.49, 5.84]	
Tedder et al., 2018	7.0%	0.73 [0.04, 13.26]	
Weiser et al., 2018	0.6%	24.14 [0.98, 594.97]	
Ziemba-Davis et al., 2019	1.3%	10.80 [0.44, 266.31]	
Total (95% CI)	100.0%	1.00 [0.52, 1.96]	+
Total events			
Heterogeneity: Chi ² = 8.99,	df = 6 (P	$= 0.17$; $I^2 = 33\%$	0.01 0.1 1 10 100
Test for overall effect: $Z = 0$.01 (P = 0	0.99)	Favours 1 Day Favours >2 Days

Forest plot of the outcome: infections

Study or Subgroup	Weight I	Odds Ratio M-H, Fixed, 95% Cl	Odds Ratio M-H, Fixed, 95% Cl	
Nelson 2016., 2016	17.8%	2.25 [0.14, 36.78]		
Otero et al., 2016	82.2%	0.87 [0.11, 6.86]		
Total (95% CI) Total events	100.0%	1.11 [0.21, 5.95]	-	
Heterogeneity: Chi ² = Test for overall effect			0.01 0.1 1 10 10 Favours 1 Day Favours >2 Days	00

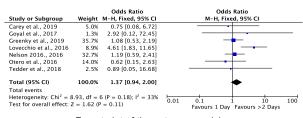
Forest plot of the outcome: mortality

Study or Subgroup	Weight	Odds Ratio M-H. Random. 95% Cl	Odds Ratio M-H. Random, 95% Cl
			M-II, Kalidolli, 55% Ci
Goyal et al., 2017	27.0%	13.24 [0.74, 238.03]	
Gromov et al., 2019	37.2%	4.96 [0.69, 35.79]	
Richards et al., 2018	35.8%	0.36 [0.04, 2.90]	
Total (95% CI)	100.0%	2.52 [0.31, 20.42]	
Total events			
Hotorogonoity: Tau2 -	2.04: Chi	$l^2 = 5.03$, df = 2 (P = 0.08); $l^2 = 6$	
			^{0,6} 0.01 0.1 1 10 100
Test for overall effect:	Z = 0.87 ((P = 0.39)	Favours 1 Day Favours > 2 Days

Forest plot of the outcome: pain

Study or Subgroup	Weight	Odds Ratio M-H, Random, 95% Cl	Odds Ratio M-H, Random, 95% Cl
Aynardi et al., 2014	4.4%	6.12 [0.32, 115.22]	,
Carey et al., 2019	13.1%	0.24 [0.15, 0.40]	
Crampet et al., 2019	8.7%	1.16 [0.25, 5.44]	
Goval et al., 2017	3.9%	2.92 [0.12, 72,45]	
Greenky et al., 2019	12.6%	0.72 [0.38, 1.35]	
Gromov et al., 2019	9.8%	7.85 [2.15, 28.61]	
Lovecchio et al., 2016	12.4%	1.30 [0.65, 2.59]	
Nelson 2016., 2016	12.0%	0.47 [0.21, 1.06]	
Otero et al., 2016	9.1%	0.86 [0.20, 3.75]	
Weiser et al., 2018	9.3%	0.23 [0.05, 0.93]	
Ziemba-Davis et al., 2019	4.6%	79.99 [4.66, 1372.61]	
Total (95% CI)	100.0%	1.11 [0.53, 2.35]	+
Total events			
Heterogeneity: Tau ² = 1.05	$Chi^2 = 50$	$0.97, df = 10 (P < 0.00001); I^2 = 80$	⁶ 0.01 0.1 1 10 100
Test for overall effect: Z = 0	0.28 (P = 0)	.78)	Favours 1 Day Favours >2 Days
			Tavours I Day Tavours >2 Days

Forest plot of the outcome: readmission



Forest plot of the outcome: revision

Figure 3. Forest plots.

4. Discussion

Results from the present meta-analysis encourage outpatient THA. Indeed, according to the main findings of the present study, the outcome of outpatient THA was similar to the procedure with longer hospitalisation. Indeed, no difference was found in pain, infections, mortality, revision, dislocation, and readmission rates between the two groups. A statistically significant greater risk of DVT in patients following outpatient THA was evidenced. This may result from the reduced care by the health care personnel once discharged [34]. According to this evidence, each centre performing outpatient THA should implement continuous homecoming welfare activity, to supervise physiotherapy and the monitoring anticoagulant therapy [14]. Comprehensive multidisciplinary cares include the orthopaedic surgeon, nurses, anaesthesiologists, and physical and occupational therapists. To support homestay recovery, the primary caregiver may help to restore and expedite patient self-care and daily living activities and to reduce mobility [35,36]. A recent systematic review evidenced a higher rate of within 60 days DVT in outpatient THA patients (outpatient 1.28% versus inpatient 1.21%) [37]. This is the only systematic review which reported a higher rate of DVT in outpatient THA patients. Expanding the literature search to articles which evaluated outpatient total joint arthroplasty, DVT-related risk rate did not significantly differ from outpatient and inpatient TJA groups [5,16,38,39]. Results from the present study indicated that outpatient THA, in appropriately selected populations, may faster restore the patient quality of life without affecting the clinical outcome. Moreover, outpatient THA significantly reduces health-care costs [1,11,38,40]. For a successful outpatient THA, proper patient selection is mandatory [38,41–45]. Comorbidities may negatively interfere with the surgical success, being associated with a greater rate of intraoperative and postoperative complications [46]. Suitable patients should be relatively young and healthy, with a solid social support and good performance status [47]. Meneghini et al. [48] developed the Outpatient Arthroplasty Risk Assessment (OARA) to identify which patients may reliably undergo outpatient arthroplasty. The OARA allows to generate risk categories stratifying patients in nine separate comorbidity layers. On the other extreme, Otero et al. [14] stated that patients with comorbidities can still be safely discharged as outpatient THA, given the pivotal importance of the solid social support and networking after discharge at homestay. Optimal pain management protocols should be adopted to be able to undertake outpatients THA in a safe and effective fashion [38]. Postoperative pain management and appropriate patients care are essential factors for outpatient THA pain control at homestay, especially during the first postoperative days [22]. Outpatient THA can be successfully achieved thanks to the advancements in pain management, regional anaesthesia, focused rehabilitation, surgical progresses, and the patient selection process [27]. The management of immediate postoperative complications is critical. Anaesthetic-related side effects such as nausea, hypotension, and urinary retention are common and should be promptly managed [22,26]. The readmission rate is important to evaluate the efficacy of outpatient THA [49,50]. Lovecchio et al. [13], in 1968, found that the most common causes for THA readmission were bleeding and DVT [13]. Springer et al. [51] evidenced increased readmission rates in outpatient THA (11.7% vs. 6.6%), mostly for pain, wound complications, and adverse reaction to pain medication. Greenky et al. [23] found a lower rate of complication and readmission within 30-days postoperatively in favour of outpatient THA. In a recent review, Shapira et al. [52] evidenced a within 3 months outpatient THA readmission rate of 0.34%. In this regard, previously published readmission rates for inpatient THA scored higher [17,53,54]. Those results reinforce the benefits for patients and the health-care system of performing outpatient THA. Regarding the revision rate, we found no difference in both protocols. Evidence with regard to the revision rate was unclear and studies reported controversial results [20]; while some evidence demonstrated a greater revision rate in outpatient THA [22], other authors found a greater rate in the longer hospitalisation cohort [14,23,26], and another still found no difference [13].

This study has limitations. The retrospective design by most of the included studies certainly represented a potential source of selection bias. Moreover, there was an over-

all lack of blinding and the length of the follow-up was not adequate by most studies. Given the lack of quantitative data, subgroup analyses according to the surgical access and exposure, stem length, and implant design were not possible to be analysed separately. However, previous meta-analyses have demonstrated that many of those factors do not impact significantly on the outcome of THA procedures [55–60]. Different surgical approaches, as well as different surgical exposures, do not significantly impact the outcomes of THA [61,62]. Comparable results were reached evaluating implant stem length-related outcomes, and no difference in terms of complications and revision rates was found [60,63]. The heterogeneity between groups with regard to baseline characteristics represents another limitation. Indeed, outpatient THA patients are healthier and younger than their inpatient counterparts. That is not surprising, given the strict inclusion criteria used to enrol outpatient arthroplasty candidates. A patient's age is a known risk factor for perioperative complications in THA [55]. Another study limitation concerns the lack of data regarding information about functional outcomes and rehabilitation programs. Given the heterogeneity of data and protocols, it was not possible to compare outpatient THA versus fast-track or other traditional inpatient lengths of hospital stay. Therefore, in light of these limitations, data from the present study must be considered with caution.

5. Conclusions

In selected patients, outpatient THA can be performed safely with optimal outcomes comparable with the inpatient group. Clear and comprehensive pre-operative planning should involve a multi-disciplinary group composed of orthopaedic surgeons, anaesthesia and rehabilitation specialists, and physiotherapists. Each centre performing outpatient THA should implement continuous homecoming welfare activity, to supervise physiotherapy and monitor anticoagulant therapy.

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