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# Improving function in people with hip-related pain: a systematic review and meta-analysis of physiotherapist-led interventions for hip-related pain

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## ABSTRACT

**Objective** To report the effectiveness of physiotherapist-led interventions in improving pain and function in young and middle-aged adults with hip-related pain.

**Design** Systematic review and meta-analysis.

**Data sources** A comprehensive, reproducible search strategy was performed on five databases in May 2019. Reference lists and grey literature were also searched.

**Eligibility criteria for selecting studies** Population: people aged  $\geq 18$  years with hip-related pain (with or without a diagnosis of femoroacetabular impingement syndrome). Intervention(s): physiotherapist-led interventions for hip pain. Comparators: sham treatment, no treatment or other treatment (eg, hip arthroscopic surgery). Outcomes: primary outcomes included patient-reported hip pain and function. Secondary outcomes included physical function measures.

**Results** 1722 papers were identified. After exclusion criteria were applied, 14 studies were included for analysis. They had varied risk of bias. There were no full-scale placebo-controlled randomised controlled trials (RCTs) of physiotherapist-led treatment. Pooled effects ranged from moderate effects (0.67 (95% CI 0.07 to 1.26)) favouring physiotherapist-led intervention over no treatment post-arthroscopy, to weak effects ( $-0.32$  (95% CI 0.57 to 0.07)) favouring hip arthroscopy over physiotherapist-led treatment.

**Conclusion** Physiotherapist-led interventions might improve pain and function in young and middle-aged adults with hip-related pain, however full-scale high-quality RCT studies are required.

**PROSPERO registration number** CRD42018089088.

## BACKGROUND

Musculoskeletal conditions, such as hip-related pain, are leading causes of pain and disability in the community, and the second largest global contributor to years lived with disability.<sup>1</sup> Hip and groin injuries are common in active individuals, for example, accounting for up to 18% of professional male football injuries.<sup>2-4</sup> The true prevalence of non-arthritic hip pain in the general population is unknown, however the burden of hip pain is high, with younger adults with hip-related pain reporting poor patient-reported outcome scores for pain, physical activity and quality of life<sup>5-8</sup> at a time of life where work and family commitments are large.

Hip-related pain may be classified into three categories, including femoroacetabular impingement (FAI) syndrome, acetabular dysplasia and other

pathology without morphological variants (labral, chondral and ligamentum teres pathology).<sup>9</sup> Of these, FAI syndrome is the most commonly diagnosed clinical condition<sup>10</sup> and is evident in 49% of people with hip pain.<sup>11</sup> Patients with FAI syndrome present with pain, a morphological variant in hip shape on radiographs, with or without intra-articular imaging findings such as labral and/or chondral pathology,<sup>12</sup> and reduced activity and quality of life.<sup>13 14</sup>

Non-surgical treatment approaches should be the first-line options for musculoskeletal pain conditions (evident from clinical guidelines for osteoarthritis (OA)<sup>15 16</sup> and low back pain,<sup>17</sup> due to the far greater costs and risks associated with surgery. Establishing the efficacy of non-surgical treatments for hip pain is critical. Physiotherapist-led interventions have the potential to reduce the burden of hip pain, with current evidence guiding physiotherapist-led treatments to target characteristic modifiable physical impairments<sup>18</sup> (strength, range of motion, functional task performance, neuromuscular/motor/movement control). At present, the level of evidence supporting the efficacy of physiotherapist-led interventions for hip pain and FAI syndrome is unclear.

## Review aim

This systematic review aimed to identify the effectiveness of physiotherapist-led interventions in improving pain and function in young and middle-aged adults who experience hip pain, when compared with sham treatment, no treatment and other treatment. This included non-operative and postoperative patient groups. This review specifically used the participants, interventions, comparators, outcomes (PICO) format.

## METHODS

This systematic review was conducted according to the Preferred Reporting Items for Systematic Reviews and Meta-Analyses guidelines. Literature search criteria and methods were proposed and agreed on by two authors (JK, SC), and were established a priori to minimise selection bias.

## Eligibility criteria for selecting studies

Studies were eligible for inclusion if they were reported in English; reported level IV evidence or above; contained human subjects with hip pain; had at least 10 participants in the overall study sample (5 per group in studies with more than one group)



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and examined the effectiveness of physiotherapist-led interventions. All quantitative study designs were considered, including randomised controlled trials (RCTs), prospective or retrospective approaches.<sup>19</sup> Studies were excluded if: hip pain was due to hip OA, dysplasia or congenital disorder; greater trochanteric pain syndrome and other conditions that are not related to the hip joint; the study evaluated other therapies that were not physiotherapist-led interventions; the study included populations of children or older adults (aged >50 years) or the study was a published abstract, non-peer-reviewed or was written in a language other than English.

#### Participants/Population

People aged 18–50 years with hip pain (based on the mean or median age of the study sample), including studies that included a diagnosis of FAI syndrome.

#### Intervention(s), exposure(s)

Studies reporting physiotherapist-led interventions for hip pain and/or function were included.

#### Comparator(s)/Control

Studies using sham treatment, no treatment or other treatment (eg, hip arthroscopy surgery) as the comparator/control treatment were included.

#### Outcomes

Primary outcomes included patient-reported hip pain and function. Secondary outcomes included: hip joint range of motion, hip muscle strength, functional task performance, electromyography (EMG) and motor control, balance and proprioception, biomechanics and gait analysis and other patient-reported outcome measures.

#### Search strategy

A comprehensive, reproducible search strategy was performed on the following databases from earliest available to 6 November 2017 and was then repeated on 20 May 2019: Medline, CINAHL, Cochrane library, EMBASE and PEDro. Reference lists of included studies were also manually searched for relevant papers. Grey literature, including the Clinical Trials database and the Australia and New Zealand Clinical Trials Registry were searched to identify potential studies that may have been published. Where data were insufficient, authors were contacted and asked to provide missing data. The search terms used PICO format and full search strategy of each database is contained in online supplementary appendix 1. The search strategy was conducted by two reviewers (JK, SC) and used the PICO format, and included:

- ▶ P=human adults (18–50 years) with hip pain.
- ▶ I=physiotherapist-led interventions.
- ▶ C=sham treatment, no treatment, other treatment (eg, surgery).
- ▶ O=pain, function, other patient-reported outcome measures. Function may include hip joint range of motion, hip muscle strength, measures of functional task performance, EMG, gait analysis.

We also used Web of Science to track the forward and backward citations and reference lists of included studies. The strategy was adapted as appropriate for each database. The full search strategy used is contained in online supplementary appendix 1.

Title, abstract and full-text screening was conducted by two independent reviewers (JK, HH) using Covidence (Veritas

Health Innovation, Australia) software. Any disagreements were resolved by a third independent reviewer (KC).

#### Risk of bias assessment

The Cochrane Collaboration Risk of Bias tool for Clinical Trials was used to appraise risk of bias. Included studies were rated by two independent reviewers (MB, MJS). Any disagreements between reviewers were discussed in a consensus meeting and an independent arbitrator (JK) was employed when consensus could not be met. Agreement between raters was determined using Cohen's Kappa ( $\kappa$ ). If risk of bias was high in >three out of five categories, overall study risk was rated as high, if risk was high in three out of five categories, study risk was moderate and if risk was high in <three out of five categories, overall study risk was rated as low.<sup>20</sup> All studies were included in subsequent analyses, and sensitivity analyses were performed as appropriate.

#### Data extraction, synthesis and analyses

All potential references were imported into Endnote X7 (Thomson Reuters, Carlsbad, California, USA) and duplicates removed. Data were extracted by two independent reviewers (JK, ABM). Any discrepancies in data extraction were resolved by an independent arbitrator (KC).

Findings were summarised and population characteristics (age, gender, type and description of hip OA, duration of symptoms), and details of outcome measures, length of follow-up and type intervention undertaken were collated. We have reported main findings only for studies where the physiotherapist-led intervention was compared with a comparator/control intervention (RCT design) in order to ensure only higher quality evidence was included.

For studies of RCT design, follow-up scores were compared with the published Patient Acceptable Symptom State (PASS) scores for that outcome (if known) and change scores were compared with the published minimal important change (MIC) score for that outcome (if known). The proportion of people who achieve a PASS from follow-up scores was estimated using previously published methods, incorporating means, SD, sample size and z-scores.<sup>21</sup> Previously published relevant PASS scores include 88 points (Hip Osteoarthritis and disability Outcome Score (HOOS)-pain)<sup>22</sup> and 83 points (HOOS-quality of life (QOL))<sup>22</sup> 1-year posthip arthroplasty; 58 points (International Hip Outcome Tool (IHOT)-33)<sup>23</sup> 1–5 years posthip arthroscopy and 98 points (Hip Outcome Score (HOS)-activity of daily living (ADL))<sup>24</sup> and 94 points (HOS-Sport)<sup>24</sup> 1-year posthip arthroscopy. Previously published MIC scores include 9 points (HOOS-pain),<sup>25</sup> 11 points (HOOS-QOL),<sup>25</sup> 10 points (IHOT-33)<sup>25</sup> 1–2 years posthip arthroscopy; 15 points (HOS-ADL)<sup>24</sup> 1-year posthip arthroscopy and 28 points (HOS-Sport) 6 months posthip arthroscopy.<sup>24</sup>

Data analyses were conducted by two investigators (AIS and JK). The 'meta' package (V.4.9–5), from the R statistical software package (V.3.5.1) was used to calculate relevant effect sizes, produce forest plots and pool data in a meta-analysis where relevant (<https://www.r-project.org/>). Standardised mean differences (SMD) were calculated for the studies of RCT design, to determine the magnitude of the effect of any interventions within groups or between groups. Where data were deemed statistically and clinically homogenous, meta-analyses were undertaken using a random effects model. In order to undertake SMD calculations in studies where non-normally distributed data were presented, the IQR was calculated.<sup>26</sup> For analysis of outcomes that reported within group (pre-intervention to post-intervention), the

standardised paired difference (SPD) was calculated from the sample size, mean and SD of the difference from pre-intervention to post-intervention time points. An additional requirement for SPD calculation was to account for the within-person correlation ( $r$ ) between the two time points. If between-limb correlation was not reported, a conservative estimate of  $r=0.5$  was used.<sup>27</sup> Standardised mean/paired difference magnitude was interpreted as:  $\geq 0.8$  large effect;  $0.5-0.79$  moderate effect and  $0.2-0.49$  weak effect.<sup>28</sup> Where SMDs could not be calculated, study conclusions were presented in tables, and best evidence synthesis was performed. For the best evidence synthesis, evidence was categorised as 'strong' if there were multiple high-quality clinical trials or cohort studies; 'moderate' if there was either one high-quality clinical trial or cohort study and more than two high-quality case-control studies or pilot clinical trials, or more than three high-quality case-control studies; 'limited' if there were either one or two case-control studies or pilot clinical trials, or multiple cross-sectional studies and 'insufficient' if there was not more than one cross-sectional study.<sup>20</sup> All data used in calculations of

SMDs and SPDs have been shared publicly at Figshare (<https://figshare.com/s/d18bcb066f1de48861cf>).

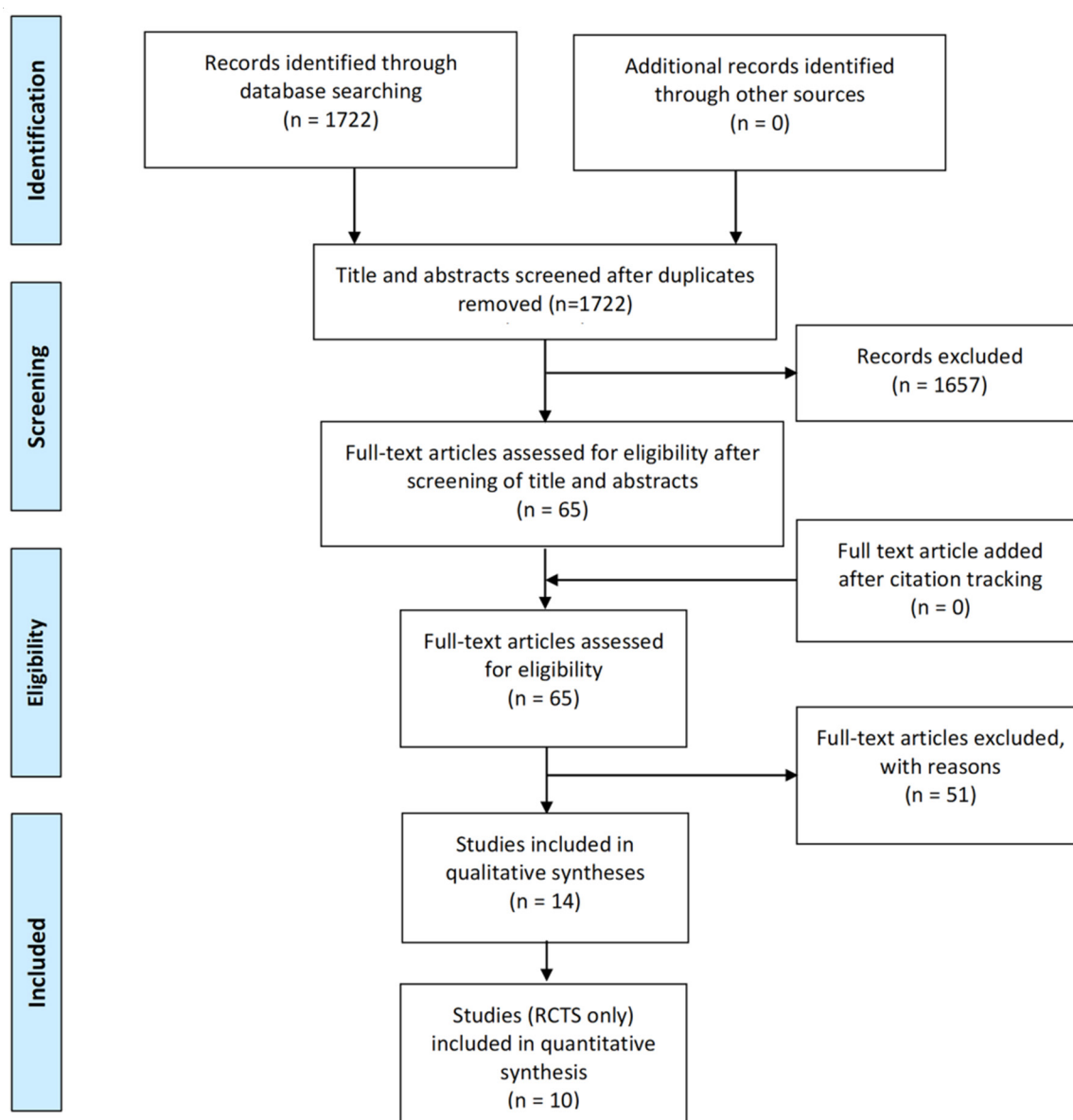
## RESULTS

### Search strategy

The search yielded 1722 titles and abstracts for screening. Sixty-five full texts were screened and 51 were excluded. There were 14 papers in the final analyses. An overview of the study identification process is provided in figure 1. Characteristics of the included studies are presented in table 1. The number of studies excluded along with reasons is provided in online supplementary appendix 2.

### Risk of bias

Online supplementary appendix 3 contains the results of risk of bias assessment using the Cochrane Risk of Bias Tool. Agreement between raters occurred on 54 out of 70 items, where  $\kappa=0.65$ , which represents moderate agreement.<sup>29</sup> Following discussion, consensus was obtained for all items. Overall results for the



**Figure 1** Preferred Reporting Items for Systematic Reviews and Meta-Analyses flow chart of study inclusion. RCT, randomised controlled trial.

**Table 1** Summary of included studies

Study	Title	Study type	Overall risk of bias	Inclusion criteria	Exclusion criteria	Number (PT/ control) at baseline	Number (PT/ control) at follow-up	Age mean (SD) (PT/ control)	Sex (M/F) (PT/ control)	BMI kg/m <sup>2</sup> mean (SD) (PT/ control)	Intervention type	Control type	Primary end point	Other follow-up	Primary outcomes	Secondary outcomes	Included in SMD calculation (Y/N)
<b>Bennell, et al 2017<sup>33</sup></b>	Efficacy of adding a physiotherapy rehabilitation programme to arthroscopic management of femoroacetabular impingement syndrome: a randomised controlled trial	Pilot RCT	Moderate	Aged >16 years; FAI; scheduled for arthroscopy	Hip OA (Tönnis >grade 1); professional athlete; concurrent injury; unable to attend study; physiotherapist unwilling to refrain from rehab	14/16	11/13	31 (7) / 29 (8)	12/2 / 12/4	24.6 (2.2) / 25.2 (3.2)	7 PT sessions (1 pre-op; 6 postoperative) education manual therapy deep hip rotator retraining, stretching, bike, pool jog	No therapy allowed	14 weeks	24 weeks	IHOT-33; HOS-Sport subscale	HAGOS subscales; Legier; HSAS; HOS-ADL	Y
<b>Coppack, et al 2016<sup>34</sup></b>	Physical and functional outcomes following multidisciplinary residential rehabilitation for prearthritic hip pain among young active UK military personnel	Case series	High	Non-surgical and postsurgical military personnel attending for physical therapy for of hip pain	Not reported	40	40	33 (7)	27/13	25.8 (4.5)	5 hours/day 3 weeks. Hip ROM and strength, core and trunk muscle function, deep hip stabiliser exercise, correct gait balance retraining, weight management, patient education	NA	3 weeks	NA	HAGOS subscales Pain VAS	Y-balance flexion ROM IR ROM; Shuttle test	N
<b>Enara, et al 2011<sup>21</sup></b>	Conservative treatment for mild femoroacetabular impingement	Case series	High	Men and women with unilateral hip pain; alpha angle <60	Aged >55 years; skeletally immature; hip disease or hip surgery; alpha angle >60; hip OA	37	37	NR	27/10	NR	Stage 1=rest, NSAIDs Stage 2=stretching exercise Stage 3 and 4=ADL modification	NA	6 months	24 months	HHS MAHS VAS	Flexion ROM Extension ROM Abduction ROM ER in flexion ROM ER in extension ROM IR in flexion ROM IR in extension ROM	N
<b>Grant, et al 2017<sup>34</sup></b>	The HAPI Hip Arthroscopy Prehabilitation Intervention study: does prehabilitation affect the outcomes in patients undergoing hip arthroscopy for femoroacetabular impingement?	Pilot RCT	Moderate	Men and women aged >18 years with FAI, awaiting hip arthroscopy	Radiographic evidence of hip dysplasia; grades 3-4 OA; AVN; rheumatology disorders; <18 years; previous hip surgery	9/9	8/8	38 (6)	4/4 / 1/7	NR	Five visits, points of interest were two pre-operative exercise for hip strength	Five visits, two pre-operative, same as intervention group except no pre-operative exercise advice	2 weeks pre-operative	NA	MAHS	Abduction strength Adduction strength Flexion strength ER strength Knee extension strength	Y
<b>Griffin, et al 2018<sup>7</sup></b>	Hip arthroscopy vs best conservative care for the treatment of femoroacetabular impingement syndrome (UK FASHION): a multicentre randomised controlled trial	RCT	Moderate	Men and women aged 16+ years with hip pain, suitable for hip arthroscopy	Hip OA (Tönnis >1) significant hip pathology past hip surgery	177/171	162/157	35 (9)/35 (10)	113/64 / 100/71	NR	Personalised hip therapy between 6 and 10 treatments over 12-24 weeks of assessment, education, exercises (core, neuromotor control, progressive strength, stretching) pain relief	Hip arthroscopy surgery as pragmatically determined by surgeon, including acetabular rim trimming, labral debridement or repair, femoral osteoplasty	12 months	6 months	IHOT-33	EQ-5D-5L EQ-5D VAS SF-12 PCS SF-12 MCS	Y
<b>Guenther, et al 2017<sup>25</sup></b>	A pre-operative exercise intervention can be safely delivered to people with femoroacetabular impingement and improve clinical and biomechanical outcomes	Case series	Moderate	FAI on MRI; positive impingement test; anterior groin pain	Hip OA; CSI within last month; hip surgery; significant lower body injuries or conditions; osteonecrosis of the hip; planned commencement or current enrollment in a hip strengthening exercise programme	20	20	30 (7)	18/2	24.1 (2.9)	Progressive phased strength programme, bilateral intervention, Exercises at home 4xweek Phase I=activation, neuromuscular control, endurance Phase II=strength and intensity Phase III=strength and function	NA	10 weeks	NA	HOS Symptoms HOS Pain HOS-ADL HOS-Sport HOS-QOL	Abduction strength Adduction strength Extension strength ER strength IR strength lined stair climb test	N
<b>Harris-Hayes, et al 2016<sup>29</sup></b>	Movement pattern training to improve function in people with chronic hip joint pain: a feasibility randomised clinic trial	Pilot RCT	Low	Aged 18-40 years; anterior hip or groin pain >3 months; positive FADIR test	Previous hip surgery or fracture; contraindication to MRI; pregnancy; neurological involvement; knee or low back pain; BMI >30	18/17	16/16	27 (5)/29 (5)	1/15 / 4/12	24.2 (3.0) / 24.4 (2.6)	Movement pattern retraining; task-specific training for basic functional tasks; progressive strengthening of hip musculature, 6x1 hour visits over 6 weeks individualised and progressed	Wait list control	6 weeks	NA	HOS Symptoms HOS-Pain HOS-ADL HOS-Sport HOS-QOL	ABD strength ER 0 strength ER 90 strength Adduction single leg squat	Y

Continued

**Table 1** Continued

Study	Title	Study type	Overall risk of bias	Inclusion criteria	Exclusion criteria	Number (PT/ control) at baseline	Number (PT/ control) at follow-up	Age mean (SD) (PT/ control)	Sex (M/F) (PT/ control)	BMI kg/m <sup>2</sup> mean (SD) (PT/ control)	Intervention type	Control type	Primary end point	Other follow-up	Primary outcomes	Secondary outcomes	Included in SMD calculation (Y/N)
Hunt <i>et al</i> 2012 <sup>22</sup>	Clinical outcomes analysis of conservative and surgical treatment of athletes with clinical indicators of prearthritic, intra-articular hip disorders	Case-control	High	Anterior or lateral hip pain; pain on activity; physical associated medial symptoms; pain at rest; positive anterior hip impingement test; FABER test; leg pain or resisted straight leg raise test	Aged >50 years; history hip surgery; inflammatory arthropathy; hip infection or tuberculosis; hip OA (Tomms 2-3)	2729	2729	33 (11)/36 (11)	2/23 27/29	26.3 (7) 23.9 (5)	Conservative interventions, patient education, activity modification, directed physical therapy protocol	Same as PT plus injection and surgery (when patient not satisfied with PT)	12 months	NA	Numeric pain scale HHS WOMAC Backde Questionnaire SF-12 PCS SF-12 MCS NAHS	NA	N
Kemp <i>et al</i> 2018 <sup>36</sup>	A pilot randomised clinical trial of physiotherapy (manual therapy, exercise and education) for early onset hip osteoarthritis posthip arthroscopy	Pilot RCT	Moderate	Aged 18-50 years; 4-14 months posthip arthroscopy for early hip OA (Chondropathy outbridge grade ≥ 1); pain ≥ 3/10	Hip buritis or tendinopathy; surgical complications; planned further surgery in the following 12 months	107	96	32 (10)/31 (6)	6/4 2/5	26.3 (5.3) 24.4 (4.7)	8x30 min sessions over 12 weeks, and home exercise programme 4xweek (i) Manual hip joint and soft-tissue mobilisation and stretching; (ii) hip muscle retraining; (iii) trunk muscle retraining; (iv) functional, proprioceptive and sports-specific or activity-specific retraining; (v) enhancing physical activity and (vi) education	8x30 min sessions over 12 weeks, education only	12 weeks	NA	HOOS-Symptoms HOOS-Pain HOOS-ADL HOOS-Sport HOOS-QOL IHOT	Abduction strength Adduction strength Extension strength Flexion strength ER Strength IR Strength Flexion ROM	Y
Kemp <i>et al</i> 2018 <sup>40</sup>	The Physiotherapy for Femoroacetabular Impingement Rehabilitation Study: a pilot randomised controlled trial	Pilot RCT	Low	Aged 18-50 years; no surgery; FAI (pain >3/10; >6 weeks, in hip or groin; alpha angle >60)	Past surgery; significant other hip disease; pregnancy; physiotherapy within past 3 months	177	146	37 (8)/28 (10)	5/12 2/5	25.1 (3.7) 26.1 (2.4)	Individualised, progressive programme 8x30 min 1:1 and 12x30min supervised gym, 2xHEP (12-week intervention), Hip strength, trunk strength, functional retraining, manual therapy, education, fitness programme	Standardised programme 8x30 min 1:1 and 12x30min supervised gym, 2xHEP (12-week intervention), Hip strength, trunk strength, functional retraining, manual therapy, education, fitness programme	12 weeks	Nil	IHOT	HOOS-QOL HOOS-Pain Adduction strength Abduction strength Extension strength ER strength Flexion ROM Single leg hop Stair bridge	Y
Mansell <i>et al</i> 2018 <sup>41</sup>	Arthroscopic surgery or physical therapy for patients with femoroacetabular impingement syndrome	RCT	Low	Military personnel and families; FAI and/or labral tear (pain in hip or groin); pain on flexion; pain on FADIR; pain relief after injection; candidate for surgery (failed 6/52 conservative; positive cross-over or alpha angle >55)	Hip OA; other concurrent hip disease; workers compensation; positive LBP findings; pregnancy; past surgery; physiotherapy within previous 6 months	40/40	33/29	31 (7)/30 (7)	21/19 26/14	27.5 (4.3) 28.2 (4.4)	Standardised, supervised PT, 2x/week for 6 weeks. Included elements—manual therapy (joint mob, MMW, soft tissue therapy), stretching, repair, femoral therapeutic and motor control exercises	Pragmatic hip arthroscopy surgery (acetabular rim trimming, labral debridement or repair, femoral osteoplasty)	2 years	6, 12 months	HOS-ADL HOS-Sport	IHOT Global rating of change	Y
Palmer <i>et al</i> 2019 <sup>41</sup>	Arthroscopic hip surgery compared with physiotherapy and activity modification for the treatment of symptomatic femoroacetabular impingement: multicentre randomised controlled trial	RCT	Low	Aged 18-60 years; referred to secondary or tertiary care. Clinically and radiologically diagnosed FAS (no imaging thresholds used. Instead surgeons qualitatively assessed morphology).	Physiotherapy in past 12 months; past hip surgery; hip OA (KLz2) hip dysplasia (LCEA <20).	110/112	88/100	36 (10)/36 (10)	37/73 38/74	26.6 (4.8)/25.9 (4.8)	Goal-based supervised PT, up to 8 sessions over 5 months. Patient goals supplemented with programme focusing on muscle strength, core stability and movement control. Avoidance of extreme ROM encouraged. No specific programme details available	Pragmatic hip arthroscopy surgery (osteochondroplasty, labral repair or debridement, chondral debridement and microfracture as deemed appropriate by surgeon)	8 months		HOS-ADL HOS-Sport NAHS OHS HAGOS IHOT-33 EQ-5D-3L PainDETECT HADS Hip flexion ROM Hip extension ROM Hip abduction ROM Hip adduction ROM Hip ER ROM Hip IR ROM		Y
Sneatham <i>et al</i> 2017 <sup>27</sup>	Does treatment by a specialist physiotherapist change pain and function in young adults with symptoms from femoroacetabular impingement? A pilot project for a randomised controlled trial	Pilot RCT	Moderate	FAI diagnosed Surgeon (FAI on X-ray; Aged 18 and 50 years; groin/anterolateral hip pain; mechanical symptoms; pain FADIR; willing for treatment by a specialist physiotherapist, and abstain from treatment outside study protocol)	Previous hip surgery to the hip or pelvis; other significant hip pathology; back symptom; chronic pain; inability to comply with the physiotherapy protocol	15/15	11/12	36/33	7/8 5/10	MR	Pragmatic care from specialist physio Any number of sessions allowed. Manual therapy and exercise to address movement deficits	Routine care (analgesia, continuation self-management or exercise previously given)	3 months	NA	NAHS HOS LEFS VAS	NA	Y

Continued

Table 1 Continued

Study	Title	Study type	Overall risk of bias	Inclusion criteria	Exclusion criteria	Number (PT/ control) at baseline	Number (PT/ control) at follow-up	Age mean (SD) (PT/ control)	Sex (M/F) (PT/ control)	BMI kg/m <sup>2</sup> mean (SD) (PT/ control)	Intervention type	Control type	Primary end point	Other follow-up	Primary outcomes	Secondary outcomes	Included in SMD calculation (Y/N)
Wright, et al 2016 <sup>38</sup>	Non-operative management of femoroacetabular impingement: a prospective, randomised controlled clinical trial pilot study	Pilot RCT	Moderate	Aged 18–50 years clinical diagnosis FAI (hip flexion <95°, internal rotation <10°, positive FADIR or FABER) radiological diagnosis FAI (alpha angle >55°, lateral centre edge angle >35°, crossover sign)	Previous hip surgery; other surgical procedure of lower limb in the prior 6 months; pre-existing hip disease pregnancy; opioid analgesia or CSI past 30 days; advanced osteoporosis; BMI >38; cardiopulmonary disease	87	8/7	31 (5)/36 (12)	3/4 1/7	25.6 (3.7) 24.1 (7.4)	Manual therapy and supervised exercise	Advice and home exercise	7 weeks	NA	HOS-ADL HOS-Sport NPRS LEFS SANE-ADL SANE-Sport	Depth of squat Triple hip Hip flexion ROM Hip flexion Strength FABER	Y

ADL, activity of daily living; AVM, avascular necrosis; BMI, body mass index; CSI, corticosteroid injection; EQ-5D-5L, EuroQol Questionnaire; ER, external rotation; F, female; FABER, flexion-abduction-external rotation test; FADIR, flexion-adduction-internal rotation test; FAI, femoroacetabular impingement; HAGOS, Copenhagen Hip and Groin Outcome Score; HHS, Harris-Hip Score; HOOS, Hip Osteoarthritis and Disability Outcome Score; HOS, Hip Outcome Score; HSAS, Hip Sports Activity Scale; IHOT, International Hip Outcome Tool; IR, internal rotation; KL, Kellgren Lawrence; LCEA, lateral centre edge angle; LEFS, lower extremity functional scale; m, metre; M, male; MCS, emotional function subscale; MBA, magnetic resonance arthrogram; n, no; NA, not applicable; NAHS, Non-Arthritic Hip Score; NPRS, Numeric Pain Rating Scale; NR, not reported; NSAIDs, non-steroidal anti-inflammatory drugs; OA, osteoarthritis; PA, physical activity; PCS, physical function subscale; PT, physiotherapy/physical therapy; QOL, quality of life; RCT, randomised controlled trial; ROM, range of motion; SANE, single assessment numeric evaluation; SF-12, Short Form-12 Questionnaire; SMD, standardised mean difference; VAS, Visual Analogue Scale; WOMAC, Western Ontario and McMaster Universities Arthritis Index; Y, yes.

risk of bias assessment are mentioned in table 1. Three studies had a high risk of bias,<sup>30–32</sup> seven studies had a moderate risk of bias<sup>7 33–38</sup> and four studies had a low risk of bias.<sup>8 39–41</sup> In the included studies, the overall risk of performance bias (blinding of participants and personnel) and detection bias (blinding of outcome assessors) was high (high in ≥ nine studies); the risk of attrition bias (incomplete outcome data) and the risk of selection bias (random sequence generation and allocation concealment) was moderate (high in six to eight studies) and the risk of reporting bias (selective reporting of outcomes) was low (high in < six studies).

**Participants**

The 14 included studies contained 542 patients (283 men, 259 women) with sample sizes of the physiotherapist-led intervention groups ranging from 8 patients<sup>38</sup> to 177 patients.<sup>7</sup> Ten studies were of RCT design. Mean participant age ranged from 27 to 38 years, while the mean body mass index (BMI) ranged from 24.1 to 27.5 kg/m<sup>2</sup>. Ten studies included participants based on a diagnosis of FAI syndrome,<sup>8 10 30 31 33–35 37 38 40 41</sup> with the remaining four studies including subjects based on a diagnosis of hip pain.<sup>30 32 36 39</sup> Methods used for diagnostic inclusion criteria comprised surgical findings,<sup>28</sup> clinical examination results<sup>8 10 14 30–32 35–41</sup> and radiological findings.<sup>31 35 37 38 40 41</sup> Two studies did not specify how FAI syndrome was diagnosed for inclusion<sup>33 34</sup> (table 1). One study provided information about level of sports/physical activity,<sup>41</sup> and no study provided detail about the duration of symptoms.

**Outcomes measured**

All included studies used a patient-reported outcome measure (PROM) as the primary outcome measure, but there was large heterogeneity in the PROMs used. The PROMs used included: the IHOT-33, the Copenhagen Hip and Groin Outcome Score (HAGOS), HOOS, the Oxford Hip Score (OHS), the Non-Arthritic Hip Score (NAHS), HOS, the Harris Hip Score (HHS), a pain Visual Analogue Scale (VAS), a Numeric Pain Rating Scale (NPRS), the Hip Sports Activity Scale (HSAS), a Global Rating of Change (GROC) score, the Hospital Anxiety and Depression Scale (HADS), the University of California, Los Angeles (UCLA) activity score, the European Quality of Life-5 Dimensions (EQ-5D) and the 36-item Short Form survey (SF-36) score.

Secondary outcomes measured were mostly measures of physical function, and included: hip muscle strength, trunk muscle strength, standardised hopping tests, measures of performance on a double-leg and single-leg squat, hip range of motion tests, the timed stair climb test and the Y-balance test. The methods used to measure these impairment-based outcomes varied widely between studies. Primary follow-up time points also varied and ranged from 3 weeks<sup>30</sup> to 2 years.<sup>8</sup> Most studies undertook a 3-month primary follow-up period (table 1).

**Physiotherapist-led interventions performed**

Seven studies included participants who had not undergone hip surgery,<sup>31 32 35 37–40</sup> while three studies examined physiotherapist-led interventions posthip arthroscopy surgery,<sup>33 34 36</sup> one study included both postsurgical and non-surgical participants<sup>30</sup> and three studies compared physiotherapist-led interventions to hip arthroscopic surgery.<sup>7 8 41</sup> The duration of physiotherapist-led interventions ranged from 3 weeks<sup>30</sup> to 5 months.<sup>34 41</sup>

There was a large variety in the types of physiotherapist-led interventions performed. Nine studies included a strengthening programme,<sup>7 10 30 34–37 39–41</sup> four studies included stretching/

ROM exercises,<sup>7 8 10 31</sup> five included 'core stability',<sup>7 30 38 39 41</sup> eight studies included manual therapy,<sup>8 30 33 34 36–38 40</sup> two studies included cardiovascular and return to sport retraining,<sup>33 40</sup> two studies included functional retraining<sup>39 40</sup> and six studies included neuromotor control exercises<sup>7 8 30 36 38 39</sup> (table 1). Five studies did not report the interventions in sufficient detail to allow replication of the interventions.<sup>30–32 37 41</sup>

There was large heterogeneity in the control/comparator interventions used in the 10 RCTs. Control/comparator interventions used included (i) sham treatments, such as exclusion of pre-operative exercise advice,<sup>34</sup> a standardised minimal intervention,<sup>40</sup> education only,<sup>36</sup> usual care (including medication and continuation of previous exercises)<sup>37</sup> and a home-based unsupervised exercise programme<sup>38</sup>; (ii) no-treatment group<sup>33</sup> or waiting-list control group<sup>39</sup> and (iii) hip arthroscopy surgery<sup>7 8 41</sup> (table 1).

### Main findings

Between-group differences were generated for physiotherapist-led interventions compared with the comparator intervention in four studies in patients who had not undergone surgery<sup>37–40</sup>; in three studies posthip arthroscopy<sup>33 34 36</sup> and compared with hip arthroscopic surgery<sup>7 8 41</sup> in three studies (table 2).

#### Between-group comparisons of physiotherapist-led interventions compared with sham/no treatment in non-surgical patients

The level of evidence was limited, with two high-quality<sup>39 40</sup> and two moderate-quality<sup>37 38</sup> pilot RCTs included. In patients with hip pain (non-surgical), physiotherapist-led interventions of 3 months duration that included targeted strengthening programmes showed moderate pooled effects for function (SMD (95% CI): 0.66 (0.09 to 1.23)) favouring the physiotherapist-led intervention group<sup>37 40</sup> (figure 2). For physiotherapist-led interventions of shorter duration (6–8 weeks), effects showed no significant differences between groups (figure 3). One study achieved a follow-up score greater than the PASS score, and change score greater than the MIC for the primary outcome.<sup>40</sup>

#### Between-group comparisons of physiotherapist-led interventions compared with sham/no treatment in post-hip arthroscopy patients

The level of evidence was limited, with two moderate-quality pilot RCTs included.<sup>33 36</sup> Data could not be pooled due to heterogeneity in outcomes assessed. Moderate positive effects for patient-reported function (0.67; 95% CI 0.07 to 1.26) were reported in the two RCTs, favouring the physiotherapist-led interventions (figure 4). Both studies achieved a follow-up score greater than the PASS score and change score greater than the MIC for the primary outcome. The proportion of participants undertaking physiotherapist-led interventions achieving a score greater than the PASS score ranged from 11%<sup>36</sup> to 90%.<sup>33</sup>

#### Between-group comparisons of physiotherapist-led interventions compared with hip arthroscopic treatment

The level of evidence was strong, with two high-quality RCTs<sup>8 41</sup> and one moderate-quality RCT<sup>7</sup> included (figure 5). In studies comparing physiotherapist-led interventions with hip arthroscopic surgery, at 8–12 months, weak positive pooled effects (−0.32; 95% CI −0.57 to −0.07) favoured hip arthroscopy surgery. At 24 months, there was only one moderate-quality RCT and thus the level of evidence is limited. There was no significant difference between groups (−0.18; 95% CI −0.64 to 0.28). For the physiotherapist-led intervention groups, no

studies achieved a follow-up score greater than the PASS score, and only one study had a change score greater than the MIC for the primary outcome.<sup>7</sup> The proportion of participants undertaking physiotherapist-led interventions achieving a score greater than the PASS score ranged from 7%<sup>41</sup> to 37%.<sup>7</sup>

#### Within-group change for patient-reported outcome measures for physiotherapist-led interventions in non-operative patient groups

Within-group effects for physiotherapist-led interventions on PROMS in patients with hip pain were able to be calculated for nine of the included studies<sup>7 8 30 32 37–40</sup> (table 3). The level of evidence overall was moderate, with one high-quality RCT and several high-quality pilot RCTs included in the analyses. Positive SPDs ranged from moderate effects for patient-reported function (0.57; 95% CI 0.03 to 1.12)<sup>37</sup> following a 3-month intervention, to large positive effects for function (3.85; 95% CI 2.91 to 4.78)<sup>31</sup> following a 6-month intervention. Data were not able to be pooled due to heterogeneity between time points and the outcomes measured. The proportion of participants undertaking physiotherapist-led interventions achieving a score greater than the PASS score ranged from 25%<sup>38 39</sup> to 86%.<sup>40</sup>

#### Within-group change for physical impairments for physiotherapist-led interventions in non-operative patient groups

Nine studies reported the effects of physiotherapist-led interventions on physical impairments on people with hip pain<sup>30 31 34 35 37–41</sup> (table 3), with SPDs able to be calculated for seven of the nine studies. The level of evidence was limited, with no high-quality, full-scale RCTs included in any of the analyses relating to physical impairments. The impairment measures included hip range of motion,<sup>30 31 38 40</sup> hip muscle strength,<sup>35 38–40</sup> depth of squat,<sup>38</sup> balance,<sup>30</sup> trunk endurance,<sup>40</sup> control during single leg squat<sup>39</sup> and hopping performance.<sup>38 40</sup> Data were not able to be pooled for within-group change in physical impairment measures, due to heterogeneity between time points and the methods by which outcomes were measured.

For hip flexion range of motion, SPDs varied, and ranged from large negative changes (2.07, 95% CI −2.64 to −1.50) following a 6-month intervention consisting of rest, stretching and activity modification<sup>31</sup> to large positive change (1.08, 95% CI 0.49 to 1.68) following a 3-month intervention comprising strengthening exercise, manual therapy and education.<sup>40</sup>

Hip muscle strength was recorded in four studies,<sup>35 38–40</sup> and SPDs ranged from weak, non-significant effects (0.09, −0.35 to 0.53) for an 10 week intervention comprising progressive strengthening exercises,<sup>35</sup> to large positive SMDs (1.19, 0.57 to 1.81) for a 12 week intervention comprising targeted strengthening and functional retraining exercises.<sup>40</sup>

There were varied within-group changes in functional task performance. Positive SPDs ranged from moderate improvements in the timed stair climb test (0.57, 95% CI 0.10 to 1.05),<sup>35</sup> single-leg hop test (0.65, 95% CI 0.12 to 1.17)<sup>40</sup> and Y-balance test (0.63, 95% CI 0.29 to 0.97)<sup>30</sup> with movement retraining and functional exercise programmes, to large improvements for trunk endurance (0.95, 95% CI 0.38 to 1.53) following a 3-month targeted trunk-strengthening programme.<sup>40</sup>

## DISCUSSION

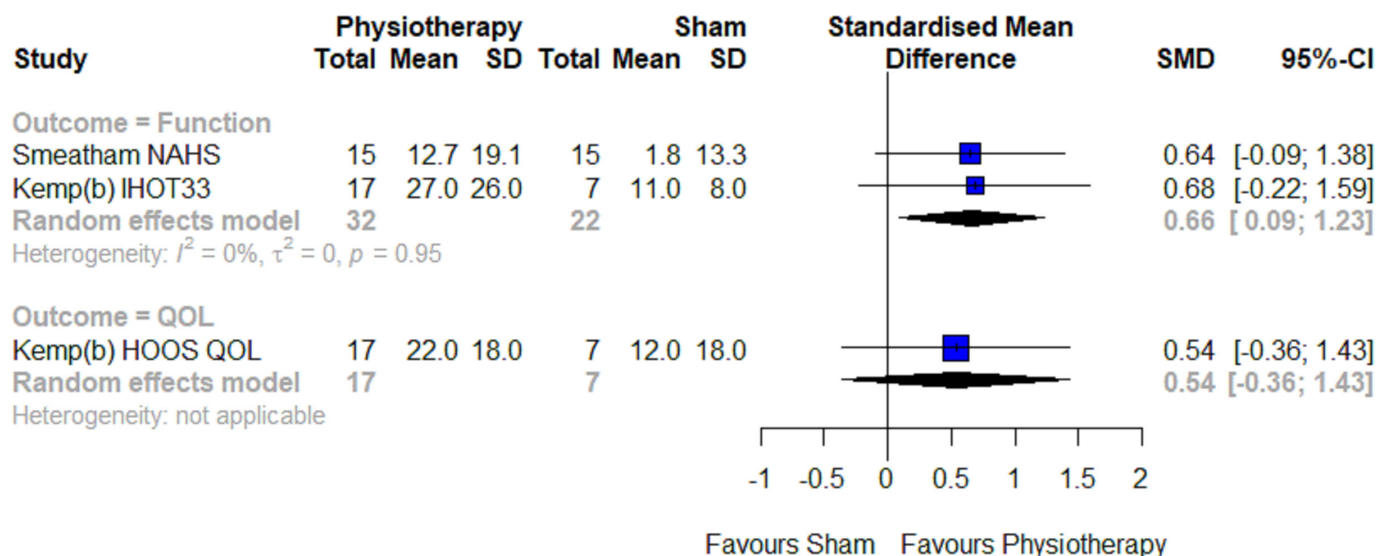
Our systematic review evaluated the effectiveness of physiotherapist-led interventions to improve pain and function in young and middle-aged adults experiencing hip-related pain, including those with FAI syndrome. The 14 studies included 7 pilot and 3 full-scale RCTs, and demonstrated considerable variability in

**Table 2** Summary of results of between-group SMD for primary outcomes of included randomised controlled trials

Study	Title, journal	Baseline M (SD) PT control	Baseline M (SD) control	Primary FU M (SD) PT	Primary FU M (SD) control	Other FU M (SD) PT	Other FU M (SD) control	Did primary FU score reach?	Proportion of participants with FU score greater than PASS score	PT group change score (primary outcome primary end point)	Control group change score primary outcome primary end point	Did primary change score reach MIC?	Between-group SMD (95% CI) for primary outcomes (positive SMD favours PT intervention)
<b>Studies comparing physiotherapist-led treatment with sham or control treatment in non-surgical cohorts</b>													
<b>Grant, et al 2017<sup>24</sup></b>	The IAPT 'Hip Arthroscopy Pre-Intervention' study: does prehabilitation affect the outcomes in patients undergoing hip arthroscopy for femoroacetabular impingement?	NAHS 60 (23.6)	NAHS 54.4 (24.5)	NAHS 56.3 (18.5)	NAHS 48.8 (20.6)	NA	NA	Unknown	NA	Author did not provide data	Author did not provide data	Unknown	NAHS NC
<b>Harris-Hayes, et al<sup>25</sup></b>	Movement pattern training to improve function in people with chronic hip joint pain: a feasibility randomised clinical trial	HOS-Pain 78.2 (13.8) HOS-QOL 65.1 (13.0)	HOS-Pain 73.9 (13.2) HOS-QOL 55.3 (18.3)	HOS-Pain 81.5 (13.2) HOS-QOL 71.3 (19.7)	HOS-Pain 75.8 (13.4) HOS-QOL 65.1 (19.7)	NA	NA	No No	5/16 (31%) 4/16 (25%)	HOS-Pain 1.9 (10.4) HOS-QOL 7.8 (11.8)	HOS-Pain 1.9 (10.4) HOS-QOL 7.8 (11.8)	No No	HOS-Pain 0.13 (-0.53 to 0.80) HOS-QOL -0.14 (-0.80 to 0.52)
<b>Kemp, et al<sup>26</sup></b>	The Physiotherapy for Femoroacetabular Impingement Rehabilitation Study: a pilot randomised controlled trial	IHOT 60 (26) HOS-QOL 54 (21)	IHOT 56 (25) HOS-QOL 50 (17)	IHOT 87 (12) HOS-QOL 76 (13)	IHOT 67 (24) HOS-QOL 62 (24)	NA	NA	Yes No	6/7 (86%) 2/7 (29%)	IHOT 11 (8) HOS-QOL 12 (8)	IHOT 11 (8) HOS-QOL 12 (8)	Yes Yes	IHOT 0.68 (-0.22 to 1.58) HOS-QOL 0.54 (-0.36 to 1.43)
<b>Sneatham, et al 2017<sup>27</sup></b>	Does treatment by a specialist physiotherapist change pain and function in young adults with symptoms from femoroacetabular impingement? A pilot project for a randomised controlled trial	NAHS Total 50.1 (19.1)	NAHS Total 54.4 (13.3)	NAHS Total 62.2 (16.1)	NAHS Total 55.2 (15.0)	NA	NA	Unknown	NA	NAHS Total 12.7 (19.1)	NAHS Total 1.8 (13.3)	Unknown	NAHS Total 0.64 (-0.09 to 1.38)
<b>Wright, et al 2016<sup>28</sup></b>	Non-operative management of femoroacetabular impingement: a prospective, randomised controlled clinical trial pilot study	HOS-ADL 74.3 (13.1) HOS-Sport 59.4 (18.0)	HOS-ADL 73.8 (11.3) HOS-Sport 51.4 (16.2)	HOS-ADL 81.1 (20.3) HOS-Sport 70.0 (29.3)	HOS-ADL 85.1 (6.0) HOS-Sport 72.4 (15.3)	NA	NA	No No	2/8 (25%) 2/8 (25%)	HOS-ADL 6.4 (13.1) HOS-Sport 10.6 (18)	HOS-ADL 11.4 (11.3) HOS-Sport 21.0 (16.2)	No No	HOS-ADL -0.34 (-1.37 to 0.68) HOS-Sport -0.57 (-1.61 to -0.47)
<b>Studies comparing physiotherapist-led treatment with sham or control treatment in postsurgical cohorts</b>													
<b>Bennell, et al 2017<sup>23</sup></b>	Efficacy of adding a physiotherapy rehabilitation programme to arthroscopic management of femoroacetabular impingement syndrome: a randomised controlled trial (FAIR)	IHOT-33 40.9 (15.7) HOS-Sport 50.9 (17.1)	IHOT-33 42.0 (17.5) HOS-Sport 52.1 (16.7)	IHOT-33 78.8 (17.8) HOS-Sport 83.6 (18.1)	IHOT-33 66.4 (20.5) HOS-Sport 70.8 (18.6)	IHOT-33 HOS-Sport 85.0 (17.8)	IHOT-33 HOS-Sport 86.0 (12.4)	Yes No	10/11 (90%) 3/11 (27%)	IHOT 22.5 (22.3) HOS-Sport 16.7 (24.5)	IHOT 22.5 (22.3) HOS-Sport 16.7 (24.5)	Yes No	IHOT 0.80 (0.05 to 1.55) HOS-Sport 0.71 (-0.03 to 1.45)
<b>Kemp, et al 2018<sup>26</sup></b>	A pilot randomised clinical trial of physiotherapy (manual therapy, exercise and education) for early onset hip osteoarthritis post hip arthroscopy	HOS-QOL 49.0 (25.0) IHOT 62.6 (23.7)	HOS-QOL 51.0 (15.9) IHOT 58.1 (17.3)	HOS-QOL 51 (23) IHOT 64 (19)	HOS-QOL 55 (20) IHOT 57 (23)	NA	NA	No Yes	1/9 (11%) 6/9 (67%)	IHOT-33 7 (23) HOS-Pain 10 (17)	IHOT-33 -4 (24) HOS-Pain -2 (17)	No No	IHOT 0.45 (-0.53 to 1.43) HOS-QOL 0.67 (-0.33 to 1.67)
<b>Studies comparing physiotherapist-led treatment with hip arthroscopy surgery</b>													
<b>Griffin, et al 2018<sup>29</sup></b>	Hip arthroscopy vs best conservative care for the treatment of femoroacetabular impingement syndrome (UK FASHoN): a multicentre randomised controlled trial	IHOT-33 35.6 (18.2)	IHOT-33 39.2 (20.9)	IHOT-33 49.7 (25)	IHOT-33 58.8 (27)	IHOT-33 45.6 (23)	IHOT-33 46.6 (25)	No	60/162 (37%)	IHOT-33 14.1 (25)	IHOT-33 19.3 (27)	Yes	IHOT-33 -0.21 (-0.43 to 0.01)
<b>Maneill, et al 2018<sup>8</sup></b>	Arthroscopic surgery or physical therapy for patients with femoroacetabular impingement syndrome	HOS-ADL 64.6 (14.2) HOS-Sport 53.2 (16.8) IHOT 29.4 (16.1)	HOS-ADL 65.6 (15.2) HOS-Sport 52.1 (18.1) IHOT 28.5 (16.1)	HOS-ADL 73.1 (37.1) HOS-Sport 57.1 (29.7) IHOT 44.9 (29.0)	HOS-ADL 68.4 (22.3) HOS-Sport 43.4 (29.3) IHOT 51.2 (28.1)	HOS-ADL 6M 68.4 (18.1) HOS-Sport 6M 53.1 (28.3) IHOT 6M 37.5 (27.7) HOS-ADL 12M 72.5 (27.8) HOS-Sport 12M 52.4 (27.8) IHOT 12M 43.9 (29.3)	HOS-ADL 6M 68.5 (18.7) HOS-Sport 6M 45.2 (28.3) IHOT 6M 43.8 (27.1) HOS-ADL 12M 67.7 (19.7) HOS-Sport 12M 51.8 (29.7) IHOT 12M 48.9 (29.0)	No No No	8/33 (24%) 4/33 (12%) 11/33 (33%)	HOS-ADL 18.2 (32.9) HOS-Sport 11.3 (45.9) IHOT NR	HOS-ADL 18.2 (32.9) HOS-Sport 11.3 (45.9) IHOT NR	Yes No	HOS-ADL -0.18 (-0.64 to 0.28) HOS-Sport -0.05 (-0.50 to 0.41) IHOT NC
<b>Palmer, et al 2019<sup>31</sup></b>	Arthroscopic hip surgery compared with physiotherapy and activity modification for the treatment of symptomatic femoroacetabular impingement: multicentre randomised controlled trial	HOS-ADL 65.7 (18.9)	HOS-ADL 66.1 (18.5)	HOS-ADL 69.2 (19.1)	HOS-ADL 78.4 (19.9)	NA	NA	No	6/88 (7%)	HOS-ADL 69.2 (19.1)*	HOS-ADL 78.4 (19.9)*	No	HOS-ADL -0.47 (-0.76 to -0.18)

\*Data reported as follow-up score rather than change score.  
ADL, activity of daily living; ER, external rotation; FABER, flexion-abduction-external rotation test; FAIR, flexion-abduction-internal rotation test; FU, follow-up time point; HAGOS, Copenhagen Hip and Groin Outcome Score; HHS, Hip Score; HOS, hip osteoarthritis disability outcome score; HOS, Hip Outcome Score; HSAS, hip sport and activity score; IHOT, International Hip Outcome Tool; IR, internal rotation; LEFS, lower extremity functional scale; 12M, 12-month follow-up; M, mean SD; 6M, 6-month follow-up; MIC, minimal important change; NA, not applicable; NAHS, Non-Arthritic Hip Score; NC, not calculated (insufficient data); NPRS, Numeric Pain Rating Scale; NR, not reported; PA, physical activity; PASS, Patient Acceptable State Score; PT, physiotherapy/physical therapy; QOL, quality of life; ROM, range of motion; SAME, single assessment numeric evaluation; SMD, standardised mean difference; VAS, Visual Analogue Scale.





**Figure 2** Between-group differences for physiotherapist-led treatment compared with sham/no treatment in non-surgical populations at 3 months follow-up. HOOS, Hip Osteoarthritis and disability Outcome Score; IHOT, International Hip Outcome Tool; QOL, quality of life; SMD, standardised mean difference; total, number of participants.

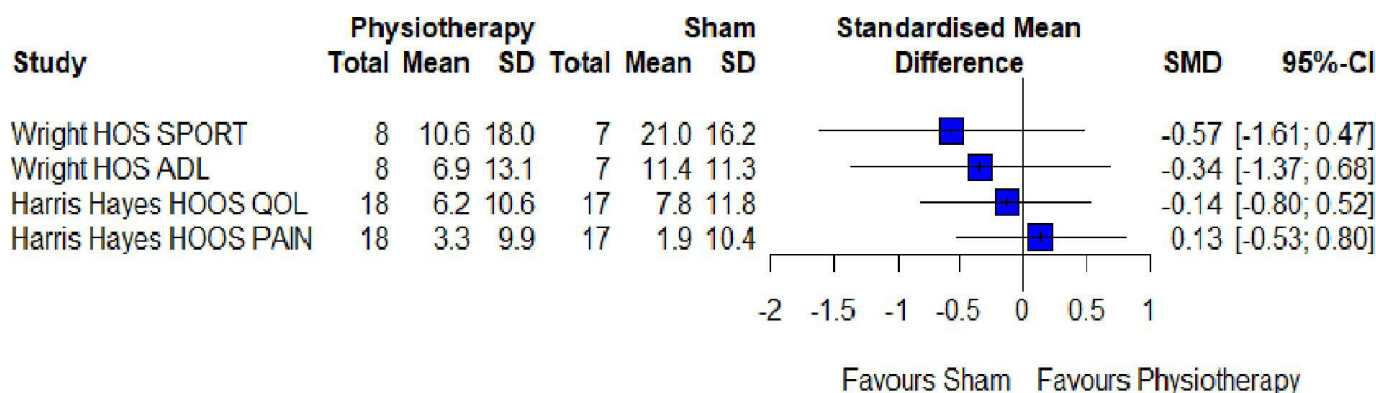
the risk of bias, the outcomes reported and the interventions performed, which limited opportunities for meta-analysis. Included studies had poor transparency in reporting of interventions, inconsistency in PROMs and methods used to measure physical impairments.

Our findings suggest that in people with hip pain, physiotherapist-led interventions may improve function and strength, however the effects on pain and QOL were unclear. There was limited evidence that interventions with targeted strengthening exercise programmes that were at least 3 months duration might have the best effect. Hip arthroscopy surgery had a small positive benefit compared with a physiotherapist-led intervention at 8–12 months. At 24 months, the level of evidence was limited indicating no difference between the hip arthroscopy surgery and physiotherapist-led interventions. Very few of the physiotherapist-led interventions in this review achieved follow-up and change scores that surpassed previously published PASS and MIC scores.

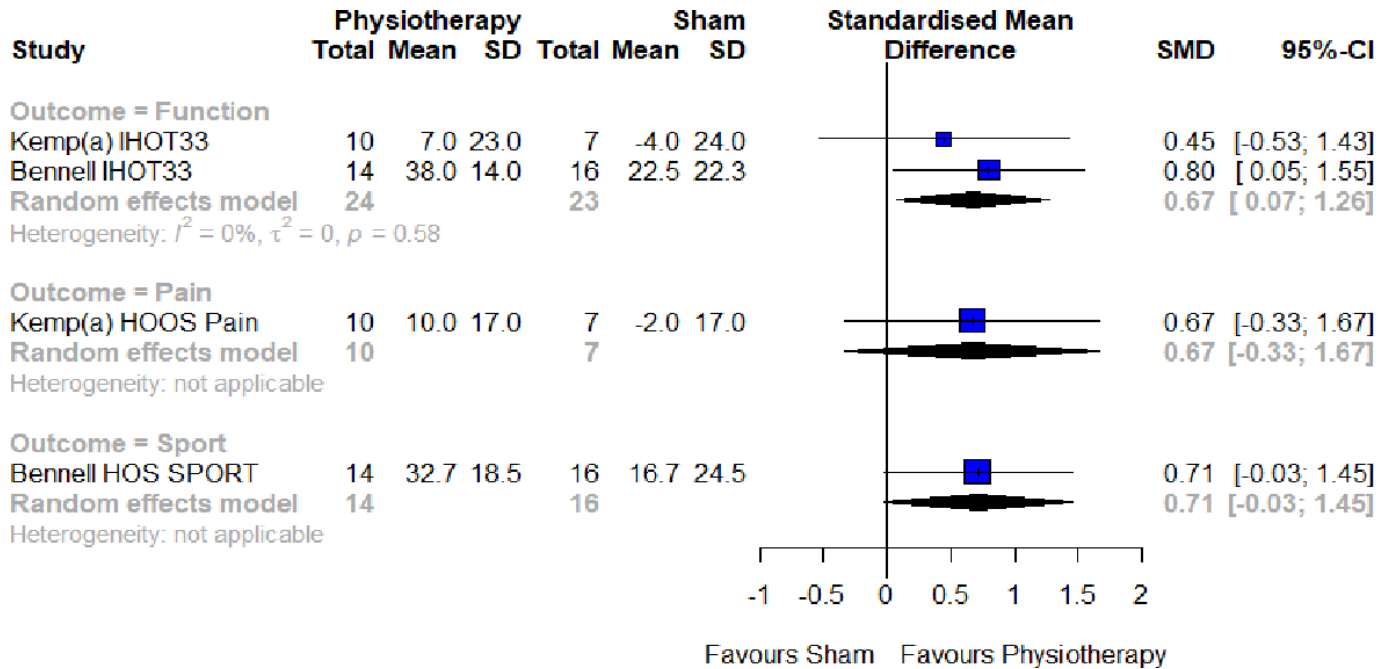
Physiotherapist-led interventions for those who had and had not undergone hip arthroscopy surgery for hip pain primarily comprised exercise therapy, where the types of exercise described included strength training, movement pattern retraining, range

of motion exercises and stretching. However, specific details of the programmes were rarely well described. The moderate effect observed for these interventions were hampered by small sample sizes and require full-scale RCTs to confirm findings. Extending the outcome measurement beyond the 3-month mark would determine whether improvements seen would be maintained in the medium-term to long-term. A recent consensus meeting reported considerable discord in the type, duration, intensity and modality of posthip arthroscopy rehabilitation provided by physiotherapists.<sup>42</sup> The consensus group suggested that full-scale RCTs are required in order to gain clarification on the composition of optimal postarthroscopic rehabilitation programmes.<sup>42</sup>

Physiotherapist-led intervention was inferior to hip arthroscopy surgery with small between-groups differences at 12-month follow-up.<sup>7 8 41</sup> Not surprisingly, despite the small difference favouring surgery, physiotherapy was far more cost-effective (£155 for physiotherapist-led treatment compared with £2372 for hip arthroscopy).<sup>7</sup> Arthroscopic surgery could be recommended as a second-line treatment for patients who have not responded adequately to a physiotherapist-led treatment programme.<sup>42</sup> However, the mean effects beyond that of physiotherapy were weak and may not be clinically meaningful.



**Figure 3** Between-group differences for physiotherapist-led treatment compared with sham/no treatment in non-surgical populations at 6 weeks follow-up. ADL, Activity of Daily Living; HOOS, Hip Osteoarthritis and disability Outcome Score; HOS, Hip Outcome Score; QOL, quality of life; SMD, standardised mean difference; total, number of participants.



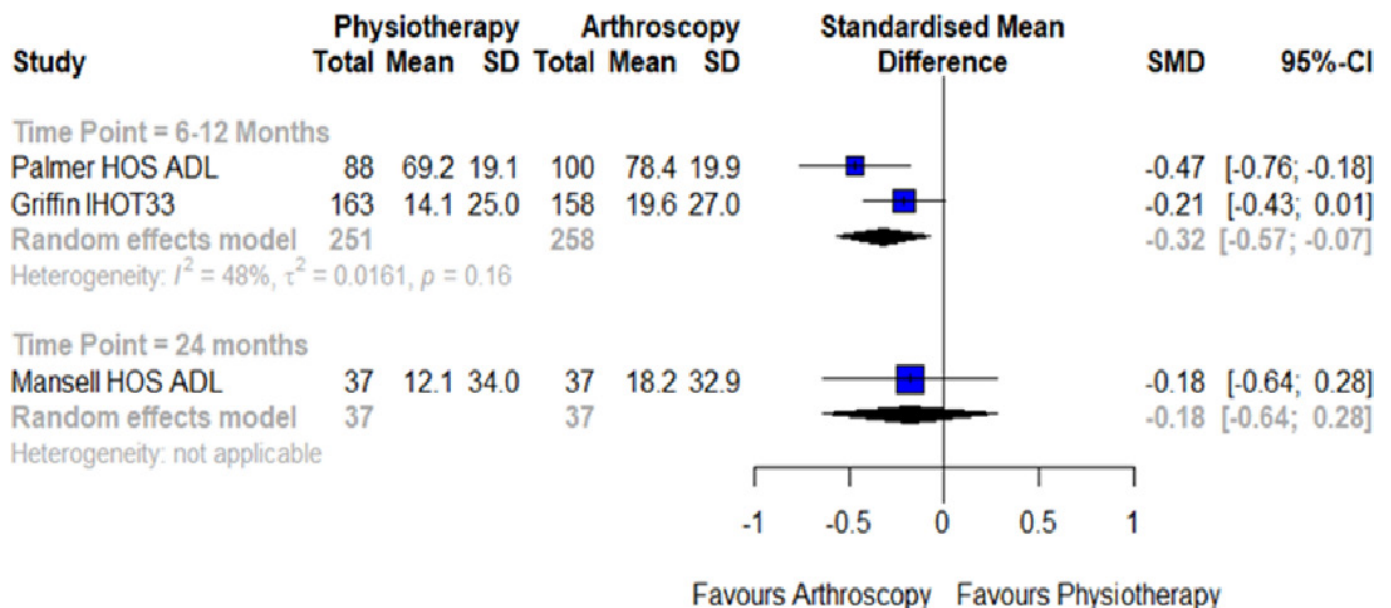
**Figure 4** Between-group differences for physiotherapist-led treatment compared with sham/no treatment in posthip arthroscopy populations at 3-month follow-up. HOOS, Hip Osteoarthritis and disability Outcome Score; HOS, Hip Outcome Score; IHOT, International Hip Outcome Tool; SMD, standardised mean difference; total, number of participants.

The within-group improvements for the physiotherapist-led interventions in these studies were modest, with only one study achieving the PASS, and they may not represent optimal treatment.<sup>18</sup> Further studies might shed more light on the relative effectiveness of surgery and physiotherapy particularly in the medium-term and long-term.<sup>43</sup>

Multimodal, physiotherapist-led interventions,<sup>7 8 30 33 34 36–38 40</sup> including exercise therapy combined with manual therapy, medication, activity modification, advice and education, were most commonly studied. Manual therapy is effective when combined with exercise for hip OA,<sup>44 45</sup> but there is debate whether

contemporary physiotherapist-led interventions for musculo-skeletal pain should include multimodal treatment additions such as manual therapy.<sup>42</sup> Further studies are required to confirm whether additional treatment elements, such as manual therapy, impart a greater benefit than exercise-therapy alone for young and middle-aged adults with hip pain.

Physiotherapist-led interventions on physical impairments had variable effects. For hip range of motion, the largest positive effects pre-physiotherapist-led to post-physiotherapist-led treatment were seen following a 3-month intervention strengthening exercise, manual therapy and education.<sup>40</sup> The greatest



**Figure 5** Between-group differences for physiotherapist-led treatment compared with hip arthroscopy surgery. ADL, Activity of Daily Living; HOS, Hip Outcome Score; IHOT, International Hip Outcome Tool; SMD, standardised mean difference; total, number of participants.

**Table 3** Summary of results of within-group standardised paired differences for physiotherapist-led treatment in non-operative patients (randomised and non-randomised studies)

Study	Title	Baseline M (SD) PT	Prim FU M (SD) PT	Within-group SPD (95% CI) for primary outcomes (positive SPD favours postintervention improvement)
<b>Randomised studies</b>				
Grant, <i>et al</i> 2017 <sup>34</sup>	The HAPI 'Hip Arthroscopy Pre-habilitation Intervention' study: does prehabilitation affect the outcomes in patients undergoing hip arthroscopy for femoroacetabular impingement?	Abduction strength 16.6 (11.8) Adduction strength 13.4 (4.8) Flexion strength 27.3 (23.1) ER strength 15.3 (5.3) Knee extension strength 37.1 (23.0)	Abduction strength 20.9 (11.6) Adduction strength 19.3 (9.9) Flexion strength 32.9 (16.6) External rotation strength 19.0 (6.3) Knee extension strength 49.0 (40.2)	Abduction strength not calculated as follow-up dataset not complete Adduction strength not calculated as follow-up dataset not complete Flexion strength not calculated as follow-up dataset not complete External rotation strength not calculated as follow-up dataset not complete Knee extension strength not calculated as follow-up dataset not complete
Griffin, <i>et al</i> 2018 <sup>7</sup>	Hip arthroscopy vs best conservative care for the treatment of femoroacetabular impingement syndrome (UK FASHIoN): a multicentre randomised controlled trial	IHOT-33 35.6 (18.2)	IHOT-33 49.7 (25)	IHOT-33 0.63 (0.46 to 0.79)
Harris-Hayes, <i>et al</i> 2016 <sup>39</sup>	Movement pattern training to improve function in people with chronic hip joint pain: a feasibility randomised clinic trial	HOOS Symptoms 75.0 (17.0) HOOS-Pain 78.2 (12.3) HOOS-ADL 90.7 (9.9) HOOS-Sport 77.1 (17.5) HOOS-QOL 65.1 (13.0) Abduction strength 6.7 (1.8) ER 0 strength 2.9 (0.9) ER 90 strength 3.4 (0.8) Flexion in single leg squat 67.4 (14.0) Adduction in single leg squat 20.2 (6.6) IR in single leg squat 2.5 (7.7)	HOOS Symptoms 85.0 (13.6) HOOS-Pain 81.5 (13.2) HOOS-ADL 93.5 (10.9) HOOS SP 84.6 (19.6) HOOS-QOL 71.3 (18.3) Abduction strength 7.2 (2.3) ER 0 strength 3.2 (0.8) ER 90 strength 3.9 (0.9) Flexion in single leg squat 61.7 (15.9) Adduction in single leg squat 17.6 (5.7) IR in single leg squat 2.7(6.3)	HOOS Symptoms 0.61 (0.11 to 1.12) HOOS-Pain 0.25 (-0.22 to 0.72) HOOS-ADL 0.26 (-0.21 to 0.73) HOOS SP 0.38 (-0.09 to 0.86) HOOS-QOL 0.36 (-0.11 to 0.84) Abduction strength 0.23 (-0.24 to 0.70) ER 0 strength 0.10 (-0.36 to 0.56) ER 90 strength 0.78 (0.25 to 1.31) Flexion in single leg squat 0.36 (0.11 to 0.84) Adduction in single leg squat 0.40 (-0.08 to 0.88) IR in single leg squat -0.02 (-0.49 to 0.44)
Kemp <i>et al</i> , 2018 <sup>40</sup>	The Physiotherapy for Femoroacetabular Impingement Rehabilitation Study: a pilot randomised controlled trial	IHOT 60(26) HOOS-QOL 54(21) Adduction strength 0.85 (0.17) Abduction strength 0.94 (0.23) Extension 0.92 (0.28) ER strength 0.48 (0.11) Flexion ROM 109(14) Single leg hop 1.14 (0.26) Side bridge 98(42)	IHOT 87(12) HOOS-QOL 76(13) Adduction strength 1.1 (0.22) Abduction strength 1.16 (0.23) Extension strength 1.17 (0.32) ER strength 0.57 (0.19) Flexion ROM 123(9) Single leg hop 1.34 (0.32) Side bridge 98(35)	IHOT 1.14 (0.53 to 1.75) HOOS-QOL 1.14 (0.53 to 1.75) Adduction strength 1.19 (0.57 to 1.81) Abduction strength 0.91 (0.35 to 1.48) Extension strength 0.79 (0.24 to 1.33) ER strength 0.52 (0.01 to 1.03) Flexion ROM 1.08 (0.49 to 1.68) Single leg hop 0.65 (0.12 to 1.17) Side bridge 0.95 (0.38 to 1.53)
Mansell, <i>et al</i> 2018 <sup>8</sup>	Arthroscopic surgery or physical therapy for patients with femoroacetabular impingement syndrome	HOS-ADL 64.6 (14.2) HOS-Sport 53.2 (16.8)	HOS-ADL 73.1 (37.1) HOS-Sport 57.1 (29.7)	HOS-ADL 0.26 (-0.06 to 0.57) HOS-Sport 0.15 (-0.16 to 0.46)
Palmer, <i>et al</i> 2019 <sup>41</sup>	Arthroscopic hip surgery compared with physiotherapy and activity modification for the treatment of symptomatic femoroacetabular impingement: multicentre randomised controlled trial	Hip Flexion ROM 95.7 (19.1) Hip extension ROM 17.9 (7.9) Hip Abduction ROM 27.5 (11.9) Hip adduction ROM 21.6 (7.9) Hip ER ROM 25.0 (11.8) Hip IR ROM 24.0 (11.2)	Hip Flexion ROM 99.7 (17.5) Hip extension ROM 15.7 (8.0) Hip Abduction ROM 29.6 (11.7) Hip adduction ROM 23.2 (8.9) Hip ER ROM 27.4 (9.7) Hip IR ROM 28.9 (11.2)	Hip Flexion ROM not calculated as follow-up dataset not complete Hip extension ROM not calculated as follow-up dataset not complete Hip Abduction ROM not calculated as follow-up dataset not complete Hip adduction ROM not calculated as follow-up dataset not complete Hip ER ROM not calculated as follow-up dataset not complete Hip IR ROM not calculated as follow-up dataset not complete
Smeatham, <i>et al</i> 2017 <sup>37</sup>	Does treatment by a specialist physiotherapist change pain and function in young adults with symptoms from femoroacetabular impingement? A pilot project for a randomised controlled trial	NAHS Total 50.1 (19.1) HOS-ADL 69 (39) HOS-Sport 51.5 (19.7)	NAHS Total 62.2 (16.1) HOS-ADL 90 (27) HOS-Sport 68 (21.6)	NAHS Total 0.64 (0.09 to 1.20) HOS-ADL 0.57 (0.03 to 1.12) HOS-Sport 0.75 (0.18 to 1.33)
Wright, <i>et al</i> 2016 <sup>38</sup>	Non-operative management of femoroacetabular impingement: a prospective, randomised controlled clinical trial pilot study	HOS-ADL 74.3 (13.1) HOS-Sport 59.4 (18.0) Depth squat 54.1 (12.5) Triple hop 11.4 (5.1) Flexion ROM 100.3 (20.5) Flexion strength 9.9 (4.2)	HOS-ADL 81.1 (20.3) HOS-Sport 70.0 (29.3) Depth squat 43.1 (15.5) Triple hop 13.3 (5.3) Flexion ROM 122.4 (18.8) Flexion strength 13.2 (4.4)	HOS-ADL 0.34 (-0.37 to 1.05) HOS-Sport 0.37 (-0.35 to 1.08) Depth squat 0.69 (0.08 to 1.46) Triple hop 0.32 (-0.39 to 1.04) Flexion ROM 1.00 (0.15 to 1.84) Flexion strength 0.68 (-0.08 to 1.45)
<b>Non-randomised studies</b>				
Coppack, <i>et al</i> 2016 <sup>30</sup>	Physical and functional outcomes following multidisciplinary residential rehabilitation for prearthritic hip pain among young active UK military personnel	HAGOS Pain 37.7 (20.9) HAGOS Symptoms 45.8 (23.2) HAGOS ADL 32.2 (24.1) HAGOS Sport 51.0 (28.1) HAGOS PA 84.7 (24.9) HAGOS QOL 69.5 (24.0) Y BALANCE 240.5 (26.9) Flexion ROM 110.2 (24.3) IR ROM 25.2 (13.7)	HAGOS Pain 35.1 (23.7) HAGOS Symptoms 46.3 (24.2) HAGOS ADL 31.0 (24.7) HAGOS Sport 48.5 (28.6) HAGOS PA 77.5 (31.2) HAGOS QOL 64.9 (23.3) Y BALANCE 256.3 (20.8) Flexion ROM 116.7 (23.3) IR ROM 29.8 (12.4)	HAGOS Pain -0.11 (-0.42 to 0.19) HAGOS Symptoms 0.02 (-0.29 to 0.33) HAGOS ADL -0.05 (-0.36 to 0.26) HAGOS Sport -0.09 (-0.40 to 0.22) HAGOS PA -0.25 (-0.56 to 0.07) HAGOS QOL -0.19 (-0.50 to 0.12) Y BALANCE 0.63 (0.29 to 0.97) Flexion ROM 0.27 (-0.05 to 0.58) IR ROM 0.34 (0.03 to 0.66)

Continued

Table 3 Continued

Study	Title	Baseline M (SD) PT	Prim FU M (SD) PT	Within-group SPD (95% CI) for primary outcomes (positive SPD favours postintervention improvement)
Emara, <i>et al</i> 2011 <sup>31</sup>	Conservative treatment for mild femoroacetabular impingement	HHS 72(6) NAHS 72(4) VAS 6 (1) Flexion ROM 95.0 (0.4) Extension ROM 4.0 (1.6) Abduction ROM 37.0 (0.4) Adduction ROM 17.0 (7.0) ER in flexion ROM 28.5 (0.5) ER in extension ROM 25.3 (0.3) IR in flexion ROM 9.4 (0.3) IR in extension ROM 15.8 (0.4)	HHS 91(4) NAHS 90(5) VAS 3 (1) Flexion ROM 88.0 (3.5) Extension ROM 3.7 (2.2) Abduction ROM 36.0 (1.4) Adduction ROM 17.0 (9.0) ER in flexion ROM 28.4 (1.2) ER in extension ROM 24.5 (1.0) IR in flexion ROM 11.3 (0.5) IR in extension ROM 15.7 (0.7)	HHS 3.52 (2.65 to 4.38) NAHS 3.85 (2.91 to 4.78) VAS 2.94 (2.19 to 3.68) Flexion ROM -2.07 (-2.64 to -1.50) Extension ROM -0.15 (-0.47 to 0.17) Abduction ROM -0.78 (-1.15 to -0.41) Adduction ROM 0.00 (-0.32 to 0.32) ER in flexion ROM -0.09 (-0.41 to 0.23) ER in extension ROM -0.88 (-1.26 to -0.50) IR in flexion ROM 4.27 (3.24 to 5.29) IR in extension ROM -0.16 (-0.48 to 0.16)
Guenther, <i>et al</i> 2017 <sup>35</sup>	A pre-operative exercise intervention can be safely delivered to people with femoroacetabular impingement and improve clinical and biomechanical outcomes	HOOS Symptoms 56.1 (13.2) HOOS-Pain 64.1 (12.3) HOOS-ADL 73.0 (14.4) HOOS-Sport 51.7 (12.2) HOOS-QOL 35.3 (17.2) Abduction strength 1.53 (0.35) Adduction strength 1.40 (0.38) Extension strength 1.81 (0.46) Flexion strength 1.89 (0.45) ER strength 0.75 (0.23) IR strength 0.76 (0.36) Timed stair climb test 2.96 (0.66)	HOOS Symptoms 63.9 (14.6) HOOS-Pain 72.5 (12.3) HOOS-ADL 83.4 (11.0) HOOS-Sport 63.4 (14.0) HOOS-QOL 42.8 (22.0) Abduction strength 1.67 (0.34) Adduction strength 1.53 (0.39) Extension strength 1.93 (0.50) Flexion strength 2.04 (0.43) ER strength 0.77 (0.18) IR strength 0.89 (0.36) Timed stair climb test 2.61 (0.46)	HOOS Symptoms HOOS-Pain HOOS-ADL HOOS-Sport HOOS-QOL Abduction strength 0.39 (-0.07 to 0.84) Adduction strength 0.32 (-0.13 to 0.77) Extension strength 0.24 (-0.21 to 0.68) Flexion strength 0.33 (-0.12 to 0.78) ER strength 0.09 (-0.35 to 0.53) IR strength 0.35 (-0.10 to 0.80) Timed stair climb test 0.57 (0.10 to 1.05)
Hunt, <i>et al</i> 2012 <sup>32</sup>	Clinical outcomes analysis of conservative and surgical treatment of patients with clinical indications of pre-arthritis, intra-articular hip disorders	HHS 61.3±13 WOMAC 29.2±16 NAHS 63.2±14	HHS 78.9±14 WOMAC 13.5±14 NAHS 81.6±12	HHS 1.26 (0.76 to 1.77) WOMAC 1.01 (0.54 to 1.47) NAHS 1.36 (0.84 to 1.89)

ADL, activity of daily living; ER, external rotation; FU, follow-up time point; HAGOS, Copenhagen Hip and Groin Outcome Score; HHS, Harris Hip Score; HOOS, Hip Osteoarthritis and disability Outcome Score; IR, internal rotation; M, mean SD; MCS, emotional function subscale; NAHS, Non-Arthritic Hip Score; NPRS, Numeric Pain Rating Scale; NR, not reported; PA, physical activity; PCS, physical function subscale; PT, physiotherapy/physical therapy; QOL, quality of life; ROM, range of motion; SF-12, Short Form-12 Questionnaire; SMD, standardised mean difference; VAS, Visual Analogue Scale.

hip muscle strength gain was seen with a strengthening exercise programme of 3 months duration,<sup>40</sup> and largest in hip adductor muscles. Greater hip adductor strength following hip arthroscopy is associated with better hip-related QOL,<sup>46</sup> suggesting that it may be an important target. However, this was a pilot study, and the most effective type, dose and progression of exercise is unknown. The American College of Sports Medicine<sup>47</sup> guidelines for exercise prescription contain information about the dosage, volume and progression of exercises that may be useful for clinicians and researchers when developing strength programmes for patients with hip pain. In studies measuring changes in functional task performance, had positive effects<sup>30 35 40</sup> with movement retraining and functional exercise programmes. These programmes may improve patient self-efficacy as well as increase the capacity for load, thus enabling participation in more challenging activity. Larger, future studies including evaluating the potential of effect mediators and moderators may provide insight into the most effective physiotherapist-led interventions to improve physical impairments.

Returning to pre-injury sport and activity is important to young and middle-aged people with hip pain, and often the reason they seek surgical and/or non-surgical treatment.<sup>10 48</sup> However, only two studies in this review had a specific return to sport/return to physical activity component within the physiotherapist-led intervention.<sup>33 40</sup> Only 17% of people returned to optimal performance and full sports participation at 33±16 months following hip arthroscopy.<sup>48</sup> Given the importance of returning to sport in this active patient group and the disappointing rates of returning to optimal performance reported,<sup>48</sup> future studies should incorporate key functional and sporting components.<sup>49</sup> These could include: valid and consistent definitions of what comprises a successful return to sport and return to activity<sup>50</sup>; fully powered RCTs that include a specific, targeted return to sport programme throughout the duration of the intervention<sup>50</sup> and inclusion of return to sport outcomes as

a continuum.<sup>40</sup> Until physiotherapist-led interventions include high-quality return to sport elements, outcomes are unlikely to improve beyond those reported by Ishoi *et al*.<sup>48</sup>

Transparency and reproducibility are critical when reporting the efficacy of clinical interventions. Guidelines such as the Consensus on Exercise Reporting Template<sup>51</sup> and Template for Intervention Description and Replication checklist<sup>52</sup> should be used in all trials reporting interventions to ensure adequate transparency and reproducibility. In addition, describing targeted strengthening interventions should use detailed procedures such as those described by Toigo and Boutellier.<sup>53</sup> The documentation of adherence to exercise programmes is also critical. Such guidelines allow researchers to evaluate interventions and clinicians to reproduce efficacious interventions in clinical practice. Very few studies used these guidelines to report interventions in the current review. As such, it was not possible to pool findings to complete a meta-analysis, limiting the scope of the review. We recommend that future studies report physiotherapist-led interventions using guidelines such as those described above to maximise transparency and utility of study findings by researchers and clinicians alike.

This review contains several limitations that should be acknowledged. First, the methodological quality of the studies was variable, with only 4/14 (29%) studies considered to have a low risk of bias. In the studies with a higher risk of bias, inflated effect sizes are possible, raising questions about the strength of the findings of these studies. Second, most of the included RCTs were pilot studies and as such were underpowered to detect statistically significant differences between groups. In addition, as there were no fully powered studies comparing physiotherapist-led interventions with sham interventions, adequately powered studies that undertake a head-to-head comparison of physiotherapist-led interventions are required to determine the optimal management strategies for hip pain in young and middle-aged adults. Furthermore, the terminology used to describe hip

## Summary box

## What is already known?

- ▶ Hip-related pain is common in young, active adults.
- ▶ Non-surgical treatments such as physiotherapist-led treatments should be first-line treatment for musculoskeletal conditions including hip-related pain, but effectiveness of these treatments is unclear.

## What are the new findings?

- ▶ There is a paucity of literature in this field.
- ▶ Physiotherapist-led interventions improve function and strength.
- ▶ Effects of physiotherapist-led interventions on pain and quality of life are uncertain.
- ▶ Targeted strengthening exercise programmes and at least 3-month duration might have the best effect.
- ▶ Hip arthroscopy surgery had a small positive benefit compared with a physiotherapist-led intervention at 8–12 months.
- ▶ At 24 months, there was limited evidence suggesting no difference between groups.

disorders is not clear, and the populations that were included in this review may be heterogenous. The recent Zurich consensus statement on hip-related pain has provided some guidance in classifying hip disorders as FAI syndrome, acetabular dysplasia and ‘other’.<sup>9</sup> At present, there are not enough studies published to be able to analyse data separately for each of these three classifications. However, as the field evolves, this may be an approach that is appropriate for future reviews. As with all reviews of intervention studies, publication bias may have existed where studies with negative findings were not published. We also excluded studies not written in English, which may have led to inclusion bias. Finally, the PROMs and physical impairment-based outcome measures used in the studies were inconsistent, which limited the pooling of data. A recent consensus<sup>54</sup> determined that the HAGOS and IHOT were the most appropriate PROM for use in young and middle-aged people with hip pain, and future studies using these measures may make stratification and pooling of data based on these measures in future reviews possible. Furthermore, it is not yet clear what is considered an acceptable level of improvement in a patient’s condition. We compared the findings of our review to previously published PASS and MIC scores, to provide some context to clinical relevance of the effects reported. We acknowledge that the previously published PASS and MIC scores were determined in studies of posthip arthroscopy and posthip arthroplasty patients. It is not known what constitutes a PASS and MIC in non-surgical cohorts of people with hip-related pain. The inclusion of PASS questions at specific time points in future studies may help determine whether patients are gaining acceptable improvement when undergoing physiotherapist-led interventions.

## CONCLUSION

There were no full-scale RCTs comparing physiotherapist-led interventions with other non-surgical treatments or sham treatments. The risk of bias in included studies, as well as limitations in included study methodology should be considered in the interpretation of the results of this systematic review. Physiotherapist-led interventions may improve pain and function in young and middle-aged adults experiencing hip pain, including those

with FAI syndrome. There was limited evidence of larger effects for interventions that included targeted strengthening exercise programmes and were of 3 months duration. Hip arthroscopy surgery had a weak positive effect compared with a physiotherapist-led intervention at 8–12 months. Future full-scale RCTs undertaking a head-to-head comparison of physiotherapist-led interventions for hip pain are required.

**Correction notice** This article has been corrected since it published Online First. An ORCID ID has been added for author Steven Chang.

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