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Abstract

Patellofemoral Pain Syndrome (PFPS) is a common musculoskeletal disorder typically occurring in physically active people aged 40 years and younger, causing pain, functional deficits and lower limb weakness. Traditional treatment has been aimed at strengthening the knee, however recent research suggests the muscles around the hip also play an important role in the development and continuity of Patellofemoral Pain Syndrome.

Purpose: To investigate the effectiveness of the addition of hip strengthening exercises to standard physiotherapy treatment (knee strengthening and stretching exercises) on reducing pain, and enhancing strength and function when compared to standard physiotherapy treatment alone in adults with Patellofemoral Pain Syndrome.

Method: A systematic search of Cochrane, CINAHL, Embase, MEDLINE®, PEDro and SportDiscus was conducted. Studies of participants aged 18 to 44, diagnosed with Patellofemoral Pain Syndrome by a healthcare practitioner, or reporting peripatellar or retropatellar pain with common functional tasks, were included. A critical appraisal, using the Critical Appraisal Skills Program for Randomised Controlled Trials (CASP) was used to assess methodological quality.

Results: Five randomised controlled trials of varying methodological quality met the inclusion criteria. The participants in these studies were aged between 18 to 40 years of age. The duration of the intervention ranged from four to six weeks consisting of 12 to 30 supervised exercise sessions. Studies used varying outcome measures for each of the three outcomes. Overall, the studies demonstrated that the addition of hip strengthening exercises to standard physiotherapy care consistently improved pain and function, but the impact on strength was variable.

Conclusion: Previously, only a small number of studies have looked at the addition of hip exercises to standard physiotherapy care for treatment of Patellofemoral Pain Syndrome. While there is a growing body of evidence for the efficacy of hip strengthening exercises for Patellofemoral Pain Syndrome, this is constrained by bias towards female participants, lack of true controls in most studies, and low methodological quality of studies overall. Hip exercises added to standard physiotherapy care shows potential as a treatment method for improving outcomes of pain and function in adults with Patellofemoral Pain Syndrome.

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ABSTRACT

Patellofemoral Pain Syndrome (PFPS) is a common musculoskeletal disorder typically occurring in physically active people aged 40 years and younger, causing pain, functional deficits, and lower limb weakness. Traditional treatment has been aimed at strengthening the knee, however, recent research suggests the muscles around the hip play an important role in the development and continuity of Patellofemoral Pain Syndrome. **Purpose:** To investigate the effectiveness of the addition of hip strengthening exercises to standard physiotherapy treatment (knee strengthening and stretching exercises) on reducing pain and enhancing strength and function when compared to standard physiotherapy treatment alone in adults with Patellofemoral Pain Syndrome. **Method:** A systematic search of Cochrane, CINAHL, Embase, MEDLINE®, PEDro, and SportDiscus was conducted. Studies of participants aged 18 to 44, diagnosed with PFPS by a healthcare practitioner, or reporting peripatellar or retropatellar pain with common functional tasks, were included. A critical appraisal, using the Critical Appraisal Skills Program for Randomised Controlled Trials (CASP) was used to assess methodological quality. **Results:** Five randomised controlled trials of varying methodological quality met the inclusion criteria. The participants in these studies were aged between 18 and 40 years of age. The duration of the intervention ranged from four to six weeks consisting of 12 to 30 supervised exercise sessions. Studies used varying outcome measures for each of the three outcomes. Overall, the studies demonstrated that the addition of hip strengthening exercises to standard physiotherapy care consistently improved pain and function, but the impact on strength was variable. **Conclusion:** Only a small number of studies have looked at the addition of hip exercises to standard physiotherapy care for the treatment of Patellofemoral Pain Syndrome. While there is a growing body of evidence for the efficacy of hip strengthening exercises for Patellofemoral Pain Syndrome, the studies tend to be constrained by bias towards female participants, lack of true controls, and low methodological quality of studies overall. Hip exercises added to standard physiotherapy care shows potential as a treatment approach for improving outcomes of pain and function in adults with Patellofemoral Pain Syndrome.

BACKGROUND

Patellofemoral Pain Syndrome (PFPS) is a common musculoskeletal disorder typically occurring in physically active people aged 40 years and younger.¹ Prevalence ranges for PFPS are reported to be from 15% to 33% in adults, and 21% to 45% in active adolescents.² This syndrome is considered the most common overuse injury of the lower limb in active individuals, accounting for 11% to 17% of all knee pain presentations in general practice.^{1,3} Common symptoms include diffuse peripatellar or localised retropatellar pain during activities such as running, moving up or down stairs, squatting, and sitting with knees bent for prolonged periods.⁴ The clinical presentation of PFPS often includes muscular weakness and altered lower limb biomechanics.⁵

Traditional treatment methods aim to improve patellar alignment and strengthen muscles surrounding and acting on the knee joint.⁶ However, a recent study by Santos et al. and best practice guidelines developed by Barton et al. suggest that hip biomechanics may play a major role in the development and continuity of PFPS.^{5,7} Hip muscle strengthening to counteract excessive hip flexion,

adduction, and internal rotation, which exert stress on the patellofemoral joint, has been proposed as additional treatment for PFPS.^{5,8}

A systematic review by Santos et al. explored the effectiveness of hip muscle strengthening in patients with PFPS.⁵ The results suggested that hip strengthening exercises were effective at improving pain and function, however, in addition to a limited number of databases and search terms being used, two included studies used comparator groups that made it difficult to clearly identify the specific effects of the hip strengthening exercises.^{5,9,10} A third study by Nakagawa et al. added transversus abdominus exercises to the hip strengthening exercises, also potentially obscuring the true effects of hip strengthening.¹¹

The functional problems associated with PFPS in young adults are increasing, particularly with the rise in physical activity to counteract sedentary lifestyles and to promote health and wellbeing.¹² Hence, this current systematic review is important in order to update the findings of Santos et al. through a more comprehensive search, rigorous selection and appraisal processes, and summary of updated high-quality literature.⁵ This can then be integrated with previous knowledge on the effect of hip strengthening exercises on PFPS to further assist clinicians with important clinical decision making. The systematic review research question was: 'In adults with Patellofemoral Pain Syndrome, what is the effectiveness of the addition of hip strengthening exercises to standard physiotherapy treatment on pain, strength, and function, when compared with standard physiotherapy care alone?'

METHODS

Search Strategy

This review was conducted and reported in line with the Preferred Reporting Items for Systematic Reviews and Meta Analyses (PRISMA) Statement.¹³ In September 2017, a search was conducted in six electronic databases: Cochrane, Cumulative Index to Nursing and Allied Health Literature (CINAHL), Embase, MEDLINE®, Physiotherapy Evidence Database (PEDro), and SportDiscus. Each database was searched from inception to the date of the search (September 2017). Only studies published in English were included. Table 1 outlines the Participant, Intervention, Comparator, Outcomes (PICO) components of the search question and keywords used in the search strategy. Details of the search strategy are included in Appendix 1.

Table 1: PICO Search Strategy

	Definition
Population	Adults (ages 18-44) with Patellofemoral Pain Syndrome (PFPS)
Intervention	Hip strengthening exercises in addition to standard physiotherapy care
Comparator	Standard physiotherapy care
Outcome	Pain, strength, and function

Eligibility Criteria

Studies were only considered for inclusion based on design if they were controlled trials, including randomised, pseudo-randomised, and non-randomised controlled clinical trials. In addition, the reference lists of all systematic reviews and the final studies were searched for further relevant trials. Other primary research evidence including case studies/series, observational studies, cohort studies, case-control studies, and all qualitative research evidence was not considered for inclusion.

Population

The population of included studies was limited to adults aged 18 to 44 years who had been diagnosed with PFPS by a healthcare practitioner or complained of: 1) the presence of anterior retropatellar knee pain during at least two of the following activities: ascending/descending stairs, squatting, hopping/running, kneeling, and prolonged sitting, 2) insidious onset of symptoms unrelated to a trauma, 3) pain on compression or palpation of the patellar and facets.^{4,5} Study exclusion occurred in the presence of a current significant injury affecting the lower extremity, a history of other knee pathology, any ankle, hip, or lower back/sacroiliac pain, or the use of corticosteroids and/or anti-inflammatory medication.

Intervention

Studies were considered for inclusion if hip strengthening exercises targeting at least two of the following muscle groups were utilised as an adjunct to standard PFPS physiotherapy treatment directed at the knee: hip external rotators, hip abductors, hip extensors. Studies with comparator groups that focused solely on hip stretching exercises or which included abdominal and other trunk muscle strengthening exercises were excluded.

Outcome Measures

Outcomes of interest were at least one or more of the following: pain, function, and strength. Only studies incorporating validated outcome measures were accepted (e.g. visual analogue scale [VAS] for pain, Lower Extremity Functional Scale [LEFS] for function, hand-held dynamometer for strength).

Study Selection

Two reviewers (FG, BJ) independently undertook the search of each database to ensure consistency and reproducibility. The reviewers compared the number of 'hits' after each database search; where discrepancies occurred, the error was identified and the search re-run until there were no errors. The results from each database were exported into the industry standard bibliographic software tool Endnote™ to separate database folders. All folders were then combined and duplicates removed. The remaining studies were imported to Covidence, a data management software for systematic reviews, where two rounds of screening took place. In the first round, both reviewers (FG, BJ) independently screened all study titles and abstracts. Any disagreement that arose was resolved using the following system: Maybe + Yes = Yes, Maybe + Maybe = Yes, Maybe + No = No, Yes + No = conflict. Both reviewers met to discuss and resolve any conflicts. Full text was obtained for all studies where there was insufficient detail in the abstract to determine eligibility, where there was no available abstract, or where the study was likely to be included based on title and abstract. A second round of screening of the full text of each study against the eligibility criteria was then undertaken by the same two reviewers (FG, BJ) and reasons for exclusions were recorded. Discrepancies about eligibility that could not be resolved by the two reviewers were resolved via a face-to-face discussion with the remaining two reviewers (KH, CE).

Risk of Bias

The methodological quality of each included study was assessed independently by all four reviewers using the Critical Appraisal Skills Program (CASP) Randomised Controlled Trial critical appraisal tool.¹⁴ Initially, reviewers met to discuss the agreed interpretation of items in the CASP, where a scoring system of, yes = 1, no and can't tell = 0, was used. Reviewers then met to compare findings of each CASP and any disagreements were discussed until consensus was reached. Potential methodological and reporting biases not formally assessed by the CASP were also independently considered by two team reviewers and then discussed to resolve any conflicts.

The hierarchy of all studies was assessed according to the National Health and Medical Research Council (NHMRC) designation of levels of evidence.¹⁵ An agreed interpretation of results was previously established by all research team members and disagreements were resolved through discussion with the project supervisor (MM).

Data Extraction

The data were extracted by two reviewers (KH, CE) and collated into a specifically customised template which included information relating to the population, intervention, comparator, outcome measures, and results. The process of data extraction using this template was previously tested with one study by both reviewers together to ensure data would be interpreted and extracted consistently. Data extraction of the remaining studies was then completed independently, and findings were compared between the same two reviewers before the process of condensing and refining was implemented. Any disagreements that arose through the extraction process were resolved through face-to-face discussion with all reviewers. A narrative analysis of included studies was subsequently performed as the small number of studies and variability in outcome measures meant a meta-analysis was not appropriate. The statistical effects of the interventions on validated measures for pain, function, and strength were presented and compared using p-values with an alpha level set at 0.05 or using 95% confidence intervals.

Data Synthesis

A review team utilised the NHMRC FORM methodology to grade and provide a framework to synthesise the evidence from the literature.¹⁵ The NHMRC FORM methodology considers the evidence provided in all studies to assist in the development of a specific recommendation. There are five main components: 1) quantity and quality of the evidence, 2) consistency, 3) clinical impact, 4) generalisability, and 5) applicability. This framework provided a basis for evidence-based recommendations for implementation in clinical practice and identified where care may need to be taken in the application of the findings.¹⁵

RESULTS

Search Results

The search strategy generated 200 'hits' with an additional one record being identified through other sources. Following removal of duplicates, title and abstract screening, and screening of full-text versions, five studies were included in the review. The authors also followed-up a Persian study by Soleimani et al. with an English abstract, but no full text version in English was available.¹⁶ There was no response to a request from the authors, so the study was excluded. Figure 1 provides an overview of the literature selection process.

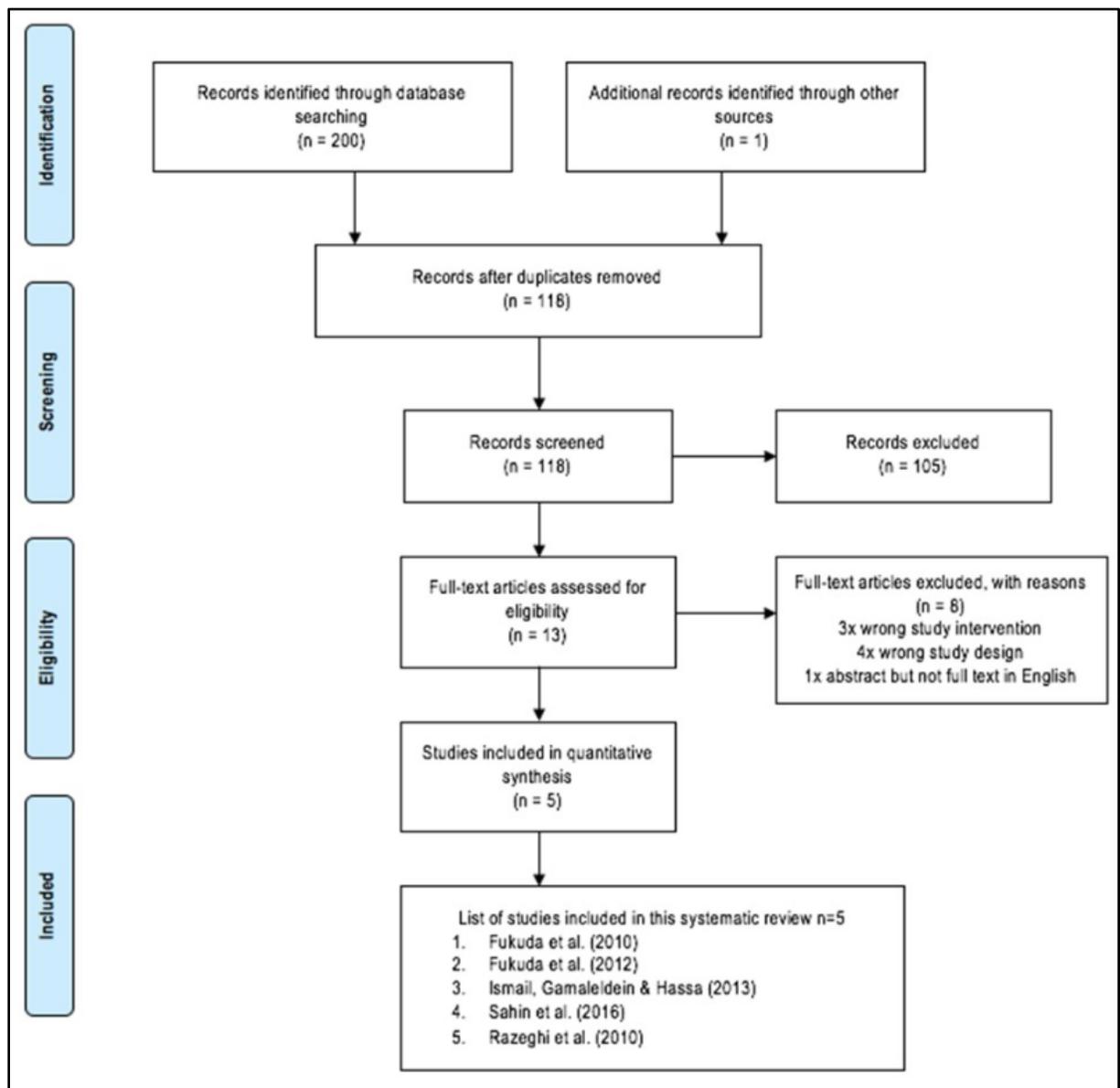


Figure 1

Ranking and Methodological Quality

An overview of the NHMRC designation of levels of evidence and a summary of the collated findings and consensus agreement from the four researchers for the critical appraisal for individual studies is presented in Table 2.¹⁴ All studies scored between five to nine out of a possible total score of 10 (Table 2).¹³

Table 2: CASP Scores of each included study

Critical Appraisal Skills Program for Randomised Controlled Trials (CASP) Question Number	Study				
	Fukuda et al. ¹⁷	Fukuda et al. ¹⁸	Ismail et al. ¹⁹	Sahin et al. ²⁰	Razeghi et al. ²¹
	RCT (NHMRC III-1)	RCT (NHMRC III-1)	RCT (NHMRC III-1)	RCT (NHMRC III-1)	RCT (NHMRC III-1)
Q1: Did the trial address a clearly focused issue?	1	1	1	1	1
Q2: Was the assignment of patients to treatment groups randomised?	1	1	1	1	0
Q3: Were all of the patients who entered the trial properly accounted for at its conclusion?	1	1	1	0	0
Q4: Were patients, health workers and study personnel 'blind' to treatment?	0	0	0	0	0
Q5: Were the groups similar at the start of the trial?	1	1	1	1	0
Q6: Aside from the experimental intervention, were the groups treated equally?	1	1	1	1	1
Q7: How large was the treatment effect?*					
Q8: How precise was the estimate of the treatment effect?	CI=95% 1	CI=95% 1	Nil CI 0	Nil CI 0	CI=95% 1
Q9: Can the results be applied in your context?	1	1	1	1	1
Q10: Were all clinically important outcomes considered?	1	1	1	1	0
Q11: Are the benefits worth the harms and costs?	1	1	1	1	1
TOTAL SCORE /10	9/10	9/10	8/10	7/10	5/10
TOTAL SCORE %	90%	90%	80%	70%	50%

Scoring: Yes = 1, No = 0, Can't tell = 0. *Question 7 did not require a score to be allocated.

The CASP analysis revealed several inherent methodological biases across the studies. All studies had focused questions, treated the intervention and comparator groups equally, and presented results that could be applied and would be considered beneficial. Except for the study by Razeghi et al, the randomisation process was undertaken well, and the groups were similar at the commencement of the trial. In no study were participants blinded to the treatment, which is not surprising due to the nature of both interventions and comparators. Studies scored "0" on the third CASP item if they did not employ an intention to treat analysis or did not provide explanations for participant dropout.¹⁴ Only studies by Sahin et al and Razeghi et al demonstrated the potential for significant attrition bias.^{20,21} Finally, all studies provided some statistical measure of the significance of results. However, the precision of results from Sahin et al was deemed inadequate due to the absence of reporting of 95% confidence intervals.²⁰

Team members acknowledged the potential for other biases not formally assessed by the CASP to affect the validity of results. All studies were potentially subject to the Hawthorne effect since both the intervention and comparator involved supervision by an educated therapist. The potential for attention bias must also be acknowledged since in each study, the intervention group received a longer supervised session than the control group. This was due to the additional exercises involved in the intervention. In addition, no explicit mention of the training provided to therapists was provided in any study, potentially contributing to proficiency bias. It was difficult to determine the impact of contamination bias due to a lack of reported restrictions preventing exposure of control groups to the intervention. The study by Ismail et al. did not demonstrate sufficient efforts to eliminate the presence of co-intervention bias.¹⁹

Study Characteristics

Publication dates of the included studies ranged from 2008 to 2016. They comprised of five pseudo-randomised controlled trials. The studies were conducted in Brazil, Egypt, Turkey, and Iran.¹⁷⁻²¹ The characteristics of each study are outlined in Table 3. Specific exercises within each group are detailed in Appendix 2.

Participant Characteristics

The number of participants in the studies ranged from 32 to 70 and the mean age across the studies ranged between 23 to 34 years.¹⁷⁻²¹ Three of the five studies evaluated sedentary females;^{17,18,20} one additional study by Razeghi et al. evaluated female university students, but it is unclear as to whether they met the sedentary classification.²¹ Ismail et al. was the only study to include both male and female participants.¹⁹

Table 3: Study Characteristics

Study	Population (n) Gender: F/M Age: mean±SD Status	Intervention (I)	Comparator (C)	Outcome: Measurements (OM)	Findings
Fukuda et al. ¹⁷	n = 70 70 F 25±7 (18-32) Sedentary	n = 23 Hip: OKC AB & ER + Knee Protocol 3x/wk, 4wks	1. n = 22 Knee Protocol: squats, Kn E, stretch 3x/wk, 4wks 2. n = 25 Control: ADLs	P: NRPS up/down stairs F: LEFS, AKPS, SLHT	I & C: P: ↓ (p<0.05, p>0.05 resp.) F: ↑ all OM (p<0.05) Control: Nil change (p>0.05) I vs C vs Control: P down stairs: ↓ (I cf C) (p<0.05) & Control (p<0.01); ↓ (C cf Control) (p>0.05) F: ↑ (I & C > Control): all OM (p<0.05) F: ↑ (I cf C): all OM (p>0.05)
Fukuda et al. ¹⁸	n = 54 54F 23±3 (20-40) Sedentary	n = 28 Hip: OKC AB, ER & CKC HE + Knee Protocol 3x/wk, 4wks	n = 26 Knee Protocol: squats, Kn E, calf raise, PKF, stretch 3x/wk, 4wks	P: NRPS up/down stairs F: LEFS, AKPS, SLHT	I & C: I: P ↓ & F ↑: all OM 12, 24, 52wks (p<0.05) C: P ↓: upstairs 24wks; downstairs 12, 24wks (p<0.05) F ↑: SLHT 12, 24, 52wks (p<0.05), others (p>0.05) I vs C: P: ↓ & F: ↑ (I > C): all OM 12, 24, 52wks (p<0.05)
Ismail et al. ¹⁹	n = 32 23F/9M 24±6 (18-30)	n = 16 Hip: OKC AB & ER + Knee Protocol 3x/wk, 6wks	n = 16 Knee Protocol: squats, Kn E, stretch 3x/wk, 6wks	P: VAS F: AKPS S: Dyn. (AB, ER)	I & C: P: ↓ (p=0.01, p=0.001 resp.) F & S: ↑ both (p<0.05, p<0.05 resp.) I vs C: P: ↓ & F: ↑ (I > C): (p=0.03) & (p=0.04) resp. S: ↑ (I cf C): (p=0.25-0.43)
Sahin et al. ²⁰	n = 55 55F 34.1±6.2 (28-40) Sedentary	n = 25 Hip: OKC AB & ER + Knee Protocol 12wks: 5x/wk 6wks supervised + 6wks home	n = 25 Knee Protocol: squats, Kn E, SLR, stretch 12wks: 6wks supervised + 6wks home	P: VAS F: (s) AKPS (o) THT, SLST & SDT S: Dyn. (KE, HF, AB, ER)	I & C: P: ↓ 6, 12wks (both p<0.001) F: ↑ all 6, 12wks (p<0.03 except THT) S: ↑ all 6, 12wks (p values not provided*) I vs C: P ↓ (I > C): 6, 12wks (both p=0.017) F: ↑ (I > C) AKPS: 6, 12wks (both p=0.017); SLST: 6wks (p<0.017); SDT: 6, 12wks (p<0.017) S: ↑ (I > C): AB + ER 6wks, ER 12wks (p≤0.027), others p>0.05
Razeghi et al. ²¹	n = 33 33F 22.6±2.7 (18-30) Students	n = 16 Hip: resistive hip ('all') + Knee Protocol	n = 16 Knee Protocol:	P: VAS S: digital myometer (HF, HE,	I & C: P: ↓ (p=0.001, p=0.005 resp.) S: not reported I vs C:

		4wks	squats, Kn E, (no stretch)	AB, AD, ER, IR, KE)	P:↓ (I > C): (p=0.032) S: not reported.
		4wks			
<p>P = pain; F = function; S = strength; AB = hip abductors; AD = hip adductors; ER = hip external rotators; IR = hip internal rotators; HE = hip extensors; HF = hip flexors; KE = knee extensors; Kn E = knee extension; PKF = prone knee flexion; NPRS = numerical pain rating scale; LEFS = lower extremity functional scale; AKPS = anterior knee pain scale; SLHT = single-leg hop test; SLST = single-limb squat test; THT = triple-hop test; SDT = step-down test; SLR = straight leg raise; OM = outcome measures; CKC = closed kinetic chain; OKC = open kinetic chain; PT = physical therapy; PFJ = patellofemoral joint; EMG = electromyography; Dyn. = dynamometer; s = subjective; o = objective; resp. = respectively; wks = weeks; *unable to interpret superscript a, b, c as Table 6 legend missing</p>					

Types of Intervention

Across all studies the treatment protocol for the comparator consisted of quadriceps strengthening and stretching of surrounding knee musculature, whilst the intervention protocol included the addition of strengthening exercises targeting the hip muscles. Four studies clearly stated the intervention to include hip abductors and external rotators;¹⁷⁻²⁰ with hip extensors included in one of these studies.¹⁸ The intervention periods across the studies for both intervention and comparator groups consisted of 4, 6 or, 12 week treatment programs, with training frequency ranging from three to five sessions per week. Sahin et al provided participants with supervised treatment sessions, however, no consistent significant differences in any of the outcomes were found between that study and the four remaining studies, which did not provide or report supervised treatment.^{20, 16,17,19,21}

Outcomes

Pain

Pain was rated in all studies, three using the VAS and two using the numerical pain rating scale (NPRS). All five studies demonstrated improvements, with decreased pain scores in those undertaking both the knee exercises only (control) and the combined hip and knee exercises (intervention), at all time points assessed (4, 6, & 12 weeks and 6 & 12 months).¹⁷⁻²¹ This indicates that knee-based treatments alone have positive effects on pain reduction. However, between group analyses showed that participants in the intervention groups had significantly more pain reduction post intervention and/or during all follow-ups, compared to the knee only exercise groups. Two studies that included a follow-up assessment found that pain continued to decrease during this time, suggesting the potential long-term benefits of strengthening both hip and knee muscles for pain reduction.^{18,20}

Function

Function was evaluated by four studies, all using the anterior knee pain scale (AKPS), which is also referred to as the Kujala questionnaire. This questionnaire considers objective knee findings and functional capabilities in conjunction with pain.¹⁷⁻²⁰ All intervention groups employing both hip and knee exercises displayed significant improvement in function, whereas findings from the control groups were more inconsistent. Between group analyses agreed that those in the intervention group performing both hip and knee exercises had greater function at all follow-up (4, 6, & 12 weeks and 6 & 12 months) than the comparator. Other outcome measures considered across studies included the LEFS, step down test, triple hop test, and single leg squat test. All studies found that both groups, irrespective of undertaking knee exercises alone or with hip exercises, had improved function at the end of their designated program.¹⁷⁻²⁰

Strength

Muscle strength (varying combinations of hip abductors, hip external and internal rotators, hip flexors and extensors, and knee extensors) was assessed by three studies.¹⁹⁻²¹ Two of these studies used a dynamometer.^{19,20} One study used a myometer.²¹ One study reported increased strength in both intervention and comparator groups in the assessed muscles at each time point measured.¹⁹ However, interpretation of results of the remaining two studies assessing strength was unclear.^{20,21} Between group findings were inconsistent between the studies. Sahin et al. found that at 6 and 12 weeks, those undertaking both hip and knee exercises had greater strength (abductors and external rotators at six weeks, external rotators only at 12 weeks) than those performing knee exercises alone.²⁰ Contrastingly, Ismail et al. found no significant difference in strength at six weeks between those performing both hip and knee exercises and those who performed knee exercises only.¹⁹ These results may be influenced by differences in treatment frequency and duration between the two studies. Sahin et al. held five sessions per week over six weeks (a total of 30 sessions) with an additional six-week home exercise program, whilst Ismail et al. only conducted three sessions per week over six weeks (a total of 18 sessions) without a home exercise program.^{19,20}

Table 4: Summary of Results

Study	Outcomes		
	Pain	Function	Strength
Fukuda et al. (2010) ¹⁷	↓*	↑*	N/A
Fukuda et al. (2012) ¹⁸	↓*	↑*	N/A
Ismail, Gamaleldein and Hassa (2013) ¹⁹	↓*	↑*	↔
Sahin et al. (2016) ²⁰	↓	↑	↑
Razeghi et al. (2010) ²¹	↓*	N/A	N/R

Key:

N/A = Not applicable (for this outcome measure in this study)

↔ = No difference (between intervention and comparator/control)

N/R = not reported

↓ = Reduction with intervention (cf comparator/control)

↑ = Increase with intervention (cf comparator/control)

* = Results are statistically significant (P<0.05)

NHMRC Body of Evidence Framework

Table 5 synthesises the results of the included studies using the NHMRC FORM framework.¹⁵ The included studies, despite being classified as high-level evidence forms, could only be considered as moderate quality due to methodological concerns involving lack of blinding and true randomisation. Furthermore, the evidence base is somewhat mixed due to the focus being on different outcomes, using a range of outcome measures, as well as differing intervention programs. Therefore, despite results being largely positive for the efficacy of the addition of hip strengthening exercises to regular physiotherapy treatment, care should be taken when considering clinical application.

Table 5: NHMRC Body of Evidence Framework

Component	Grade	Comments
Evidence base	C Satisfactory One or two level III studies with a low risk of bias	Quantity: total of five studies Level III: six studies Quality: Moderate
Consistency	B Good Most studies consistent and inconsistency may be explained	Consistent study design Some variety in gender/s of population studied Some variety in intensity, frequency and duration of intervention Some variety in outcome measures
Clinical Impact	B Substantial	Duration of therapy required to achieve effects across all studies is clinically feasible All intervention protocols were well described All outcome measures reported on were of substantial clinical significance Findings for pain were consistent across all studies No adverse effects were reported No long-term regression was reported by the two studies that included follow up
Generalisability	C Satisfactory Population/s studied in body of evidence differ to target population for guideline but it is clinically sensible to apply this evidence to target population	Population is consistent with the target population in terms of age however not gender Three of the five studies specifically investigated sedentary females Three studies conducted power calculations where power was achieved Studies conducted in four different countries, all with different cultural and religious beliefs

Applicability	B Good Applicable to Australian healthcare context with few caveats	No additional or specialised treatment or equipment required for implementation No studies conducted in Australia, however it can be assumed that, based on mode and nature of treatment, these treatment protocols would be well adopted in Australian healthcare Attitudes, beliefs (religious or otherwise), may be contrasting and have an effect on compliance and results
Grade of Recommendation	B Body of evidence can be trusted to guide practice in most situations	The studies were of moderate quality Homogenous findings across studies for at least one of the three outcome measures The current evidence is mixed with some discrepancies in outcome measures, baseline characteristics and intervention programs Results of the review are most relevant to a sedentary female population

DISCUSSION

The five studies included in the systematic review investigated the effectiveness of the addition of hip strengthening exercises to standard physiotherapy knee treatment on pain, strength, and function when compared to standard physiotherapy care alone in adults with PFPS.¹⁷⁻²¹ All five studies demonstrated significant improvements in pain scores in intervention compared to comparator groups, and the four studies that assessed function also reported significant comparative functional improvements. Of the three studies that measured strength, only Sahin et al. reported a significant increase in the hip strength for the abductors and external rotators in the intervention compared to the comparator group.²⁰ No adverse effects of treatment were reported in any of the five studies.¹⁷⁻²¹

The results of this review are most applicable to young sedentary females. Although Ismail et al.¹⁹ included male participants in their study, across all studies the overall sample sizes were small, the power calculations limited, and males were underrepresented, making up only 9 out of a total of 244 (3.7%) participants. The four other studies only included female participants classified as either 'sedentary' or 'not involved in professional sports activity'.^{17,18,20,21} Sahin et al. had a slightly older group of participants with mean age 34.1 years compared to all other studies with mean ages ranging from 22.6 to 25 years.²⁰ Thus, the transferability of the results of the review is most appropriate for the young sedentary female population.

Exercise type, Prescription & Method of Delivery

The results of this review indicate that additional open or closed-kinetic chain strengthening exercises targeting hip abductors, hip external rotators, and hip extensors can be effective in improving pain and function. Although there was variation in the frequency of interventions across studies and session durations were largely unreported, it may be useful for clinicians to know that there is moderate level evidence to suggest that significant and long-lasting improvements in both outcomes can be achieved with as little as 12 therapist-supervised sessions over four weeks. One of the great benefits for clinicians incorporating this management strategy is that it requires no additional training or equipment and can be undertaken at minimal cost. However, patients considering this form of management need to be motivated and willing to invest the time and costs associated with approximately 12, 30-minute visits over four weeks. At this stage, there is insufficient evidence to determine the effectiveness of a group-based exercise class or a similar home-based exercise program. Based on a study by Sahin et al. which included five sessions per week for six weeks with an additional six-week home exercise program, there is a suggestion that gains in strength may also be achieved with a more frequent and intense program.²⁰

Comparison with Previous Studies

The recent systematic review by Santos et al. investigating a similar research question reported comparable findings.⁵ In addition to updating the search from September 2014 to September 2017, the current review searched a broader range of databases, yielding one additional study by Sahin et al.²⁰ Furthermore, a more rigorous eligibility criteria was applied which resulted in the exclusion of three studies included in the Santos et al. review.⁹⁻¹¹ In Dolak et al., the patients with PFPS undertook either a hip strengthening program or a knee strengthening program; then, both groups performed similar functional weight bearing exercises.⁹ In Khayambashi et al. study, the intervention group undertook hip abductor and external rotator strengthening and the control group was placed on an Omega 3 diet with no exercise.¹⁰ In neither study was it considered that the effects of hip strengthening compared

to standard PFPS management could be discerned.^{9,10} In Nakagawa et al. study, the intervention group participated in a program consisting of both transverse abdominus exercises and hip strengthening exercises, which does not separate the isolated effect of hip strengthening exercises from standard physiotherapy care for PFPS.¹¹ It is possible that the inclusion of these three studies may have inflated the results of the review by Santos et al.⁵ While there was agreement across the findings between the current review and that by Santos et al., the more focused research question and stricter exclusion criteria of the current review provides greater confidence in the results and application of these to clinical practice.⁵

Limitations

The systematic review has several limitations related to review processes, study outcomes, and methodology. A comprehensive search strategy including searching additional studies from related systematic reviews gleaned five studies that met the inclusion criteria. Another study by Soleimani et al. with an English abstract appeared to be current and relevant; however full text was accessible in Persian only, and was therefore excluded from the review.¹⁶

Common methodological concerns included lack of blinding to treatment, which is difficult in an ongoing physiotherapy program. Included studies were also mostly low-level research designs (NHMRC Level III-1 Intervention). They were unable to be classified as Level II randomised controlled trials due to the use of an envelope-based pseudo-randomisation process as opposed to a computer generated randomisation.¹⁵ Furthermore, one study was limited in participant continuity and between-group participant homogeneity, as well as in considering all clinically important outcomes.²¹ For the three main outcomes, pain, function, and strength, different outcome measures were used, resulting in an inability to undertake a meta-analysis, thus preventing combining the data from a number of studies to identify the overall effect. Power calculations were utilised and achieved in three of the five included studies.^{17,18,21} In a fourth study, the sample size was based on a previous study.¹⁹ All studies utilised minimal clinically important difference (MCID) to detect clinical significance, however not for all outcome measures. Although two studies used MDIC to determine improvement in both pain and function (and strength was not assessed)^{17,18}, the remaining three studies only utilised it to determine clinical significance of pain, not function or strength.¹⁹⁻²¹ It was not specified in any of the three studies assessing dynamometer or myometer strength whether the findings were normalized to measures of body size (e.g. body mass or height, to remove body-size dependence).^{19,20,21} Although findings were persuasive, some caution is therefore required in drawing indisputable clinical conclusions.

With respect to participant status, the majority of studies focused on females alone, with three recruiting specifically sedentary females. While Ismail et al. considered both genders, the majority of participants were female.¹⁹ As a result, evidence within this systematic review is limited regarding efficacy of application to a male population. Finally, research team members recognise the possibility of reporting bias, as all included studies had positive findings for at least one outcome.

CONCLUSIONS

Implications for Clinical Practice

Overall, it can be concluded that within the limitation of poorer quality of some of the included studies, the addition of hip strengthening exercises to standard physiotherapy treatment results in greater improvements in both pain and function for adults with PFPS, compared with standard physiotherapy care alone. There is less confidence in the findings for changes in strength. More specifically, significant improvements in both pain and function can be gained through a program including knee and hip exercises undertaken for as little as three times per week over four weeks, with potential for additional benefit of improved strength to occur with a more regular program of five times per week over six weeks. It should be noted, however, that the majority of studies in this review focused on sedentary females, so results are most relevant to this population.

Implications for Future Research

With reference to individual studies, recommendations for future research include allocation of both male and female participants with recruitment of a sufficiently large sample to allow for subgroup analysis of gender differences. The use of a true control in addition to intervention and comparator groups is also recommended. Further considerations include addressing all clinically important outcomes (pain, strength and function), utilising minimal clinically important difference (MCID) for all outcome measures used to consider clinical significance, providing normalised strength values for better interpretation of the findings, refining the randomisation process, and consistent use of power calculations. It may also be beneficial to further explore the effectiveness of a home exercise program following supervised and group classes for the first four weeks, which may offer future direction for clinicians. In terms of methodology, accepting only a particular measure for inclusion for each main outcome would increase consistency of results and improve the ability to form clearer comparisons between studies.

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Appendix 1: Search Strategy

Primary Databases						
E-Databases	Search Terms					
	P	I	C	O	Limits	Specification
MEDLINE Embase	Patellofemoral Pain Syndrome/ ("PFPS" OR "PFJS" OR "patellofemoral pain syndrome" OR "patellofemoral joint syndrome" OR "runner* knee")	Exercise Therapy/ Stretching/ Strengthening/ ("hip strength*" exercise*" OR "hip exercis*" (prev. exercises*) OR "hip strengthen*" (prev. strengthening") OR ("hip abduct*" OR "hip extens*" OR "hip external rotat*" OR "hip lateral rotat*" OR "gluteus" OR "TFL" OR "tensor fasciae latae" OR "tensor fascia lata") adj3 (strength* OR exercis*))	<i>None used in search strategy</i>	Treatment Outcome/ (pain OR function* OR "hip strength**")	English Language Humans Adult (19-44 years)	MeSH headings (Patellofemoral Pain Syndrome) and Keywords Truncations Phrase searching (in intervention, used "adj3" = within 3 words)
SportDiscus CINAHL	"PFPS" OR "PFJS" OR "patellofemoral pain syndrome" OR "patellofemoral joint syndrome" OR "runner* knee"	"hip strength*" exercise*" OR "hip exercis*" (prev. exercises*) OR "hip strengthen*" (prev. strengthening") OR ("hip abduct*" OR "hip extens*" OR "hip external rotat*" OR "hip lateral rotat*" OR "gluteus" OR "TFL" OR "tensor fasciae latae" OR "tensor fascia lata") N3 (strength* OR exercis*)		pain or function* or "hip strength**"	English language	Subject headings (PLICA syndrome) Truncations Phrase searching (in intervention, used "N3" = within 3 words of)
Secondary Databases						
Cochrane	PFPS OR PFJS OR patellofemoral pain syndrome OR patellofemoral joint syndrome	hip strength* exercise* OR hip exercise* OR hip strengthening	<i>None used in search strategy</i>	pain or function* OR hip strength*	Nil	Subject headings and Keywords Truncation
PEDro	Patellofemoral pain syndrome OR patellofemoral joint syndrome	Strength training Lower leg or knee Musculoskeletal		Strength	Clinical trial	Subject headings and Keyword

Appendix 2: Exercise Prescription

Study	Intervention (I)	Comparator (C)
Fukuda et al. 2010 ¹⁶	[12 sessions, 3x/wk, 4wks] HIP Abd: stand against elastic resistance 10RM 3x10 reps; sidelying with weight 70% 1RM 3x10 reps; side step against elastic resistance 3x1min ER: sit against elastic resistance 10RM 3x10 reps. (10RM adjusted weekly) + KNEE (see comparator)	[12 sessions, 3x/wk, 4wks] KNEE Stretching 3x30s (ITB, HS, Q, G) Iliopsoas NWB 70% 1RM 3x10 reps Sitting knee E (90-45°) 70% 1RM 3x10 reps Leg press (0-45°) 70% 1RM 3x10 reps Mini-squat (0-45°) 70% 1RM 3x10 reps
Fukuda et al. 2012 ¹⁷	[12 sessions, 3x/wk, 4wks] HIP Abd: stand against elastic resistance 10RM 3x10 reps; sidelying with weight 70% 1RM 3x10 reps ER: sit against elastic resistance 10RM 3x10 reps E: machine, 70% 1RM 3x10 reps + KNEE (see comparator)	[12 sessions, 3x/wk, 4wks] KNEE Stretching 3x30s (ITB, HS, Q, G) Iliopsoas NWB 70% 1RM 3x10 reps Sitting knee E (90-45°) 70% 1RM 3x10 reps Leg press (0-45°) 70% 1RM 3x10 reps Mini-squat (0-45°) 70% 1RM 3x10 reps Single-leg calf raise 70% 1RM 3x10 reps PKF 70% 1RM 3x10 reps
Ismail, Gamaleldein and Hassa 2013 ¹⁸	[18 session, 3x/wk, 6wks] HIP Abd: sidelying with ankle cuff weight, 60% 10RM 2x10 reps, 6s hold ER: sit with ankle cuff weight 60% 10RM, 2x10 reps, 6s hold (10 RM adjusted weekly) + KNEE (see comparator)	[18 session, 3x/wk, 6wks] KNEE Mini-squats 0-40° with ball squeeze 6s hold f/w and lateral step up (8inch height) 6s hold Standing knee E 30° 6s hold against elastic resistance Stretching 3x30s (ITB, HS, Q, G)
Sahin et al. 2016 ¹⁹	[30 sessions, 5x/wk, 6wks, 30 min] All 100% of 10RM, 1x10 reps HIP Abd: stand against elastic resistance (0 to 30-35°) 3.5s hold, 5 reps x2 daily (5reps added weekly) ER: sitting elastic resistance (0 to 30°) 3.5s hold, 5 reps x2 daily (5reps added weekly) + KNEE (see comparator) + 6wks at home exercise program	[30 sessions, 5x/wk, 6wks, 30 min] KNEE Stretching 3x10s 2x daily (ITB, HS, Q, G) Supine isometric Q with towel 10s hold 20 reps x2 daily (5reps added weekly) SLR 3.5s hold 10reps x 2 daily Mini-squats 30-45° 10s hold, 10 reps x 2 daily Sitting resisted knee E against resistance band 3.5s hold, 5 reps x 2 daily (5reps added weekly) + 6wks at home exercise program
Razeghi et al. 2010 ²⁰	[4wks (no further detail)] HIP: progressive resisted hip exercises for 'all' hip muscles (no further detail on exercise or prescription) + KNEE: (see comparator)	[4wks (no further detail)] KNEE: mini-squats, resisted knee E terminal and 90° to 50° (no further detail on prescription)

RM = rep maximum; s = seconds; wk = week; HS = hamstrings; Q = quadriceps; ITB = iliotibial band; G = gastrocnemius; NWB = non-weight-bearing; E = extension, PKF = prone knee flexion; f/w = forward; SLR=straight leg raise