Individuals With Patellofemoral Pain Have Impaired Self-Reported and Performance-Based Function: Systematic Review With Meta-Analysis and Meta-Regression

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Objectives: To determine impairments in self-reported and performance-based function in individuals with patellofemoral pain (PFP) and determine physical and nonphysical factors potentially related to these impairments.

Data Sources: We searched MEDLINE, Embase, CINAHL, Web of Science, and SPORTDiscus databases between inception and January 2024.

Study Selection: Included studies compared self-reported and performance-based measures of function between PFP-affected and pain-free limbs in individuals with unilateral PFP or between individuals with PFP and pain-free individuals.

Data Extraction: The key information from each study was extracted by 1 independent researcher and reviewed by another researcher.

Data Synthesis: We performed meta-analyses for each self-reported and performance-based measure of function and meta-regressions to identify factors that might explain outcomes of meta-analyses. We assessed the certainty of evidence using the Grading of Recommendations Assessment, Development, and Evaluation. We included 83 studies (2807 individuals with PFP and 2518 pain-free individuals). We identified very-low— to high-certainty evidence that individuals with PFP have reduced self-reported (large effect sizes; standardized

mean difference [SMD] = -1.99; 95% CI = -2.41, -1.57 to SMD = -4.87; 95% CI = -6.97, -2.77) and performance-based (small to large effect sizes; SMD = -0.30; 95% CI = -0.58, -0.02 to SMD = -0.80; 95% CI = -1.11, -0.50) measures of function compared with pain-free individuals, but no differences were found between limbs in individuals with unilateral PFP for most performance-based measures of function (small to moderate effect sizes; SMD = -0.20; 95% CI = -0.68, 0.27 to SMD = -0.49; 95% CI = -1.02, 0.03). Age, body mass index, duration of symptoms, and self-reported pain did not explain self-reported function, and age did not explain performance-based function (R^2 range, <0.01-0.02; P range, .15-.91).

Conclusions: Our results highlight the negative effect of PFP on self-reported and performance-based function, which seems to also affect the pain-free limb. Self-reported and performance-based measures of function should be considered when assessing individuals with PFP. None of the factors investigated explained impaired self-reported and performance-based function.

Key Words: clinical tests, functional capacity, patient-reported outcome measures, physical function, subjective function

Key Points

- Individuals with patellofemoral pain (PFP) had impaired function compared with pain-free individuals; thus, function measures should be considered primary outcomes in the management of PFP.
- No function differences were observed between limbs in individuals with unilateral PFP; therefore, caution is warranted when comparing function between PFP and pain-free limbs.
- Age, body mass index, duration of symptoms, and self-reported pain did not explain function.

Individuals with patellofemoral pain (PFP) frequently present to orthopaedic and sports clinics given the high prevalence of PFP in active adolescents and young adults. These individuals report diffuse anterior knee pain during daily living or sporting activities such as stair ascent and descent, squatting, and hopping. Reductions in health-

related quality of life,^{3–5} psychological well-being,^{6,7} and physical activity and sport participation,⁸ as well as impairments in self-reported and performance-based function,^{9,10} have been reported in individuals with PFP. Self-reported measures (eg, patient-reported outcome measures) indicate how individuals with PFP perceive their functional limitations,

whereas performance-based measures of function (eg, single-leg hop test [SLHT]) represent the actual objectively measured functional limitation. Both provide clinically relevant and complementary information that can help guide the development of effective interventions.

Measures of function have been considered one of the key determinants of PFP and its prognosis. 11,12 Self-reported function has been related to pain severity, kinesiophobia, and psychological well-being, 7,13 and poor self-reported function has predicted unfavorable recovery 5 to 8 years after treatment.11 Performance-based measures of function, such as hopping and stepping tasks, have been related to hip and knee strength, 9,14 which are key targets of PFP management.¹² A greater understanding of the potential magnitude of functional impairments may help inform preferable outcome measures for decision-making processes. Despite the importance of measures of function for PFP, no researchers have systematically synthesized the literature to compare selfreported function between individuals with PFP and pain-free individuals and performance-based function between individuals with PFP and pain-free individuals or the PFP-affected and pain-free limbs of individuals with unilateral PFP. In addition, no systematic review has been conducted to identify factors that may explain poor self-reported and performancebased function in individuals with PFP. Determining which measures of function are impaired as well as which physical and nonphysical factors may underline these deficits is important given that function improvement is a common target of PFP rehabilitation.¹⁵

The purposes of our systematic review were to (1) systematically review and meta-analyze the literature comparing self-reported and performance-based function between individuals with PFP and pain-free individuals or the PFP-affected and pain-free limbs of individuals with unilateral PFP and (2) investigate physical and nonphysical factors that might explain poor self-reported and performance-based function in individuals with PFP via meta-regression.

METHODS

We conducted this systematic review in accordance with Preferred Reporting Items for Systematic Reviews and Meta-Analyses guidelines¹⁶ and registered it in the International Prospective Register of Systematic Reviews database (CRD42021234911).¹⁷ Protocol deviations are summarized in Supplemental Material 1.

Search Strategy

We conducted the initial electronic search in MEDLINE, Embase, CINAHL, Web of Science, and SPORTDiscus between inception and February 2021 and updated it in January 2024. We combined the keywords and medical subject headings related to PFP and self-reported and performance-based measures of function with search filters to develop the search strategy. We created the primary search for MEDLINE and adapted it to the other databases through pilot searches (Supplemental Table 1). We did not search the gray literature.

Selection Criteria

One author (A.F.B.B.) imported identified studies into Covidence (Veritas Health Innovation), and duplicates

were removed.¹⁸ Two authors (A.F.B.B. and J.C.P.S.) independently screened titles and abstracts for eligibility using the criteria presented in Table 1. We retrieved full-text articles of potentially relevant abstracts for further review. When the full text was not available, we requested it from the corresponding authors via email. If authors were unable to provide the full text, we excluded the study. When studies from the same author groups presented similar descriptive values of measures of function, we included only the first study published after confirmation with the corresponding author that both publications included the same cohort. Disagreements were resolved by consulting a senior author (R.V.B.).

Data Extraction

One author (R.V.B.) extracted study and participant characteristics (eg, lead author, year of publication, sample size, sex, and participant age), self-reported and performancebased measures of function (eg, Anterior Knee Pain Scale [AKPS], Lower Extremity Functional Scale [LEFS], Knee injury and Osteoarthritis Outcome Score [KOOS], KOOS for PFP and osteoarthritis [KOOS-PF], forward step-down test [FSDT], and hop or balance tests), and predictors of interest to be included in the meta-regression. We selected physical (eg, body mass index [BMI] and strength) and nonphysical (eg, kinesiophobia and pain catastrophizing) predictors of interest based on recommended items from Reporting of Quantitative Patellofemoral Pain (REPORT-PFP), the biomechanical and psychological consensus of PFP. 6,19,20 A second author (J.C.P.S.) reviewed all extracted data. We extracted means, SDs, and sample sizes for all outcomes and used them for data analysis. When data were missing, we contacted corresponding authors for further information via email up to 3 times. If authors were unable to provide the data or did not respond to the requests and missing data could not be calculated using Review Manager 5.4 (The Cochrane Collaboration), we did not enter the study in the meta-analyses. For these studies, we only performed an individual study analysis by calculating the standardized mean differences (SMDs) and discussing them. We provide details on data-extraction management in Supplemental Material 2.

Methodologic Quality Assessment and Risk of Bias

We assessed internal and external validity of observational and nonrandomized interventional studies with a domain-based evaluation using the modified Downs and Black checklist, as performed by Hart et al.²¹ We assessed internal validity across the following 5 domains: performance bias (items 14, 15, and 19), reporting bias (item 16), detection bias (items 17, 18, and 20), selection bias (items 21-25), and attrition bias (item 26). We assessed external validity using items 11 through 13. Items were scored as yes, no, or unable to determine. Overall quality classification for each study was based on concerns across all applicable items and domains rather than the numeric summary score. Studies were classified across domains and external validity as low, moderate, or high quality based on item evaluation. A similar classification was performed for internal validity based on domain evaluation. We assessed the methodologic quality of the only randomized clinical trial²²

	Criteria
Inclusion	
Design	Observational prospective or cross-sectional/case-control studies, pretest-posttest studies, and randomized or nonrandomized clinical trials written in English, Portuguese, or Spanish.
Participants	Individuals with insidious unilateral or bilateral PFP of both sexes, age < 50 y, without any other previous or concomitant knee or lower limb condition reported.
Comparisons	Pain-free control group or pain-free contralateral limb of individuals with unilateral PFP.
Outcomes	Self-reported function as measured using questionnaires or scales.
	Physical performance during clinical tests.
Exclusion	Retrospective comparative cohort studies, review papers, theses, editorials, abstracts, and letters. Studies without a comparator (pain-free group or limb).

Abbreviation: PFP, patellofemoral pain.

included using the Physiotherapy Evidence Database (PEDro) scale²³ and its risk of bias using the Cochrane Risk of Bias for Randomized Trials 2 (RoB2) following Cochrane Handbook for Systematic Reviews of Interventions recommendations.²⁴ The 10-item PEDro scale consists of a score ranging from 0 to 10 and is used to rate trials according to the presence or absence of some methodologic quality criteria.²³ The score classifications are high quality (>7 of 10), moderate quality (4–6 of 10), and low quality (<3 of 10). The RoB2 comprises the following 5 domains: randomization process, deviations from intended interventions, missing outcome data, measurement of the outcome, and selection of the reported study. For each domain, the tool comprises a series of signaling questions scored as yes, probably yes, probably no, no, and no information. We classified each domain as low, high, or some concerns of risk of bias based on the tool's algorithm.²⁵ We determined the overall risk of bias using the worst-scorecounts method, which takes the lowest rating across all the domains.26 Further details regarding study-quality assessment and risk of bias are provided in Supplemental Material 3 and Supplemental Tables 2 and 3. Two authors (A.F.B.B. and M.C.W.) independently assessed all studies, and disagreements were resolved by consulting a senior author (R.V.B.).

Data Synthesis and Analysis

One author (R.V.B.) performed meta-analyses using Review Manager 5.4 and random-effects models when >2 studies investigated the same outcome and comparator (pain-free individuals or pain-free limb of individuals with unilateral PFP). Another author (A.F.B.B.) reviewed all meta-analyses. We calculated SMDs with 95% CIs (Hedges g) once different scales or units of measurement were reported across studies, even in those using the same questionnaire or test (eg, Goharpey et al,27 Peeler and Anderson, ²⁸ and Guimaraes Araujo et al²⁹). We classified SMDs as small (≥ 0.2), moderate (0.5–0.79), and large (≥ 0.80) effects.³⁰ We quantified statistical heterogeneity for pooled results using the I^2 statistic and defined it as not important (<50%), moderate (50%-75%), or high (>75%).³¹ We estimated publication bias using the Egger regression test. For data that were not included in the meta-analyses, we calculated the SMDs with 95% CIs for individual comparisons and discussed them. Confidence intervals excluding zero were considered statistically significant.

We performed meta-regressions to identify predictors that could explain the SMDs (Hedges g) of function outcomes. Random-effects meta-regressions were performed using Comprehensive Meta-Analysis software (BioSTAT Consultants, Inc) when at least 10 studies included in a meta-analysis presented data for the same predictor.²⁵

Certainty of Evidence

Two authors (A.F.B.B. and M.C.W.) assessed the certainty of evidence using a modified Grading of Recommendations Assessment, Development, and Evaluation approach.32,33 Given the observational nature of the research question of our systematic review, certainty of evidence started as high and was downgraded or upgraded according to Grading of Recommendations Assessment, Development, and Evaluation Handbook recommendations, which are described in Table 2.32,34 We defined levels of certainty of evidence as follows: high when further research is very unlikely to change confidence in the estimate of the effect; *moderate* when further research is likely to have an important effect on confidence in the estimate of the effect and may change the estimate; low when further research is very likely to have an important effect on confidence in the estimate of the effect and is likely to change the estimate; and very low when there is very little confidence in the effect estimate.32

RESULTS

Our systematic search identified 28 797 titles and abstracts for screening (Figure 1). After duplicates were removed, 21 648 studies underwent title and abstract screening, then 475 studies underwent full-text screening. We included 83 studies in the review: 73 observational studies, 8-10,27,29,35-102 8 pretest-posttest studies, 28,103-109 1 randomized clinical trial, 22 and 1 crossover study. 110 We grouped studies with cross-sectional and case-control designs because, despite being similar, they did not consistently report the design and most exhibited characteristics of both designs.

Study and Participant Characteristics

Study and participant characteristics are summarized in Supplemental Material 4. A total of 2807 individuals with

Table 2. Outcome Level of Certainty of Meta-Analyses (GRADE Approach)

					Downgrad	ling Domain ^a		Upgrading Domain	
	No. of Individuals			Risk of			Publication	Large	Level of
Outcome	(Studies)	SMD (95% CI)	I^{2} , %	Biasb	Inconsistency ^c	Imprecision ^d	Biase	Effect ^f	Certainty
Self-reported function	n: PFP \times pain-free	groups							
AKPS	2414 (40)	-3.45(-3.84, -3.06)	88	-1	-1	0	0	+1	Moderate
LEFS	593 (9)	-3.83(-5.10, -2.55)	95	-1	-1	0	NA	+1	Moderate
FIQ	337 (7)	-4.87 (-6.97, -2.77)	96	-1	-1	0	NA	+1	Moderate
KOOS	255 (5)	-1.99(-2.41, -1.57)	43	-1	0	0	NA	+1	High
ADLS	375 (5)	-2.79(-3.49, -2.08)	83	-1	-1	0	NA	+1	Moderate
KOOS-PF	124 (4)	-2.66(-3.47, -1.86)	60	-1	-1	0	NA	+1	Moderate
Lysholm	102 (3)	-2.23(-3.51, -0.96)	82	-1	-1	0	NA	+1	Moderate
Performance-based f	function: PFP $ imes$ pai	n-free groups							
Balance tests	789 (12)	-0.66(-1.12, -0.19)	88	-1	-1	0	0	0	Low
FSDT	737 (9)	-0.80(-1.11, -0.50)	68	-1	-1	0	NA	+1	Moderate
SLHT	711 (7)	-0.42(-0.57, -0.27)	0	-1	0	0	NA	0	Moderate
SLTHT	196 (2)	-0.30 (-0.58, -0.02)	0	-1	0	0	NA	0	Moderate
Bilateral squat test	70 (2)	-1.21 (-2.71, 0.29)	86	-1	-1	0	NA	+1	Moderate
	function: painful limb	b imes contralateral pain-fre	ee limb						
Balance tests	70 (2)	-0.20 (-0.68, 0.27)	0	-1	0	-1	NA	0	Low
FSDT	106 (2)	-0.36 (-1.11, 0.38)	72	-1	-1	-1	NA	0	Very low

Abbreviations: ADLS, Activities of Daily Living Questionnaire; AKPS, Anterior Knee Pain Scale; FIQ, Functional Index Questionnaire; FSDT, forward step-down test; GRADE, Grading of Recommendations Assessment, Development, and Evaluation; KOOS, Knee Injury and Osteoarthritis Outcome Score; KOOS-PF, KOOS for PFP and osteoarthritis; LEFS, Lower Extremity Functional Scale; Lysholm, Lysholm Knee Scoring Scale; NA, not applicable; PFP, patellofemoral pain; SLHT, single-leg hop test; SLTHT, single-leg triple hop test; SMD, standardized mean difference.

PFP and 2518 pain-free individuals were included. Mean \pm SD ages for PFP and pain-free individuals were 22.91 \pm 6.55 years and 23.24 \pm 6.23 years, respectively. Mean \pm SD BMIs for PFP and pain-free individuals were 23.04 \pm 3.58 and 22.30 \pm 3.05, respectively. The most common self-reported measures of function were the AKPS (42 studies*), LEFS (9 studies³ $^{37,60,65,68-73}$), Functional Index Questionnaire (7 studies² $^{28,74-77,105,106}$), KOOS (7 studies² $^{29,84-87}$), Activities of Daily Living Questionnaire (5 studies² $^{29,84-87}$), KOOS-PF (5 studies° 9,79,88,89,110), and Lysholm Knee Scoring Scale (5 studies° 60,85,90,91,107). The most common performance-based measures of function were balance tests, including the Star Excursion Balance Test (5 studies° 60,92,93,108,109), Y-Balance Test (YBT; 8 studies° $^{22,29,42,88,89,94-96}$), FSDT (11 studies†), and SLHT (8 studies° 9,42,45,61,62,93,99,101).

Methodologic Quality and Risk of Bias

Methodologic quality and risk-of-bias assessment of the included studies is provided in Supplemental Material 3 and Supplemental Tables 2 and 3. We rated nearly 76% (63 studies^{\$)} of the studies as low quality for performance bias, 20% (17 studies^{\$)} as low quality for reporting bias, 2% (2 studies^{27,51}) as low quality for detection bias, 53% (44 studies¹) as low quality for selection bias, and 5% (4 studies^{28,62,94,105}) as low quality for attrition bias. Overall, we judged most studies to have low quality for internal (87%, 72 studies¹) and external (98%, 81 studies^{8-10,27-29,35-68,70-110}) validity. A single study was assessed using the PEDro scale and RoB2 and was classified as *moderate quality* (6 of 10) and high risk of bias, respectively.²²

Publication Bias

We could only assess risk of publication bias for the AKPS and balance tests meta-analyses. No publication bias was detected (Supplemental Figure 1).

^a As the inclusion and exclusion criteria were rigorous and only studies with populations and outcomes that exactly fit the review question were included, the indirectness domain was not applied.

^b The domain was downgraded 1 level when >25% of participants from studies were judged as having one-half or a majority of domains with high risk of bias in the assessment tool.

[°] The domain was downgraded 1 level when $I^2 > 50\%$.

^d The domain was downgraded 1 level when the difference of the effect on the patient would differ depending on use of the upper vs lower boundary of the CI.

 $^{^{\}rm e}$ The domain was downgraded 1 level when P < .05 in the Egger regression test.

^f The domain was upgraded 1 level when pooled results had large effects (>0.80).

^{*} References 8-10, 27, 29, 35-67, 103-105, 110.

[†] References 9, 27, 45, 61, 62, 93, 97–100, 105.

[‡] References 8, 10, 27, 29, 35–40, 46, 48, 49, 51–53, 55, 57–72, 74–77, 79, 80, 82, 84–88, 90–92, 94–107, 110.

[§] References 27, 42, 53, 62, 66, 85, 89, 94–98, 100, 104–107.

[&]quot;References 10, 27, 37, 38, 40, 42, 45, 46, 50–52, 57, 58, 60, 64–66, 68, 69, 71, 73–79, 82, 85, 87, 89–91, 95–98, 100, 101, 104, 105, 107–109.

¹References 8, 10, 27–29, 35–40, 42, 45, 46, 48–53, 55, 57–80, 82,84–92, 94–110.

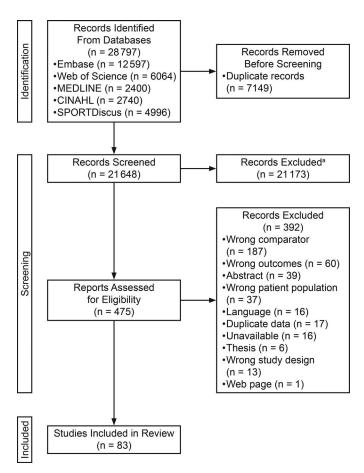


Figure 1. Flow of studies through the review.

Data Findings

We pooled 78 studies# in meta-analyses and presented their level of certainty outcomes in Table 2. We could not pool 5 studies^{82,91,100,102,107} due to missing descriptive or parametric data^{82,91,100,102,107} or lack of sufficient studies. 100,102 We pooled only part of the outcomes of 11 studies** that reported multiple function measures due to missing descriptive or parametric data,^{62,110} lack of sufficient studies,^{9,10,29,59,66,67,97,99} or duplicate data.⁶¹ We present a synthesis of unpooled studies with SMDs and 95% CIs in Supplemental Material 5, except for the betweenlimbs comparisons of performance-based measures of function that are presented in the Performance-Based Measures of Function subsection. We were unable to synthesize the AKPS and KOOS data from 2 studies^{62,110} due to missing descriptive or parametric data or both, and we did not synthesize the AKPS data of 1 study⁶¹ to avoid duplication of information, as it presented the same data as the study by Ferreira et al.⁴⁷

Self-Reported Function

Anterior Knee Pain Scale. Moderate certainty of evidence from 40 studies (2414 individuals) showed that

individuals with PFP have reduced self-reported function measured with the AKPS compared with pain-free individuals (large effect size, SMD = -3.45; 95% CI = -3.84, -3.06; $I^2 = 88\%$; P < .001; Figure 2A).

Lower Extremity Function Scale. Moderate certainty of evidence from 9 studies (593 individuals) showed that individuals with PFP have reduced self-reported function measured with the LEFS compared with pain-free individuals (large effect size, SMD = -3.83; 95% CI = -5.10, -2.55; $I^2 = 95\%$; P < .001; Figure 2B). $^{37,60,65,68-73}$

Functional Index Questionnaire. Moderate certainty of evidence from 7 studies (337 individuals) showed that individuals with PFP have reduced self-reported function measured using the Functional Index Questionnaire compared with pain-free individuals (large effect size, SMD = -4.87; 95% CI = -6.97, -2.77; $I^2 = 96\%$; P < .001; Figure 2C).^{28,74–77,105,106}

Knee injury and Osteoarthritis Outcome Score. High certainty of evidence from 5 studies (255 individuals) showed that individuals with PFP have reduced self-reported function measured using the KOOS compared with pain-free individuals (large effect size, SMD = -1.99; 95% CI = -2.41, -1.57; $I^2 = 43\%$; P < .001; Figure 2D). ^{78–81,83}

Activities of Daily Living Questionnaire. Moderate certainty of evidence from 5 studies (375 individuals) showed that individuals with PFP have reduced self-reported function measured using the Activities of Daily Living Questionnaire compared with pain-free individuals (large effect size, SMD = -2.79; 95% CI = -3.49, -2.08; $I^2 = 83\%$; P < .001; Figure 2E).

KOOS for PFP and Osteoarthritis. Moderate certainty of evidence from 4 studies (124 individuals) showed that individuals with PFP have reduced self-reported function measured using the KOOS-PF compared with pain-free individuals (large effect size, SMD = -2.66; 95% CI = -3.47, -1.86; $I^2 = 60\%$; P < .001; Figure 2F). 9,79,88,110

Lysholm Knee Scoring Scale. Moderate certainty of evidence from 3 studies (102 individuals) showed that individuals with PFP have reduced self-reported function measured using the Lysholm Knee Scoring Scale compared with pain-free individuals (large effect size, SMD = -2.23; 95% CI = -3.51, -0.96; $I^2 = 82\%$; P < .001; Figure 2G). $I^2 = 82\%$ (100) Figure 2G).

Performance-Based Measures of Function

Forward Step-Down Test. Moderate certainty of evidence from 9 studies (737 individuals) showed that individuals with PFP have fewer repetitions in the FSDT

^{**}References 8–10, 22, 27–29, 35–81, 83–90, 92–99, 101, 103–106, 108–110.

^{**} References 9, 10, 29, 59, 61, 62, 66, 67, 97, 99, 110.

^{††} References 8–10, 27, 29, 35–60, 63–67, 103–105, 110.

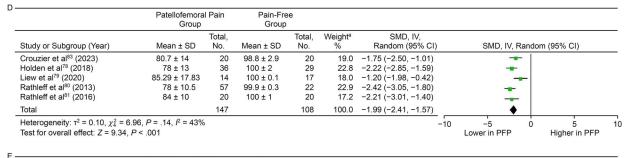
^{**} References 29, 42, 60, 88, 89, 92–96, 108, 109.

	Patellofemoral Group	Pain	Pain-Fre Group	е			
Ot and a ser Oak sure and O/2 and	Marris OD	Total,	Mara LOD	Total,	Weighta	SMD, IV,	CMD IV Denders (059/ CI)
Study or Subgroup (Year)	Mean ± SD	No.	Mean ± SD	No.	%	Random (95% CI)	SMD, IV, Random (95% CI)
Antunez et al ⁶³ (2023)	78.17 ± 8.12	75	100 ± 0.1	18	2.7	-2.96 (-3.63, -2.28)	-
Baellow et al ³⁷ (2020)	74.47 ± 7.8	15	100 ± 0.1	15	2.1	-4.50 (-5.92, -3.09)	
Baellow et al ³⁸ (2022)	77.26 ± 9.8	35	100 ± 0.1	35	2.7	-3.25 (-3.97, -2.52)	
Bley et al ³⁹ (2014)	80.2 ± 4.9	20	99.5 ± 1.2	20	2.2	-5.30 (-6.67, -3.93)	
Branco et al ⁶⁷ (2023)	83.77 ± 10.41	26	95.08 ± 6.06	24	2.8	-1.29 (-1.91, -0.68)	-
Carlson et al ⁴⁰ (2017)	61.1 ± 16.5	12	100 ± 0.1	13	2.3	-3.30 (-4.56, -2.03)	
Coelho et al ⁴² (2021)	76.1 ± 9.2	48	100 ± 0.1	48	2.8	-3.64 (-4.31, -2.98)	-
de Albuquerque et al ³⁵ (2021)	74.04 ± 7.2	26	98.96 ± 2.48	24	2.4	-4.48 (-5.55, -3.41)	
de Albuquerque et al ³⁶ (2022)	74.8 ± 11.7	40	100 ± 0.1	40	2.8	-3.02 (-3.67, -2.37)	-
de Moura Campos Carvalho E Silva et al ⁴¹ (2014)	78.9 ± 17.2	20	98.4 ± 2.3	20	2.7	-1.56 (-2.27, -0.84)	-
de Oliveira Silva et al ⁴³ (2015)	72.6 ± 9.2	31	100 ± 0.1	31	2.6	-4.16 (-5.06, -3.25)	
de Oliveira Silva et al44 (2018)	71.96 ± 9.53	158	100 ± 0.1	165	2.9	-4.20 (-4.59, -3.81)	-
de Oliveira Silva et al ⁴⁵ (2018)	72.9 ± 10	65	98.4 ± 2.5	53	2.8	-3.33 (-3.89, -2.76)	-
Felicio et al46 (2012)	77.9 ± 8.8	19	99 ± 2.3	20	2.5	-3.25 (-4.24, -2.27)	-
Ferreira et al ⁴⁷ (2021)	69.34 ± 11.7	38	100 ± 0.1	38	2.7	-3.67 (-4.42, -2.92)	
Ferreira et al ⁶⁴ (2023)	75.27 ± 7.98	11	99.69 ± 0.76	13	2.0	-4.36 (-5.93, -2.79)	
Gallina et al48 (2018)	74.3 ± 8.1	36	100 ± 0.1	20	2.6	-3.89 (-4.81, -2.96)	
Glaviano et al ⁸ (2017)	75.3 ± 7.7	20	100 ± 0.1	20	2.3	-4.45 (-5.64, -3.25)	
Goharpey et al ²⁷ (2007)	78.26 ± 4.43	15	103 ± 1.79	15	1.6	-7.12 (-9.19, -5.06)	
Guimaraes Araujo et al29 (2023)	74.5 ± 10.5	50	99.2 ± 1.7	50	2.8	-3.26 (-3.86, -2.65)	-
Jaffri and Baellow ⁶⁵ (2023)	77.22 ± 9.2	30	100 ± 0.1	30	2.6	-3.46(-4.27, -2.64)	-
Jeon et al ¹¹⁰ (2023)	76.3 ± 12.8	12	99.2 ± 2.5	12	2.4	-2.40 (-3.49, -1.31)	
Kalytczak et al ⁴⁹ (2016)	80.43 ± 4.78	14	99.5 ± 1.29	14	1.9	-5.29 (-6.96, -3.62)	
Kızılkaya and Ecesoy ⁵⁰ (2019)	70.57 ± 8.37	30	98.58 ± 2.05	31	2.5	-4.57 (-5.55, -3.59)	
Muniz et al ⁵¹ (2023)	86.8 ± 6.8	13	99.7 ± 0.8	18	2.5	-2.84 (-3.89, -1.80)	
Novello et al ⁵² (2018)	67.35 ± 9.33	34	100 ± 0.1	34	2.5	-4.89 (-5.86, -3.92)	
Nunes et al ⁹ (2019)	76.3 ± 12.5	16	99.5 ± 1.1	16	2.5	-2.55 (-3.51, -1.59)	
O'Sullivan et al ¹⁰³ (2012)	78 ± 9	12	100 ± 0.1	12	2.2	-3.34 (-4.64, -2.03)	
Ott et al104 (2011)	82.8 ± 9.98	20	99.4 ± 1.85	20	2.6	-2.27 (-3.08, -1.46)	
Pavone et al ⁵³ (2022)	98.47 ± 12.96	38	98.9 ± 15.3	13	2.8	-0.03 (-0.66, 0.60)	+
Pazzinatto et al ⁵⁴ (2017)	80.45 ± 4.55	20	100 ± 0.1	20	2.1	-5.95 (-7.46, -4.45)	
Pazzinatto et al ⁵⁵ (2019)	70.3 ± 11.84	30	97.7 ± 6.27	30	2.7	-2.85 (-3.59, -2.12)	-
Plastaras et al ⁵⁶ (2016)	85.7 ± 10.1	21	97.6 ± 5.7	36	2.8	-1.54 (-2.16, -0.93)	-
Sanchis-Alfonso et al ⁶⁶ (2023)	49.2 ± 17	44	98.4 ± 2.7	50	2.7	-4.14 (-4.87, -3.41)	-
Souza et al ¹⁰⁵ (2017)	72.3 ± 7.14	14	99.2 ± 1.92	10	2.0	-4.62 (-6.26, -2.97)	
van der Heijden et al ⁵⁷ (2018)	66.3 ± 11.6	64	99.5 ± 1.7	70	2.8	-4.07 (-4.67, -3.47)	-
Willson and Davis ⁵⁸ (2008)	80.2 ± 8.5	20	100 ± 0.2	20	2.5	-3.23 (-4.20, -2.26)	
Yilmaz Yelvar et al ⁵⁹ (2017)	49.09 ± 15.78	22	100 ± 0.1	22	2.4	-4.48 (-5.63, -3.34)	
Zamboti et al ⁶⁰ (2017)	78.4 ± 16.5	10	98.9 ± 2.02	10	2.5	-1.67 (-2.72, -0.62)	
Zamboti et al ¹⁰ (2021)	70.5 ± 11.61	20	96.7 ± 4.8	20	2.6	-2.89 (-3.80, -1.98)	
Total	70.0 1 11.01	1244	50.7 2 1.0	1170	100.0	-3.45 (-3.84, -3.06)	•
Heterogeneity: $\tau^2 = 1.33$, $\chi^2_{39} = 335$ Test for overall effect: $Z = 17.30$, P		88%					-10 -5 0 5 Lower in PFP Higher in PF

	Patellofemoral Group	Pain	Pain-Fre Group	е			
		Total,		Total,	Weighta	SMD, IV,	
tudy or Subgroup (Year)	Mean ± SD	No.	Mean ± SD	No.	%	Random (95% CI)	SMD, IV, Random (95% CI)
aellow et al ³⁷ (2020)	80.66 ± 10.92	15	100 ± 0.1	15	11.1	-2.44 (-3.41, -1.46)	- [
oglund et al ⁶⁸ (2018)	68.17 ± 7.82	36	79.72 ± 0.78	36	11.6	-2.06 (-2.63, -1.48)	-
affri and Baellow ⁶⁵ (2023)	63.03 ± 8.17	30	80 ± 0.1	30	11.4	-2.90 (-3.64, -2.16)	-
im et al ⁶⁹ (2020)	60 ± 4.3	26	78.9 ± 0.8	30	10.5	-6.25 (-7.56, -4.94)	-
im and Park ⁷⁰ (2022)	55.2 ± 2.7	60	79.1 ± 0.5	48	9.9	-11.62 (-13.25, -10.00)	-
im et al ⁷³ (2023)	58.8 ± 9.1	22	79.2 ± 1.3	19	11.2	-2.97 (-3.88, -2.06)	-
im et al ⁷² (2024)	55.44 ± 11.06	55	79.3 ± 1.83	55	11.6	-2.99 (-3.54, -2.44)	-
an Cant et al ⁷¹ (2017)	58.6 ± 11.1	20	78.7 ± 1.9	76	11.4	-3.78 (-4.52, -3.05)	-
amboti et al ⁶⁰ (2017)	74.7 ± 11.32	10	79.4 ± 0.7	10	11.2	-0.56 (-1.46, 0.34)	
otal		274		319	100.0	-3.83 (-5.10, -2.55)	•

Mean ± SD	Total,		T-1-1					
	No.	Mean ± SD	Total, No.	Weight ^a %	SMD, IV, Random (95% CI)	SMD, IV, Rando	m (95% CI)	
12.3 ± 0.89	30	16 ± 0.1	30	14.5	-5.77 (-6.95, -4.59)			_
10.2 ± 2.3	14	15.7 ± 0.7	14	14.5	-3.14 (-4.30, -1.99)			
12.1 ± 0.78	20	16 ± 0.1	20	13.8	-6.87 (-8.58, -5.17)			
11.58 ± 0.77	24	16 ± 0.1	24	13.7	-7.92 (-9.67, -6.17)			
79 ± 16.6	43	96 ± 10.3	40	15.0	-1.21 (-1.68, -0.74)	-		
12.03 ± 0.89	27	16 ± 0.1	27	14.3	-6.18 (-7.50, -4.86)			
11.35 ± 1.72	14	16 ± 0.1	10	14.3	-3.39 (-4.72, -2.07)	-		
	172		165	100.0	-4.87 (-6.97, -2.77)	•		
1	10.2 ± 2.3 12.1 ± 0.78 11.58 ± 0.77 79 ± 16.6 12.03 ± 0.89 11.35 ± 1.72	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$

Figure 2. Forest plots for self-reported function meta-analyses comparing individuals with patellofemoral pain and pain-free individuals. A, Anterior Knee Pain Scale. B, Lower Extremity Functional Scale. C, Functional Index Questionnaire. D, Knee injury and Osteoarthritis Outcome Score. E, Activities of Daily Living Questionnaire. F, Knee injury and Osteoarthritis Outcome Score for PFP and osteoarthritis. G, Lysholm Knee Scoring Scale. Abbreviations: IV, inverse variance; SMD, standardized mean difference. Continued on next page.



	Patellofemora Group	l Pain	Pain-Fre Group								
Study or Subgroup (Year)	Mean ± SD	Total, No.	Mean ± SD	Total, No.	Weight ^a %	SMD, IV, Random (95% CI)		SMD, IV,	Rando	m (95% CI)	
de Moura Campos Carvalho E Silva et al ⁸⁴ (2016)	74.4 ± 7.2	25	99.5 ± 0.8	30	15.3	-5.07 (-6.19, -3.95)		-			
Guimaraes Araujo et al29 (2023)	76.8 ± 12.8	50	99.5 ± 1.8	50	21.9	-2.46 (-2.99, -1.94)		-			
Kiliç et al85 (2021)	61.4 ± 11.5	30	79.8 ± 0.6	30	20.6	-2.23 (-2.88, -1.58)		-	-		
Magalhães et al ⁸⁶ (2010)	73.6 ± 17.16	50	99.8 ± 0.8	50	22.2	-2.14 (-2.64, -1.65)		-	-		
Piva et al ⁸⁷ (2005)	64.5 ± 18.4	30	99.8 ± 0.08	30	19.9	-2.68 (-3.39, -1.97)		•			
Total		185		190	100.0	-2.79 (-3.49, -2.08)		•			
Heterogeneity: $\tau^2 = 0.51$, $\chi_4^2 = 23$.	24, P < .001, I ² =	83%					-10	-5	0	5	10
Test for overall effect: $Z = 7.77$, P	< .001							Lower in PFP		Higher in PFP	

	Patellofemora Group	Patellofemoral Pain Group		e						
Study or Subgroup (Year)	Mean ± SD	Total, No.	Mean ± SD	Total, No.	Weight ^a %	SMD, IV, Random (95% CI)		SMD, IV, Ra	andom (95% CI)	
Eckenrode et al88 (2023)	76.79 ± 7.62	17	98.89 ± 1.73	20	22.0	-4.09 (-5.27, -2.91)			1	_
Jeon et al110 (2023)	65.8 ± 17.3	12	96.3 ± 7.8	12	24.4	-2.19 (-3.24, -1.15)		-		
Liew et al79 (2020)	68.99 ± 18.9	14	99.47 ± 1.71	17	26.5	-2.33 (-3.28, -1.39)		-		
Nunes et al ⁹ (2019)	67.3 ± 19.3	16	98.9 ± 1.7	16	27.2	-2.25 (-3.16, -1.34)		-		
Total		59		65	100.0	-2.66 (-3.47, -1.86)		•		
Heterogeneity: $\tau^2 = 0.41$, $\chi_3^2 = 7$ Test for overall effect: $Z = 6.48$,)%					-10	-5 Lower in PFP	0 5 Higher in PFF	10

	Patellofemora Group	Patellofemoral Pain Group		ee							
Study or Subgroup (Year)	Mean ± SD	Total, No.	Mean ± SD	Total, No.	Weight ^a %	SMD, IV, Random (95% CI)		SMD, IV, R	ando	m (95% CI)	
Ingram et al ⁹⁰ (2016)	73.3 ± 13.7	12	98 ± 6.3	10	31.4	-2.16 (-3.25, -1.06)		-	- 1		
Kiliç et al85 (2021)	70.4 ± 11.5	30	98.6 ± 3.5	30	35.4	-3.27 (-4.06, -2.48)		-			
Zamboti et al ⁶⁰ (2017)	82.7 ± 18.42	10	99 ± 2.11	10	33.1	-1.19 (-2.16, -0.22)		-	-		
Total		52		50	100.0	-2.23 (-3.51, -0.96)		•	-		
Heterogeneity: $\tau^2 = 1.03$, $\chi_2^2 = 1.03$ Test for overall effect: $Z = 3.44$		82%					-10	-5 Lower in PFP	0	5 Higher in PFP	10

Figure 2. Continued from previous page.

compared with pain-free individuals (large effect size, SMD = -0.80; 95% CI = -1.11, -0.50; $I^2 = 68\%$; P < .001; Figure 3).§§ Very low certainty of evidence from 2 studies (106 individuals) showed no differences between limbs for repetitions in the FSDT in individuals with unilateral PFP (small effect size, SMD = -0.36, 95% CI = -1.11, 0.38; $I^2 = 73\%$; P = .34; Figure 4).^{97,99}

Single-Leg Hop Test. Moderate certainty of evidence from 7 studies (711 individuals) indicated a shorter distance in the SLHT for individuals with PFP than for pain-free individuals (small effect size, SMD = -0.42; 95% CI = -0.57, -0.27; $I^2 = 0\%$; P < .001; Figure 3). $^{9.42,45,61,62,93,101}$ Evidence from 1 high-quality study showed a shorter distance in the SLHT for the PFP limb compared with the pain-free limb of individuals with unilateral PFP, but this result was not confirmed via the calculated SMD and 95% CI (small effect size, SMD = -0.29; 95% CI = -0.86, 0.28; P = .32; Figure 4). 99

Single-Leg Triple-Hop Test. Moderate certainty of evidence from 2 studies (196 individuals) showed a shorter distance in the single-leg triple-hop test (SLTHT) in individuals with PFP than pain-free individuals (small effect size, SMD = -0.30, 95% CI = -0.58, -0.02; $f^2 = 0\%$; P = .04; Figure 3).^{29,42}

Bilateral Squat Test. Moderate certainty of evidence from 2 pooled studies (70 individuals) showed no difference in the number of repetitions in the bilateral squat test between individuals with PFP and pain-free individuals (large effect size, SMD = -1.21; 95% CI = -2.71, 0.29; $I^2 = 86\%$; P = .11; Figure 3).

Between-Limbs Comparisons for Other Performance-Based Measures. Evidence from 1 study showed fewer repetitions for the anteromedial lunge in the PFP limb compared with the pain-free limb of individuals with unilateral PFP (moderate effect size, SMD = -0.64; 95% CI = -1.17, -0.11; P = .02) (Figure 4).⁹⁷ The same study showed fewer repetitions for the balance-and-reach test and lower scores in the single-leg press test in the PFP limb compared with the pain-free limb of individuals with unilateral PFP; however, this result

^{§§} References 9, 27, 45, 61, 62, 93, 97, 98, 105.

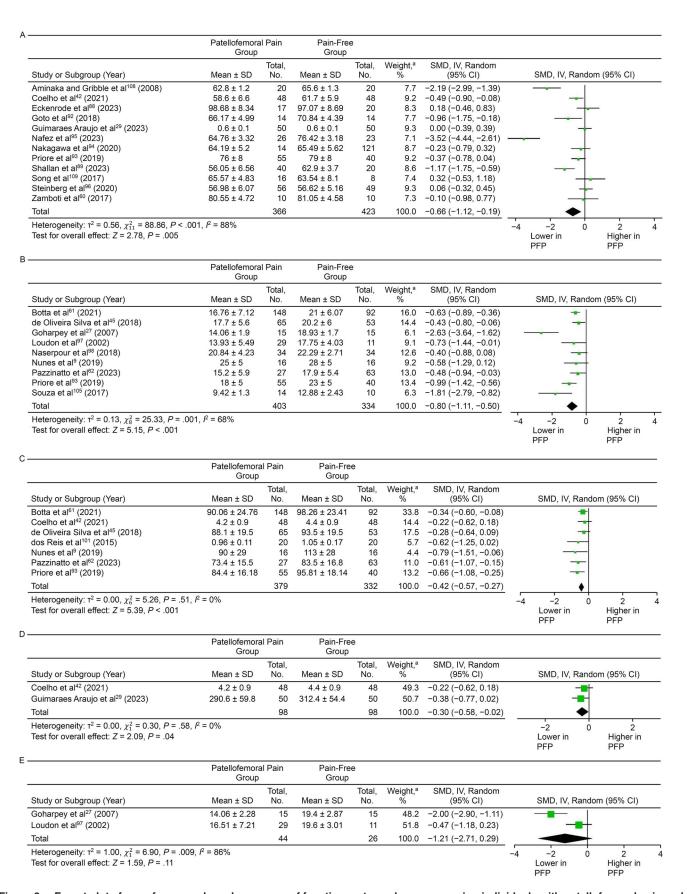


Figure 3. Forest plots for performance-based measures of function meta-analyses comparing individuals with patellofemoral pain and pain-free individuals. A, Balance tests. B, Forward step-down test. C, Single-legged hop test. D, Single-legged triple-hop test. E, Bilateral squat test. Abbreviations: IV, inverse variance; SMD, standardized mean difference.

		Patellofemora Group	l Pain	Pain-Free Group	Э			
	Study or Subgroup (Year)	Mean ± SD	Total, No.	Mean ± SD	Total, No.	Weight, %	SMD, IV, Random (95% CI)	SMD, IV, Random (95% CI)
A	Eckenrode et al ⁸⁸ (2023) Miller et al ²² (2013)	98.68 ± 8.34 91.08 ± 9.47	17 18	98.94 ± 9.19 94.84 ± 10.16	17 18	49.1 50.9	-0.03 (-0.70, 0.64) -0.37 (-1.03, 0.29)	-
	Subtotal		35		35	100.0	-0.20 (-0.68, 0.27)	•
	Heterogeneity: $\tau^2 = 0.00$, χ^2 Test for overall effect: $Z = 0$		$I^2 = 0\%$					
3	Kaya et al ⁹⁹ (2011)	9.9 ± 24	24	9.5 ± 2.3	24	49.2	0.02 (-0.54, 0.59)	
	Loudon et al ⁹⁷ (2002)	13.93 ± 5.49	29	17.31 ± 3.28	29	50.8	-0.74 (-1.27, -0.20)	
	Subtotal Heterogeneity: $\tau^2 = 0.21$, χ^2	² = 3.68 <i>P</i> = 0.6	53 $p^2 = 73^{\circ}$	%	53	100.0	-0.36 (-1.11, 0.38)	
	Test for overall effect: $Z = 0$		7 - 73	70				
С	Kaya et al ⁹⁹ (2011)	86.4 ± 15.9	24	91.1 ± 16.1	24	100.0	-0.29 (-0.86, 0.28)	-
	Subtotal		24		24	100.0	-0.29 (-0.86, 0.28)	•
	Heterogeneity: not applicable Test for overall effect: Z = 1							
)	Loudon et al ⁹⁷ (2002)	17.93 ± 5.55	29	19.9 ± 5.6	29	100.0	-0.35 (-0.87, 0.17)	-
	Subtotal		29		29	100.0	-0.35 (-0.87, 0.17)	•
	Heterogeneity: not applicable Test for overall effect: $Z = 1$							
Ξ	Loudon et al ⁹⁷ (2002)	11.72 ± 3.07	29	13.56 ± 2.58	29	100.0	-0.64 (-1.17, -0.11)	-
	Subtotal		29		29	100.0	-0.64 (-1.17, -0.11)	•
	Heterogeneity: not applicable Test for overall effect: $Z = 2$							
F	Loudon et al ⁹⁷ (2002)	13.69 ± 4.69	29	16 ± 4.52	29	100.0	-0.49 (-1.02, 0.03)	-
	Subtotal		29		29	100.0	-0.49 (-1.02, 0.03)	•
	Heterogeneity: not applicable Test for overall effect: $Z = 1$							
	Test for subgroup difference	es: χ ₅ ² = 1.76, <i>P</i> :	= .88, <i>f</i> ²	= 0%				-2 0 2
		10.70						Lower Higher in PFP in PFP Limb Limb

Figure 4. Forest plots for performance-based measures of function meta-analyses comparing patellofemoral pain and contralateral painfree limbs of individuals with unilateral patellofemoral pain. A, Balance tests. B, Forward step-down test. C, Single-legged hop test. D, Balance-and-reach test. E, Anterior lunge. F, Single-legged press test. Abbreviations: IV, inverse variance; SMD, standardized mean difference.

was not confirmed by the SMDs and 95% CIs (small effect sizes, SMD = -0.35; 95% CI = -0.87, 0.17; P = .19 and SMD = -0.49; 95% CI = -1.02, 0.03; P = .06, respectively) (Figure 4).⁹⁷

Meta-Regressions

We could only perform meta-regressions for self-reported function measured using the AKPS and the following predictors: age (in years), BMI, duration of symptoms (in months), and self-reported pain (score). Meta-regressions could be performed only for performance-based function measured using balance tests and age (in years) (Supplemental Figure 2).

Meta-regression results indicated no relationship between self-reported function and age (40 studies||||; $R^2 < .01$, P = .54); BMI (20 studies||||; $R^2 < .01$, P = .18); duration of

^{***}References 8–10, 27, 29, 35–60, 63–67, 103–105, 110.

[¶] References 10, 27, 35, 36, 39, 40, 44, 45, 47–50, 52, 55, 57, 59, 60, 64, 103, 104.

symptoms (21 studies***; $R^2 < .01$, P = .86); and worst level of pain in the last month, week, or 24 to 72 hours (15 studies***; $R^2 = .02$, P = .15) (Supplemental Figure 2). No relationship was observed between performance-based function and age (12 studies; $R^2 < .01$, P = .91; Supplemental Figure 2).††

DISCUSSION

We identified 83 studies investigating self-reported or performance-based measures of function in individuals with PFP. Moderate to high certainty of evidence demonstrated that individuals with PFP have impaired selfreported function compared with pain-free individuals, regardless of instrumentation. Very low to moderate certainty of evidence also demonstrated between-groups (ie, individuals with PFP vs pain-free individuals) but not between-limbs (ie, painful vs pain-free limb of individuals with PFP) differences for most of the performance-based measures of function. Reduced performance between individuals with PFP and pain-free individuals was observed in tasks simulating dynamic balance (eg, YBT), stepping (eg, FSDT), or hopping (eg, SLHT). Our results highlight the negative effect of PFP on self-reported and performancebased measures of function. However, none of the predictors investigated in our study (ie, age, BMI, duration of symptoms, and self-reported pain) could explain impaired self-reported or performance-based function in individuals with PFP.

Self-Reported Function

We identified that individuals with PFP have impaired self-reported function compared with pain-free individuals. This finding is based on large effects across 7 different questionnaires/scales and supports previous evidence considering self-reported measures of function as determinants of treatment success or patient recovery after nonoperative management of PFP. ^{12,15} Impaired self-reported function as a consequence of having PFP not only affects individuals' perception and perspectives about their physical function^{5,111} but also can predict persistent or recurrent PFP (ie, poor prognosis) in the medium and long term. ^{11,112,113} Along with previous evidence, our results highlight the need to consider self-reported function as one of the primary condition-specific outcome measures during rehabilitation of individuals with PFP.

The assessment of self-reported function is clinician friendly and strongly recommended by the REPORT-PFP. PRecommended questionnaires include the AKPS (also known as the *Kujala scale*) and the KOOS-PF. Although the AKPS is by far the most commonly used questionnaire (42 studies included in our review), the recently developed KOOS-PF. seems to have better content validity, reliability, construct validity, and responsiveness. The KOOS-PF is also more feasible for clinical use due to the its ease of administration and scoring, smaller number of

items, and short time to complete. 116 Changes in KOOS-PF ranging from 16 to 17.2 have been suggested to detect meaningful differences postintervention. 116 However, only 5 studies included in our review used the KOOS-PF. 9,79,88,89,110 More studies using the KOOS-PF to assess self-reported function of individuals with PFP are needed to further support the recommendation for using this tool instead of the AKPS.

Physical Performance

Performance-based measures of function can complement information from self-reported measures by objectively quantifying functional impairments using physical performance tests that are clinically accessible, low-cost, and time efficient. 117,118 We identified that individuals with PFP have impaired physical performance compared with pain-free individuals during balance tests FSDT, SLHT, and SLTHT. These tests are easily measured in clinical settings, and they represent aspects of daily function or sport and simulate common pain-provoking tasks (eg, stepping, jumping, and landing). 93,94,119 Given that self-reported function does not fully reflect the magnitude of performance deficits, we recommend using performance-based measures of function when assessing individuals with PFP. Whereas balance tests and FSDT may be more useful for evaluating sedentary or lower-functioning individuals due to their reduced physical demand, the SLHT may be used for athletic populations as it is more challenging. In a recent review, Berg et al also recommended the use of the SLHT for assessing performance deficits of youth and young adults, given its sufficient intrarater reliability, construct validity, and responsiveness. 117

Although we observed that performance during balance tests FSDT, SLHT, and SLTHT was impaired when compared with that of controls, we found no differences between the painful and pain-free limbs of individuals with unilateral PFP. This suggested that the functional performance of the pain-free limb may be also compromised in individuals with PFP, as recently reported by Waiteman et al. 120 Individuals with PFP have reduced physical activity, which may result in reduced bilateral lower limb muscle strength and physical performance, regardless of pain laterality.^{8,121} Reduced pain-free limb performance may also be a consequence of fear-avoidance belief, a commonly reported trait in this population. 7,93 Caution is warranted when comparing limbs using performance-based measures of function in clinical practice, as the pain-free limb of individuals with unilateral PFP may not be an accurate comparator. 122 In the absence of reference values of performance-based measures of function for individuals with PFP, we encourage pretest and posttest bilateral assessments to aid clinicians in their decision-making processes because changes may occur bilaterally. 123,124 Future research is needed to provide reference values from ageand sex-matched pain-free samples, as the use of the contralateral limb does not seem appropriate.

Predictors of Self-Reported or Performance-Based Function

We performed meta-regressions to investigate physical and nonphysical factors that might explain poor self-reported and

^{***} References 8–10, 35, 37, 38, 41, 42, 46, 47, 50, 51, 53–55, 57, 59, 60, 63, 64, 110.

^{***} References 8, 9, 36–39, 43, 44, 47, 49, 52, 54, 55, 65, 105.

^{†††} References 29, 42, 60, 88, 89, 92–96, 108, 109.

^{‡‡‡} References 8–10, 27, 29, 35–67, 103–105.

performance-based function in individuals with PFP. We observed no relationships between self-reported function and age, BMI, or duration of symptoms. Similarly, no relationship between performance-based function and age was observed. Although researchers have reported a direct relationship between these factors and clinical outcomes in PFP (eg, higher BMI was related to lower functional capacity), our findings showed that this relationship does not explain differences between groups. 125,126 This means that individuals with PFP have lower self-reported and performancebased function compared with pain-free controls regardless of age, BMI, or duration of symptoms. Another reason that we did not observe relationships between function and these variables is that, as commonly reported in PFP, most individuals from the studies included in this review had normal BMI (mean \pm SD BMI of individuals with PFP = 23.04 \pm 3.58 kg/m^2 and of pain-free individuals = $22.30 \pm 3.05 \text{ kg/m}^2$) and were young adults (mean ± SD age of individuals with PFP = 22.91 \pm 6.55 years and of pain-free individuals = 23.24 \pm 6.23 years). This results in a constrained range of age and BMI across studies and may have influenced statistical analysis. Other factors, such as quadriceps strength, kinesiophobia, or both, may be more associated with impaired self-reported or performance-based function, as previously reported. 13,127,128 Quadriceps strength and kinesiophobia have been reported to be associated with pain intensity in individuals with PFP, which plays an important role in the perception of disability and function. ^{7,129,130} These uncontrolled factors may also be the source of potential heterogeneity between the studies. More studies are needed to better understand what physical and nonphysical factors might explain impaired self-reported and performance-based function in individuals with PFP. Furthermore, more longitudinal studies are necessary to investigate how function changes across time in individuals with PFP.

We also did not observe a relationship between self-reported function and self-reported pain. Glaviano et al reported high pain variability in individuals with PFP over 10 days, which explained almost 60% of the variance in self-reported function. One should assume that traditional methods of assessing pain (eg, worst levels of pain in the last month or week) used in the studies included in our review may not be sensitive to pain variation and are also more susceptible to recall bias. Longitudinal pain assessments of daily pain variability over a period may be better suited to investigate the relationship between self-reported pain and function in individuals with PFP versus isolated pain observations.

Limitations

The design of studies included in the review did not allow us to infer causality of self-reported and performance-based measures of function in individuals with PFP. We did not review the gray literature; thus, relevant but unpublished studies may have been excluded from our findings. In addition, the limited number of studies did not allow us to investigate whether other important physical and nonphysical predictors (eg, hip and knee strength, physical activity level, and psychological factors) may be more associated with self-reported and performance-based measures of function. More studies following the REPORT-PFP guidelines are needed to fill this gap in the literature.

CONCLUSIONS

Individuals with PFP have impaired self-reported and performance-based function compared with pain-free individuals. Our results also suggest a negative effect of PFP on performance-based measures of function on the pain-free limb of individuals with unilateral PFP. No physical or non-physical factors explain impaired self-reported function in individuals with PFP. Both self-reported and performance-based measures of function should be clinically assessed when treating individuals with PFP.

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SUPPLEMENTAL MATERIAL

Supplemental Figure 1. Funnel plots assessing publication bias in studies included in meta-analyses of A, Anterior Knee Pain Scale of individuals with patellofemoral pain compared with pain-free individuals, and B, balance tests of individuals with patellofemoral pain compared with pain-free individuals.

Supplemental Figure 2. Meta-regressions for self-reported function measured using A, Anterior Knee Pain Scale and age, B, Anterior Knee Pain Scale and body mass index, C, Anterior Knee Pain Scale and duration of symptoms, and D, Anterior Knee Pain Scale and self-reported pain. E, Meta-regression for performance-based function measured using balance tests and age.

Supplemental Material 1. Protocol deviations.

Supplemental Material 2. Data-extraction management.

Supplemental Material 3. Internal and external validity of studies assessed using the modified Downs and Black checklist.

Supplemental Material 4. Summary of included studies.

Supplemental Material 5. Synthesis of unpooled data.

Supplemental Table 1. Search strategy.

Supplemental Table 2. Study quality of Miller et al¹ assessed using Physiotherapy Evidence Database scale.

Supplemental Table 3. Risk of bias of Miller et al²² assessed using Cochrane Risk of Bias in Randomized Trials 2.

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