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This file contains both the full text version of record and the supplementary files. The supplementary material appears at the end of this file, after the references section in the main text.





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What are small, medium and large effect sizes for exercise treatments of tendinopathy? A systematic review and meta-analysis

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ABSTRACT

Objective To quantify and describe effect size distributions from exercise therapies across a range of tendinopathies and outcome domains to inform future research and clinical practice through conducting a systematic review with meta-analysis.

Design Systematic review with meta-analysis exploring moderating effects and context-specific small, medium and large thresholds.

Eligibility criteria Randomised and quasi-randomised controlled trials involving any persons with a diagnosis of rotator cuff, lateral elbow, patellar, Achilles or gluteal tendinopathy of any severity or duration.

Methods Common databases, six trial registries and six grev literature databases were searched on 18 January 2021 (PROSPERO: CRD42020168187). Standardised mean difference (SMD_{pre}) effect sizes were used with Bayesian hierarchical meta-analysis models to calculate the 0.25 (small), 0.5 (medium) and 0.75 quantiles (large) and compare pooled means across potential moderators. Risk of bias was assessed with Cochrane's Risk of Bias tool. Results Data were obtained from 114 studies comprising 171 treatment arms 4104 participants. SMD_{pre} effect sizes were similar across tendinopathies but varied across outcome domains. Greater threshold values were obtained for self-reported measures of pain (small=0.5, medium=0.9 and large=1.4), disability (small=0.6, medium=1.0 and large=1.5) and function (small=0.6, medium=1.1 and large=1.8) and lower threshold values obtained for quality of life (small=-0.2, medium=0.3 and large=0.7) and objective measures of physical function (small=0.2, medium=0.4 and large=0.7). Potential moderating effects of assessment duration, exercise supervision and symptom duration were also identified, with greater pooled mean effect sizes estimated for longer assessment durations, supervised therapies and studies comprising patients with shorter symptom durations.

Conclusion The effect size of exercise on tendinopathy is dependent on the type of outcome measure assessed. Threshold values presented here can be used to guide interpretation and assist with further research better establishing minimal important change.

WHAT IS ALREADY KNOWN ON THIS TOPIC

⇒ Exercise therapy, in particular resistance exercise, is frequently used in the management of tendinopathy and is known to have general effectiveness across a range of important outcome domains. There is, however, a lack of research comparing effectiveness across different tendinopathies and outcome domains.

WHAT THIS STUDY ADDS

⇒ This large and comprehensive meta-analysis shows that exercise therapy results in relatively wide change distributions relative to baseline. Distributions of standardised mean difference effect sizes appear consistent across the most common tendinopathies. In contrast, substantive differences exist in the distributions of standardised mean difference effect sizes across outcome domains.

HOW THIS STUDY MIGHT AFFECT RESEARCH, PRACTICE OR POLICY

⇒ The results of this study provide researchers and clinicians with important information regarding how individuals should be expected to respond to exercise therapy for the management of tendinopathies, thereby influencing decisions regarding the effectiveness of any intervention and how to power future research studies.

BACKGROUND

Tendinopathy is a common musculoskeletal condition associated with degenerative changes and characterised by a combination of pain impaired movement and reduced function that typically requires extended periods for recovery. Tendinopathy can affect any muscle-tendon unit in the body, however, it is most frequently reported in the Achilles, patellar, lateral elbow, rotator cuff and hip tendons. Surveys of prevalence of lower extremity tendinopathy in the general population have reported rates of 11.8 and



10.5 per 1000 person-years, while prevalence for upper limb tendinopathies have been estimated between 1.3% and 21.0%. Solvential Costs to the individual, the health service and economy (due to absenteeism and loss of productivity) are substantial such that identifying effective interventions is a priority.

Exercise therapy is the mainstay of conservative management of tendinopathy and has focused largely on resistance training, and in many instances eccentric strengthening techniques.¹¹ The rationale of exercise therapy is to improve load tolerance and possibly structural adaptation of the musculotendinous unit to restore function. 12 13 In the early phase of rehabilitation, flexibility exercises are often initiated and incorporated into strengthening regimes to facilitate improvements in mobility.¹¹ Effective exercise therapy may also require targeting a range of contributing factors, which not only include muscle weakness and decreased flexibility, but also corticospinal and neuromuscular adaptations resulting from persistent pain. 14 As such, proprioceptive exercise interventions have been used to retrain normal patterns of muscle recruitment in the rehabilitation of shoulder-related tendinopathies including impingement. 14-17 The time course of tendinopathy recovery is usually slow, the degree of recovery may be incomplete and there may be differences between tendinopathies and across outcome domains. For example, quality of life may improve less quickly for people with rotator cuff tendinopathy compared with those with Achilles tendinopathy, whereas recovery may be faster for pain and the relative magnitudes of these improvements may not be equivalent. Quantifying any tendinopathy or domainspecific differences in expected improvements would help guide efforts to develop consensus concerning optimal management by enabling better intervention comparisons.

A recent scoping review identified a lack of effective tools to draw general conclusions across the large tendinopathy and exercise therapy research base (~450 primary studies), which featured a wide range of interventions across different tendinopathies, populations and outcome domains. 18 At present, one of the main tools to synthesise information and therein draw general conclusions include the use of meta-analyses. Most previous meta-analyses have attempted to quantify the effectiveness of interventions using standardised mean difference (SMD) effect sizes and Cohen's standard benchmarks (small=0.2, medium=0.5 and large=0.8) irrespective of the tendinopathy location, population or outcome domain. 19-27 Despite Cohen's recommendations that these general benchmarks should only be used where more relevant context-specific information is unavailable,²⁸ use of these standard benchmarks is ubiquitous through behavioural, social and health sciences. However, recent attempts have been made across a range of disciplines to use empirically derived effect size distributions to generate context-specific benchmarks providing better means of establishing the effectiveness of different

interventions and drawing general conclusions.²⁹⁻³⁵ Results have frequently demonstrated substantive differences between Cohen's benchmarks and those derived empirically, with examples of both underestimation and overestimation, and even differences across subdomains within a discipline. ²⁹ In addition, SMDs are used frequently as a means of informing the minimal important change (MIC) for patients, especially when preferred anchorbased approaches using external criterions such as global ratings of change are not available.³⁶ Given the range of tendinopathies and outcome domains commonly investigated, there is potential that the distribution and subsequent appropriate interpretation of therapy effects will be diverse and could benefit from the generation of context-specific benchmarks. Therefore, the purpose of this meta-analysis was to perform a large synthesis of the available research creating empirically derived thresholds to benchmark the effectiveness of exercise therapies and explore potential differences across tendinopathies and outcome domains. The analysis also investigated the potential for moderating effects of commonly reported features including assessment duration of outcomes, therapy supervision (supervised vs unsupervised) and symptom duration of patients. The results of this analysis will provide clinicians and researchers with tendinopathyspecific and domain-specific indicators of effect sizes with which to better interpret intervention outcomes.

METHODS

This meta-analysis is part of a project funded by the National Institute for Health Research; Health Technology Assessment 129 388 Exercise therapy for the treatment of tendinopathies. The inclusion criteria were influenced by the project aims, the results of our initial scoping review mapping the exercise and tendinopathy literature, as well as stakeholder workshops. The overall structure of systematic reviews and meta-analyses to address effectiveness were registered in the PROSPERO database (CRD42020168187) and individual full protocols made publicly available prior to any analyses. The review was conducted according to the PRISMA 2020 statement with checklist provided in online supplemental file 1.

Protocol deviations

Multiple protocol deviations occurred due to pragmatic considerations and reflecting on processes from previous work packages in the larger project. Originally, it was intended to extract data not in duplicate but to quantify reliability based on a random 10% sample. Given the large number of reviewers extracting data it was decided to perform extraction in duplicate with agreement following differences when required. Originally, it was intended to conduct risk of bias using the ROBINS-I tool for quasi-experimental studies. As outlined in the following sections, due to pragmatic considerations Cochrane's Risk of Bias tool was used for both randomised and non-randomised designs.



Stakeholder involvement

People with lived experience of exercise for tendinopathy were involved in all stages of the review, from inception to dissemination activities, and were recruited via National Health Service (NHS) public involvement networks and social media. Two people who had received exercise for Achilles tendinopathy and rotator cuff related shoulder pain (RCRSP) in NHS and private settings contributed to the design stage by influencing the review questions. One of these people (female, RCRSP) went on to contribute to the oversight committee throughout the review and assisted with reviewing dissemination materials. We also held a stakeholder workshop to inform the direction of the review; four females with lived experience of RCRSP or patellar tendinopathy, including one high performance athlete, took part. We had anticipated greater public involvement; timing of activities (during COVID-19 lockdown) and conducting them solely online may be contributing factors. Nonetheless, contributions were helpful in informing the review.

Inclusion criteria

Participants

This meta-analysis included people of any age or gender with a diagnosis of RCRSP, lateral elbow, patellar, Achilles or gluteal tendinopathy of any severity or duration. Studies that included participants with tendinopathy in the absence of full thickness or large tears were included. We accepted trial authors' diagnoses where a clearly verifiable group of clinical features is reported including: pathognomonic location of pain; a symptom altering response to applied load and/or stretch, with there being a specific test for most tendinopathies; strategies to rule out differential diagnoses; ultrasound or MRI confirmation of structural change. We included studies with mixed groups where there was clear reporting of the tendinopathic group, or those participants comprised >90% of the investigated cohort.

Intervention

The intervention being assessed is exercise therapy comprising five different therapy classes: (1) resistance, (2) plyometric, (3) vibration, (4) flexibility and (5) proprioception. Definitions for each therapy class are presented in online supplemental file 2. Interventions combining exercise with other active therapies (eg, laser, shockwave, manual therapy or injection) were not included. We included exercise therapies delivered in a range of settings and delivered by a range of health, exercise professionals or support workers. We also included both supervised and unsupervised exercise therapies. As part of the inclusion criteria, we required studies to report sufficient information regarding the exercise intervention to enable appropriate identification of treatment duration, therapy class and exercise dose. In clinical settings, it has been recommended that exercise dose is determined by duration, frequency and intensity. To be included in the review, we required studies

to provide sufficient information to describe at least two of the three exercise dose parameters. Where sufficient information was not presented in the main text of a study, a search was made of the publishers' website to check for online supplemental files that may include relevant information.

Comparator

No comparators were included, with effect sizes used to quantify the intervention effectiveness of exercise only therapies based on change relative to baseline.

Outcomes

Based on the results of our initial scoping review and subsequent stakeholder workshops, ¹⁸ we included outcomes that assessed six domains: (1) disability, (2) physical function capacity (PFC), (3) function, (4) pain (on loading/activity, over a specified time or without further specification), (5) quality of life and (6) range of motion (ROM) (shoulder joint only). Definitions of each domain and example tools are presented in online supplemental file 3.

Types of studies

We included randomised controlled trials (RCTs) and non-RCTs where at least one intervention arm comprised an exercise therapy that could be categorised according to the therapy classes outlined.

Context

The context included primary care, secondary care or community locations in nations defined as very high or high on the Human Development Index (top 62 countries at the time of protocol development) for the findings to be relevant to the UK context.⁴¹

Exclusion criteria

In addition to coding treatment arm interventions according to exercise criteria, we also coded non-active (eg, wait-and-see, placebo and sham) and non-exercise (eg, electrotherapy, biomechanics, manual therapy, injection therapy and surgery) categories according to preset definitions (online supplemental file 4). Where any of the non-active or non-exercise categories could be assigned to a treatment arm intervention, these were excluded from the review.

Search strategy

The search strategy used for this study was part of a larger search conducted to scope the entire tendinopathy and exercise therapy research base. The search comprised three steps. First, a limited search of MEDLINE and CINAHL using initial keywords was conducted to develop a full search strategy. Second, the full search strategy was adapted to each database and applied systematically to: MEDLINE, CINAHL, AMED, EMBase, SPORTDiscus, Cochrane library (Controlled trials, Systematic reviews), JBI Evidence Synthesis, PEDRo and Epistemonikos (search terms for each database are presented in online

supplemental file 5. In addition, six trial registries, five grey literature data bases and Google Scholar were searched (online supplemental file 5). Finally, the third step involved conducting a search of cited and citing articles using Scopus and handsearching a total of 130 systematic reviews that were identified to include information relevant to exercise therapy and tendinopathy. No limit was placed on language, with research studies reported in languages other than English translated via Google Translate or via international collaborations of the review team members. Searches were initiated from 1998 as (1) the heavy load eccentric calf-training protocol for Achilles tendinosis by Alfredson et al was published in 1998 and may be considered seminal work in the field of tendinopathy⁴² and (2) there has been a proliferation of research on exercise interventions for tendinopathies post 1998. The final date of the search was 18 January 2021.

Study selection

Proquest Refworks was used to manage references and remove duplicates before importing to Covidence (Melbourne, Australia) to facilitate screening and initiate a second deduplication process. Titles/abstracts were reviewed, independently, by two members of the research team. Full-text copies of all studies included at title/abstract screening stage were retrieved and also reviewed, independently, by two members of the research team. Conflicts were resolved by discussion or by input from a third reviewer.

Data extraction

Data were extracted independently by eight members of the review team (PAS/KC/LA/RM/LG/EP/JSCS/AVP) into preiloted Excel sheets. Data were independently coded as described in the accompanying extraction codebook (online supplemental file 6) by two members from the review team. Differences in entries were detected through automatic checking in MS Excel and then agreed between the same two reviewers. Where pre–post intervention data were not presented in text but in figures, data were extracted using digitsation software (PlotDigitizer V.2.6.8 Windows).

Risk of bias

We used Cochrane's Risk of Bias tool⁴⁰ and six outcome domains: (1) selection bias (random sequence generation and allocation concealment), (2) performance bias (blinding of participants), (3) detection bias (blinding of outcome assessors), (4) attrition bias (incomplete outcome data), (5) reporting bias (selective reporting) and (6) other bias, to assess risk of bias for RCTs and domains 2–6 for non-randomised trials. Risk of bias was recorded for each outcome and time point within each study. When obtaining a summary risk of bias for each domain within a study, the mode category across all outcomes and timepoints was selected. The Cochrane's Risk of Bias tool⁴⁰ was selected as a recent review of

popular risk of bias tools in tendinopathy management highlighted none were superior ⁴³ and Cochrane's Risk of Bias tool could be semiautomated with RobotReviewer, ⁴⁴ a machine learning system software. RobotReviewer was used to make initial assessments on selection bias and performance bias domains, with manual validation made on the relevant free texts extracted to support the final selection of low, high or unclear risk of bias. This semiautomated process was more efficient, and pragmatic given the large number of included studies and provided an additional element of consistency in the review process.

Statistical analysis overview

In general, meta-analyses attempt to pool data across studies quantifying effectiveness by focusing on a weighted mean value. Due to the use of different outcome domains and different tests within a specific outcome domain, pooling of data requires initial standardisation. The most common metric used to standardise effectiveness is the SMD_{pre} effect size, which divides the group change by the preintervention SD. Meta-analyses seek to describe the underlying population effect size based on a normal distribution with the mean representing the most likely value across studies, and the SD representing the dispersion that can be expected across individual studies. However, a focus on a single central value provides limited description of the overall distribution that can be expected. By providing estimates of the 0.25 quantile (the value which 25% of observed results are expected to be below), the 0.5 quantile and the 0.75 quantile, a more detailed description is obtained. In addition, these estimates can be used to provide benchmarks to interpret the effectiveness of future interventions, with the traditional qualitative labels of 'small' (0.25 quantile), 'medium' (0.5 quantile) and 'large' (0.75 quantile) used to provide a simple scale.

Most meta-analyses are conducted within a frequentist framework where a focus is placed on the mean value and hypothesis testing with CIs to identify whether they overlap a zero effect. In contrast, analyses performed within a Bayesian framework are more flexible enabling, for example, quantile values to be estimated and can be interpreted intuitively through reporting of subjective probabilities. 45 Given the purpose of the present metaanalysis to describe the overall distribution of effect sizes and explore differences among potentially relevant factors (eg, tendinopathy location and outcome domains), a Bayesian framework was selected. Finally, many meta-analyses only extract a single outcome from each study per-analysis to avoid complexities due to relationships within data. However, more precise estimates of effects may be obtained by including all relevant data but accounting for covariances of multiple study outcomes within the meta-analysis model. A recommended approach to account for hierarchical structures (eg, multiple measurements from the same study, multiple measurements made from difference outcomes within the same study and multiple measurements made from

the same outcome in the same study) is to apply multilevel meta-analysis models, which were included for the present study.

Statistical analysis details

 $\mathrm{SMD}_{\mathrm{pre}}$ effect sizes were calculated by dividing the relevant mean difference by the preintervention SD and including a small sample bias correction.46 Where required, SMD_{pre} values were reflected by multiplying by -1 to ensure that positive values represented an improved clinical effect. Where outcomes were assessed at multiple time points following baseline measurement, all possible SMD_{pre} values were calculated and included in the meta-analysis models. Where means and SDs were not presented but included combinations of the median, range or IOR, values were estimated by the calculations presented by Wan et al. 47 Where sufficient information was not available to estimate SDs, these were imputed through simple linear regression of the log transformed SD and means obtained from all other studies.⁴⁸ Separate regressions were performed for preintervention and postintervention data.

All meta-analyses were conducted using a nested four-level model. 49 The series of nestings included the individual study (level 4), the outcome (level 3), the measurement occasion (level 2) and the sampling variance (level 1). Standard distributional assumptions were used to calculate the sampling variance of SMD_{pre} values. 46 However, the calculation requires an estimate of the pre-post correlation which is rarely reported in studies. To account for uncertainty in the sampling variance, values within the model were allowed to vary and were estimated by including an informative Gaussian prior approximating correlation values centred on 0.7 and ranging from 0.5 to 0.9. The relative contributions of variance sources were described by variance partition coefficients (VPCs) which were calculated by dividing each estimated variance level by the total sum. Therefore, the higher the VPC for the outcome and measurement levels the greater the covariances within the data. Analyses were only completed where a minimum of 50 effect sizes were available to appropriately describe distributions. Sensitivity analyses checking the potential influences of study type (RCTs vs quasi-RCTs) and study quality (low risk of bias: >50% 'low risk' vs high risk of bias: 50% 'low risk') on effect size thresholds were conducted through subset comparisons.

It was determined a priori to assess the influence of tendinopathy location, outcome domain, assessment duration, symptom duration and supervised versus nonsupervised exercise on effect sizes. This was achieved by subset comparisons for tendinopathy location and outcome domain, and meta-regressions for assessment duration, symptom duration and supervision. Meta-regressions were presented by selecting one level of the factor as a reference to make comparisons with the median and 95% CrI ($\beta_{\rm Reference:Comparison}$ = Median (95% CrI: lower bound to upper bound), such that $\beta{>}0$

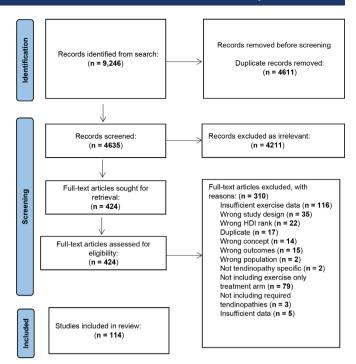


Figure 1 Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) flow diagram describing study selection.

indicates an increased effect of the comparison relative to the reference).

The importance of removing outliers to obtain accurate estimates of meta-analysis parameters was identified in a previous and similar large meta-analysis of exercise SMD_{pre} values. ⁴⁹ Outlier SMD_{pre} values were identified by adjusting the empirical distribution by a Tukey *g*-and-*h* distribution and obtaining the 0.0125 and 0.9875 quantiles, with values beyond these points removed prior to further analysis. ⁵⁰ Meta-analyses were conducted using the R wrapper package brms interfaced with Stan to perform sampling. ⁵¹ Convergence of parameter estimates were obtained for all models with Gelman-Rubin R-hat values below 1.1. ⁵² Data used for the analyses and R code are presented in online supplemental files 7 and 8.

RESULTS

Study selection

The search strategy identified a total of 9246 potential studies, with 4635 remaining following removal of duplicates (figure 1). After title and abstract screening 4210 studies were removed leaving 425 studies obtained for full text screening. Of these studies, a further 311 were excluded based primarily on insufficient description of the exercise stimulus (116 studies) and not including exercise-only treatment arms (75 studies). In total, data from 114 studies (100 RCTs and 14 quasi-experimental) comprising 171 treatment arms and 4104 participants were included in the meta-analyses. A table of includes studies along with reference lists of included and excluded studies are presented in online supplemental files 9 and 10. Risk of bias expressed for each individual study are



Table 1 Risk of bias assessment for randomised (top; N=100) and non-randomised trials (bottom; N=14) with percentages of low-risk, unclear-risk and high-risk evaluations expressed relative to the number of treatment arms (upper value) and the total number of data points (lower value)

		Random sequence allocation	Allocation concealment	Blinding of participants	Blinding of outcome assessors	Incomplete outcome bias	Selective reporting	Other bias
Randomised controlled trials	Low risk	78% 80%	61% 61%	28% 29%	58% 62%	63% 69%	28% 35%	46% 51%
	Unclear	18% 16%	34% 34%	30% 31%	24% 16%	23% 17%	67% 60%	7% 7%
	High risk	4%	5%	42% 40%	18% 22%	14%	5% 5%	47% 42%
Non-randomised controlled trials	Low risk	NA	NA	39% 55%	39% 31%	77% 89%	15% 18%	8% 26%
	Unclear	NA	NA	39% 32%	61% 69%	23% 11%	85% 82%	15% 19%
	High risk	NA	NA	22% 13%	0% 0%	0%	0%	77% 55%

presented in online supplemental file 11 with summaries presented in table 1. For RCTs, risk of bias was highest for 'other bias' (47% high risk of bias), blinding of participants (42% high risk of bias) and selective reporting (67% unclear risk of bias). For quasi-experimental trials, risk of bias was also highest for 'other bias' (77% high risk of bias) and reporting quality was also lower with high percentages of unclear risk of bias identified for selective reporting (85% unclear risk of bias) and blinding (participants: 39% unclear risk of bias; outcome assessors: 61% unclear high risk of bias).

Descriptions of the study characteristics, tendinopathy location and outcome domains are presented in table 2. Outcomes obtained from studies investigating gluteal tendons were not included in the analysis based on the number of effect sizes falling below the a priori threshold set to generate accurate estimates of the population effect size distribution.

Description of effect size distributions

From the initial 1454 outcomes extracted, a total of 38 outliers were removed from the analysis with a lower

bound threshold of -0.82 (6 effect sizes below) and an upper bound threshold of 7.0 (32 effect sizes above). Across all outcomes and tendinopathy locations, direct calculation of the 0.25 (small), 0.5 (medium) and 0.75 quantiles (large) from the complete empirical data returned the following SMD $_{\mathrm{pre}}$ values: 0.37, 0.77 and 1.31, respectively. Application of the meta-analysis model across the data with borrowing of information across studies resulted in similar but shrunken estimates (0.25 quantile_{0.5} = 0.34 (95% CrI: 0.31 to 0.37); 0.5 quantile_{0.5} = 0.73 $(95\% \text{ CrI: } 0.70 \text{ to } 0.77) \text{ and } 0.75\text{-quantile}_{0.5} = 1.21 (95\%)$ CrI: 1.17 to 1.27)). A forest plot of effect sizes illustrating effect sizes across studies is presented in online supplemental file 12. Sensitivity analyses checking the potential influences of study type and study quality are presented in online supplemental file 13. No evidence was obtained of greater effect sizes with quasi-experimental designs or with studies identified as high risk of bias.

Analyses of effect size distributions across the different tendinopathy locations are illustrated in figure 2 with numerical values presented in table 3. Analyses were



Table 2 Distribution (percentiles) of study characteristics, tendinopathy locations and outcome domains calculated across treatment arms

Study characteristic	0%	10%	20%	30%	40%	50%	60%	70%	80%	90%	100%
Participants per group	5	10	13	16	19	20	24	28	31	44	70
Mean age	22.5	29.5	39.8	42.6	44.0	46.0	47.9	48.7	49.9	51.9	62.1
Mean symptom duration (months)	0.85	4.4	6.0	7.8	11.6	17.5	19.5	24.1	29.7	37.4	98.5
Publication year	1998	2005	2007	2009	2012	2014	2015	2017	2017	2019	2020
Intervention length (weeks)	2	4	4	4	6	8	12	12	12	12	21
Measurement duration (weeks)	0.7	3	4	4	5	6	6	9	12	13	104
Tendinopathy location	No of TA	A (%)	No of eff	fects (%)		Outcom	e domain	No of TA	\ (%)	No of eff	ects (%)
Rotator cuff	77 (45.0))	817 (56.	2)		Disability	У	142 (83.	0)	447 (30.7	7)
Achilles	45 (26.3))	321 (22.	1)		Pain		122 (71.	3)	406 (27.9	9)
Lateral elbow	29 (17.0))	227 (15.	6)							
Patellar	20 (11.7))	89 (6.1)			PFC		59 (34.5))	320 (22.0	O)
Dominant therapy class	No of TA	A (%)	No of eff	fects (%)		ROM		30 (17.5))	159 (10.9	9)
Resistance	124 (72.	5)	1014 (69	0.7)		Function	1	29 (17.0))	68 (4.7)	
Flexibility	25 (14.6))	215 (14.	8)							
Proprioception	21 (12.3))	223 (15.	3)		QoL		12 (7.0)		54 (3.7)	
Vibration	1 (0.6)		2 (0.1)								

Pain, pain without further specification; Pain time, pain over a specified time; PFC, physical function capacity; QoL, quality of life; ROM, range of motion; TA, treatment arms.

pooled across all outcome domains as a means to compare tendinopathy locations with the largest amount of data possible. Analysis of the modelled small, medium and large thresholds showed considerable overlap in small and medium thresholds across all tendinopathy locations (0.25 quantile_{0.5} ranged from 0.28 to 0.38; 0.5 quantile_{0.5} ranged from 0.70 to 0.82). However, greater

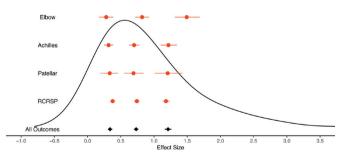


Figure 2 Effect size distributions across tendinopathy locations with identification of small, medium and large thresholds. black curve represents density plot of empirical effect size distribution. Diamonds with intervals represent small, medium and large thresholds with credible intervals (black: all outcomes; red: tendinopathy specific). RCRSP, rotator cuff related shoulder pain.

divergence was identified for large threshold estimates, with the greatest values estimated for the elbow $(0.75 \text{ quantile}_{0.5} \text{ ranged from } 1.18 \text{ to } 1.49)$.

Analyses of effect size distributions across outcome domains are presented in figure 3 with numerical values presented in table 4. A clear split was identified between the domains of quality of life and the objective measures of PFC and ROM, versus the subjective measures of function, disability and pain. The lowest threshold values were estimated for quality of life, PFC and ROM, with the small threshold for quality of life estimated to be below zero $(0.25 \text{ quantile}_{0.5} = -0.21 (95\% \text{ CrI: } -0.32 \text{ to } -0.09)).$ In contrast, the greatest effect sizes values were obtained for outcomes measuring disability, pain and function with the reduced amount of data for function resulting in wider credible intervals. Central estimates indicated that small threshold estimates for domains with the greatest effect sizes were situated between the medium and large threshold estimates for domains with the lowest effect sizes (figure 3).

Moderator analyses

Moderator analyses investigating potential changes in the mean effect size across all outcomes and tendinopathies

Table 3 Meta-analysis results for all outcomes pooled across different tendinopathy locations

Crl, credible interval; RCRSP, rotator cuff related shoulder pain; VPC, variance partition coefficient.

Tendinopathy Location	Small (95%CrI)	Medium (95%CrI)	Large (95%Crl)	Study VPC (75%CrI)	Outcome VPC (75%Crl)	Measurement VPC (75%CrI)
All	0.34	0.73	1.21	0.73	0.27	0.01
	(0.31 to 0.37)	(0.70 to 0.77)	(1.17 to 1.27)	(0.68 to 0.77)	(0.22 to 0.32)	(0.00 to 0.02)
RCRSP	0.38	0.74	1.18	0.55	0.45	0.01
	(0.34 to 0.42)	(0.70 to 0.78)	(1.13 to 1.24)	(0.48 to 0.62)	(0.37 to 0.52)	(0.00 to 0.02)
Achilles	0.32	0.70	1.22	0.71	0.27	0.01
	(0.25 to 0.38)	(0.62 to 0.78)	(1.10 to 1.34)	(0.59 to 0.81)	(0.18 to 0.39)	(0.00 to 0.03)
Elbow	0.28	0.82	1.49	0.90	0.10	0.01
	(0.17 to 0.39)	(0.72 to 0.92)	(1.31 to 1.68)	(0.84 to 0.94)	(0.06 to 0.15)	(0.00 to 0.02)
Patellar	0.33	0.69	1.21	0.46	0.52	0.01
	(0.19 to 0.46)	(0.55 to 0.84)	(1.00 to 1.43)	(0.13 to 0.69)	(0.29 to 0.85)	(0.00 to 0.05)

are presented in table 5. Evidence of a moderator effect was identified for assessment duration (short: ≤ 12 weeks, medium: 13–52 weeks and long duration: >52 weeks), with results showing a hierarchy and greater mean estimate with increased time from baseline assessment ($\beta_{\text{Short:Medium0.5}} = 0.28$ (95% CrI: 0.23 to 0.33), p>0.999; $\beta_{\text{Short:Long0.5}} = 0.37$ (95% CrI: 0.27 to 0.47), p>0.999). These estimates remained consistent after meta-regressions were controlled for tendinopathy location and outcome domain ($\beta_{\text{Short:Medium0.5}} = 0.28$ (95% CrI: 0.23 to 0.33), p>0.999; $\beta_{\text{Short:Long0.5}} = 0.38$ (95% CrI: 0.27 to 0.48), p>0.999). Consistent evidence of a moderating effect was also obtained for exercise supervision, with a greater mean

Small: 0.25 quantile; medium: 0.5 quantile; large: 0.75 quantile.

Function

Disability

Pain

Physical function capacity

QoL

ROM – Shoulder

PFC

Outcomes

-1.0 -0.5 0.0 0.5 1.0 1.5 2.0 2.5 3.0 3.5

Effect Size

Figure 3 Effect size distributions across outcome domains with identification of small, medium and large thresholds. Each curve represents density plot of empirical effect size distribution for specific outcome domain. Density curves and effect size thresholds are presented in same colours. PFC, physical function capacity; QoL, quality of life; ROM, range of motion.

estimate for supervised exercise therapies ($\beta_{Unupervised:Supervised0.5} = 0.38$ (95% CrI: 0.05 to 0.71), p=0.989), which increased in estimate after controlling for tendinopathy location and outcome domain ($\beta_{Unupervised:Supervised0.5} = 0.66$ (95% CrI: 0.35 to 0.98), p>0.999). Finally, some evidence was obtained indicating a hierarchy of effects with regards to symptom duration and greater mean estimates with patients reporting shorter symptom durations ($\beta_{Medium:Short0.5} = 0.33$ (95% CrI: -0.12 to 0.81), p=0.920; $\beta_{Long:Short0.5} = 0.51$ (95% CrI 0.08 to 0.96), p=0.989). However, differences between the symptom duration levels were close to zero after controlling for tendinopathy location and outcome domain ($\beta_{Medium:Short0.5} = -0.05$ (95% CrI: -0.44 to 0.36), p=0.411; $\beta_{Long:Short0.5} = 0.06$ (95% CrI: -0.34 to 0.47), p=0.619).

DISCUSSION

The present analysis represents the largest quantitative synthesis of exercise therapy interventions for the management of tendinopathies to date. Data from a total of 114 studies were included, with results demonstrating that substantive improvements in outcomes from baseline are generally obtained. With important clinical relevance, the meta-analyses identified clear differences in the distribution of effects sizes across outcome domains. The greatest values were generally obtained for subjective patient reported outcomes including disability and pain-related outcomes, with considerably lower values obtained for measures of quality of life and objective measures including PFC and ROM. Considerable overlap in effect size distributions were identified across the different tendinopathy location investigated, indicating that similar substantive improvements can be obtained with exercise therapies. Moderator analyses provided consistent evidence of increased improvement in outcomes as time increased from baseline assessment and with supervised compared with non-supervised exercise therapies. Some evidence was obtained for greater



Table 4 Meta-analysis results for all tendinopathy locations pooled across different outcome domains

Outcome domain	Small (95%Crl)	Medium (95%Crl)	Large (95%Crl)	Study VPC (75%Crl)	Outcome VPC (75%Crl)	Measurement VPC (75%Crl)
Function	0.62	1.05	1.78	0.43	0.55	0.01
	(0.45 to 0.77)	(0.88 to 1.23)	(1.53 to 2.09)	(0.03 to 0.81)	(0.17 to 0.94)	(0.00 to 0.04)
Disability	0.61	1.04	1.51	0.78	0.15	0.05
	(0.55 to 0.68)	(0.98 to 1.11)	(1.43 to 1.60)	(0.68 to 0.88)	(0.08 to 0.24)	(0.00 to 0.13)
Pain	0.53	0.94	1.45	0.26	0.71	0.02
	(0.45 to 0.61)	(0.87 to 1.02)	(1.32 to 1.58)	(0.06 to 0.46)	(0.51 to 0.90)	(0.00 to 0.07)
ROM	0.21	0.42	0.65	0.86	0.11	0.01
	(0.14 to 0.27)	(0.36 to 0.48)	(0.58 to 0.73)	(0.76 to 0.93)	(0.05 to 0.20)	(0.00 to 0.06)
PFC	0.16	0.38	0.64	0.73	0.27	0.01
	(0.11 to 0.21)	(0.34 to 0.43)	(0.59 to 0.71)	(0.61 to 0.82)	(0.18 to 0.38)	(0.00 to 0.02)
QoL	-0.21	0.25	0.68	0.84	0.14	0.01
	(-0.32 to -0.09)	(0.12 to 0.38)	(0.48 to 0.93)	(0.65 to 0.93)	(0.05 to 0.33)	(0.00 to 0.04)

Small: 0.25 quantile; medium: 0.5 quantile; large: 0.75 quantile.

Crl, credible interval; PFC, physical function capacity; QoL, quality of life; ROM, range of motion; VPC, variance partition coefficient.

improvements with patients reporting symptoms for shorter durations, however, differences did not remain after controlling for outcome domain and tendinopathy location.

Of the 114 studies included in the analysis, 100 (88%) were RCTs and 14 (12%) were quasi-experimental trials. Therapies predominantly focused on resistance exercise, with 70% of outcomes obtained from a treatment arm where this was the dominant class. Tendinopathy has been clinically defined as persistent pain and loss of function related to mechanical loading on a degenerative tendon. Therefore, resistance exercise that focuses on restoration of loading ability is established as the mainstay

of rehabilitation, particularly for lower limb tendinopathies, and is in keeping with current guidance. Across the included studies flexibility and proprioceptive training were frequently combined with resistance exercise. However, these alternative therapy classes were rarely the dominant exercise class, and where they were, this tended to be restricted to management of RCRSP which accounted for over 50% of outcomes measured in the studies. Given the focus on resistance exercise and the desire to change the mechanical properties of the tendon, the standard duration of exercise therapies featured in the included studies is potentially a major limitation. While clear reporting of therapy duration

Moderator		Estimate of mean (95%Crl)	Probabilities	Study VPC (75%CrI)	Outcome VPC (75%CrI)	Measurement VPC (75%CrI)
Assessment duration	Short (≤12 weeks) 923 outcomes from 144 trials	0.97 (0.85 to 1.1)	p(Medium>Short) > 0.999	0.71 (0.66 to 0.75)	0.27 (0.23 to 0.32)	0.02 (0 to 0.04)
	Medium (13–52 weeks) 442 outcomes from 23 trials	1.2 (1.1 to 1.4)	p(Long>Medium) = 0.980			
	Long (>52 weeks) 51 outcomes from two trials	1.4 (1.2 to 1.6)	<i>p</i> (Long>Short) > 0.999			
Symptom duration	1 year or less 308 outcomes from 44 trials	1.4 (1.1 to 1.7)	p(1yr>2yr) = 0.920	0.81 (0.76 to 0.84)	0.17 (0.14 to 0.21)	0.03 (0.00 to 005)
	2 years or less 258 outcomes from 30 trials	1.1 (0.74 to 1.4)	p(2yr>+2yr) = 0.772			
	Over 2 years 381 outcomes from 33 trials	0.88 (0.56 to 1.2)	p(1yr>+2yr) = 0.989			
Supervision	Supervised 354 outcomes from 35 trials	1.4 (1.1 to 1.7)	p(Supervised> Unsupervised)	0.77 (0.73 to 0.81)	0.20 (0.17 to 0.24)	0.03 (0.00 to 0.05)
	Unsupervised 914 outcomes from 112 trials	0.99 (0.83 to 1.2) = 0.989				

only occurred in 49% of studies, where it was reported, the median duration was only 8 weeks and over 90% of studies included durations of 12 weeks or shorter. Tendon healing is known to be a complex and lengthy process, with remodelling only beginning 1–2 months postinjury and extending beyond 1 year, suggesting that longer duration exercise interventions are required, which is in keeping with guidelines recommending a minimum 12 weeks duration. ⁵⁵ It is also relevant to note that in clinical practice, education and exercise programmes are frequently provided for patients to continue therapy after the initial intervention which may also contribute to the findings observed here including evidence of greater improvements with increased time from baseline assessment.

Review of previous meta-analyses and individual studies investigating the effectiveness of exercise therapy for tendinopathy shows that Cohen's standard benchmarks are most frequently used to interpret effectiveness. 19-27 56 The results of the present study show that across most outcomes Cohen's standard benchmarks of 0.2 (small), 0.5 (medium) and 0.8 (large) would tend to result in an overestimation of effectiveness and for quality of life may result in an underestimation.²⁸ Based on the analyses performed in the present meta-analysis, more appropriate thresholds include ~0.4 (small), ~0.8 (medium) and ~1.3 (large), highlighting a systematic shift by one category (eg, what would have been referred to as a medium/large effect is more representative of a small/ medium effect). However, the results of the analyses clearly show that effect size distributions are strongly dependent on the outcome domain, with more subjective patient-reported outcomes (eg, pains scales and patient-rated levels of function and disability) producing substantively larger effects compared with more objective assessments (eg, ROM and quantitative measures of PFC). A wide range of outcome measures are used for tendinopathies, with little consensus in the literature to date, ¹⁸ although this should improve in future with work ongoing to identify core outcome sets for specific tendinopathies. International consensus reported that nine outcome domains should be considered core for tendinopathy.⁵⁷ The majority (8/9) of the consensus domains are patient reported, with only PFC being objectively measured in the clinical setting (eg, number of hops/ squats, dynamometry). The results of this study highlight that when using patient-centred self-reported outcomes, substantial changes should be expected from most exercise therapies.

The lowest effect sizes were obtained for quality of life outcomes with the small threshold indicating that over 25% of exercise therapies represented by those investigated will result in patients' reporting a poorer quality of life. However, it is possible that the generic nature of quality of life instruments (eg, EuroQol-5D instrument) insufficiently reflects tendinopathy-specific symptoms or that potential limitations such as ceiling effects limit the usefulness of these instruments. Tendinopathy may be

acute but is typically due to chronic overuse and degeneration, exacerbated by overloading; therefore, patients may have developed specific coping strategies over time. Concern was raised when developing core domains that there is no tendon-specific quality of life measure, ⁵⁷ but that the overall well-being of patients was important to assess. Further research is required to better understand the factors that influence quality of life assessments when managing tendinopathies and the best measurement tool to use.

Considerable overlap in the effect size thresholds across the five tendinopathies assessed suggests that exercise therapies commonly used to manage the different tendinopathies result in similar profiles of improvement. This contrasts previous perspectives that responses to interventions are variable across tendons and even within tendon sites.⁵⁸ It is likely that the interactions between a single exercise therapy, the tendinopathy and the outcome domain are complex and when better understood can improve patient care. The results of the present analysis, however, indicate that exercise therapies commonly used to manage RCRSP, Achilles, patellar and lateral elbow tendinopathies result in overall similar responses with most causing relatively large changes in relation to baseline SDs. A limitation of the analysis was that it pooled data across all outcome domains to provide sufficient information to compare distributions across the tendinopathies. As the amount of primary data increases, more refined analyses should be conducted to better investigate potential differences across the most common tendinopathies and those which are at present are under researched.

Despite almost all exercise therapies being of short duration (≤12 weeks), a substantial proportion of outcomes were measured beyond the intervention at a medium duration (13-52 weeks), and a relatively small number of instances of longer-term follow-up (>52 weeks). Moderator analysis demonstrated an ordered effect with the smallest mean pooled effect size obtained for short durations (1.0 (95% CrI: 0.87 to 1.1)), and evidence of greater pooled means for medium (1.3 (95%) CrI: 1.1 to 1.4); p>0.999) and long durations (1.4 (95%) CrI: 1.2 to 1.6) p>0.999). While the absolute magnitude of the differences were relatively small (~0.2 to 0.4 increase) and there were only two studies that included long term observations, the finding are in keeping with previous research,59 and provide support for longer duration interventions as well as follow-up periods in studies. Moderator analyses also provided evidence of a greater mean pooled effect with supervised compared with unsupervised exercise therapies ($\beta_{Unupervised:Supervised.5}$ = 0.70 (95% CrI: 0.39 to 1.0), p=0.797). Previous systematic reviews specifically investigating supervised versus unsupervised exercise therapy for the rotator cuff have suggested that both approaches are likely to lead to similar improvements. 60 61 While previous reviews employed greater control and focused on more homogeneous comparisons and considerably smaller number



of outcomes, the findings in the present analysis and the consistency of results after controlling for tendinopathy location and outcome, demonstrate that in general supervised exercise therapies are likely to provide clinically meaningful improvements beyond unsupervised exercise therapies. Finally, limited evidence was obtained to indicate that greater improvements may be obtained with patients reporting shorter symptom durations. However, differences in pooled mean estimates were close to zero after controlling for tendinopathy location and outcome domains, indicating the need for further research, and highlighting limitations of large modelling studies such as those included here, where systematic differences in a range of potential moderating factors can bias results and generate spurious associations.

Other limitations to consider when interpreting and evaluating the results from the present study include precision of estimates given the reliance on published data and limitations of effect sizes to identify clinical significance. Whist attempts were made to include results from unpublished studies, none met the inclusion criteria and it is known that biases influencing individual studies such as publication bias can result in misleading results in meta-analyses.⁶² Publication bias, however, is most relevant to analyses of pairwise effect sizes, studies employing cohort designs and analyses focusing on null hypothesis testing. When comparing two interventions that are both effective, pairwise comparisons may fail to identify significant differences thereby reducing the chances of dissemination in peer-reviewed journals. In the present study, analyses were conducted on non-controlled effect sizes which are less likely to be influenced by this bias, except in cohort designs that were excluded. Multiple statistical techniques have been developed to correct for potential biases that can skew the results of meta-analyses.⁶² In general, these approaches are best implemented in traditional metaanalyses where the mean effect and confidence intervals are used to make statements regarding a null hypothesis. In contrast, in the present study, inferences focused on describing the majority of the effect size distribution and mitigating biases including potential overestimation of effect sizes through a meta-analysis model that shrunk estimates based on borrowing of strength across studies and accounted for dependencies in the data such that large single values from individual studies had less influence. The use of quantiles to describe distributions and attempts to remove outliers is also likely to have reduced the influence of biases. The importance of removing outliers was highlighted in a previous analysis of SMD effect sizes following exercise interventions based on implausible values due to large underestimations of SD. 49

In addition to challenges associated with estimates, the present analysis is limited in the ability to address clinical significance. While benchmarking effect sizes using empirical values within a specific context is important, ⁶³ the labelling of thresholds as small, medium and large remains somewhat arbitrary. In contrast,

anchor-based approaches that use an external criterion such as global rating of change are viewed as a superior method to establish important thresholds such as the MIC. 36 64 Where anchor-based thresholds have not been developed, attempts have been made to use effect sizes such as those presented here (distribution-based approach) as a surrogate. Research, however, has identified a lack of consistency between distribution and anchor-based approaches where it has been argued effect size thresholds should only be used pending development of well-established anchor-based MID values.³⁶ Further research is required to establish MIC values for the management of tendinopathy, where the effect sizes reported in the present study may be used to inform values where results from anchor-based approaches diverge.³⁶ To assist with clinical interpretations, the threshold values presented can also be transformed into the original scales of measurement.⁶⁵ Using suitable estimates of population means and SDs, effect sizes can be transformed into typical change scores. For example, previous studies of tendinopathy patients have reported baseline Victorian Institute of Sport Assessment-Achilles questionnaire (VISA-A) disability scores of 64±17, and EQ-5D quality of life scores of 0.75±0.15.6667 From table 4, we can compute that a medium effect size for VISA-A scores would reflect an increase from 64 to $64+1.04\times17\approx82$, and that a small effect size for EQ-5D-5L would reflect a decrease from 0.75 to 0.75-0.21×0.15 \approx 0.72.

In conclusion, the results from this large meta-analysis show that relative to baseline assessment, a reasonably wide distribution of changes and in general, improvements, should be expected following exercise therapy to manage tendinopathy. The magnitudes of improvement appear somewhat independent of the location of the tendinopathy, but are strongly influenced by the outcome domain, with the greatest improvements measured in subjective patient-reported outcomes (eg, disability, function, pain) and the smallest improvements measured in quality of life and more objective outcomes (eg, PFC and ROM). When interpreting the effectiveness of exercise therapies for the management of tendinopathies, clinicians and researchers should be aware of these factors and can use the context-specific information presented here as a guide. Further research is required to better establish clinical significance using MIC and anchorbased approaches, where the information presented here may assist should divergent results be obtained.

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and clinical interpretation of results. All authors contributed to the final manuscript. KC is the overall quarantor of this work.

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Supplementary File 1: PRISMA 2020 Checklist and abstract checklist.

Section and Topic	Item #	Checklist item	Location where item is reported
TITLE			
Title	1	Identify the report as a systematic review.	1
ABSTRACT			
Abstract	2	See the PRISMA 2020 for Abstracts checklist.	SF-1
INTRODUCTION	0		
Rationale	3	Describe the rationale for the review in the context of existing knowledge.	2
Objectives	4	Provide an explicit statement of the objective(s) or question(s) the review addresses.	5
METHODS Eligibility criteria	5	Specify the inclusion and exclusion criteria for the review and how studies were grouped for the syntheses.	6
Information sources	6	Specify all databases, registers, websites, organisations, reference lists and other sources searched or consulted to identify studies. Specify the date when each source was last searched or consulted.	8
Search strategy	7	Present the full search strategies for all databases, registers and websites, including any filters and limits used.	SF-5
Selection process	8	Specify the methods used to decide whether a study met the inclusion criteria of the review, including how many reviewers screened each record and each report retrieved, whether they worked independently, and if applicable, details of automation tools used in the process.	8
Data collection process	9	Specify the methods used to collect data from reports, including how many reviewers collected data from each report, whether they worked independently, any processes for obtaining or confirming data from study investigators, and if applicable, details of automation tools used in the process.	8
Data items	10a	List and define all outcomes for which data were sought. Specify whether all results that were compatible with each outcome domain in each study were sought (e.g. for all measures, time points, analyses), and if not, the methods used to decide which results to collect.	7
	10b	List and define all other variables for which data were sought (e.g. participant and intervention characteristics, funding sources). Describe any assumptions made about any missing or unclear information.	7
Study risk of bias assessment	11	Specify the methods used to assess risk of bias in the included studies, including details of the tool(s) used, how many reviewers assessed each study and whether they worked independently, and if applicable, details of automation tools used in the process.	9
Effect measures	12	Specify for each outcome the effect measure(s) (e.g. risk ratio, mean difference) used in the synthesis or presentation of results.	12
Synthesis methods	13a	Describe the processes used to decide which studies were eligible for each synthesis (e.g. tabulating the study intervention characteristics and comparing against the planned groups for each synthesis (item #5)).	8
	13b	Describe any methods required to prepare the data for presentation or synthesis, such as handling of missing summary statistics, or data conversions.	9
	13c	Describe any methods used to tabulate or visually display results of individual studies and syntheses.	12
	13d	Describe any methods used to synthesize results and provide a rationale for the choice(s). If meta-analysis was performed, describe the model(s), method(s) to identify the presence and extent of statistical heterogeneity, and software package(s) used.	10
	13e	Describe any methods used to explore possible causes of heterogeneity among study results (e.g. subgroup analysis, meta-regression).	14
	13f	Describe any sensitivity analyses conducted to assess robustness of the synthesized results.	11
Reporting bias assessment	14	Describe any methods used to assess risk of bias due to missing results in a synthesis (arising from reporting biases).	NA
Certainty assessment	15	Describe any methods used to assess certainty (or confidence) in the body of evidence for an outcome.	14
RESULTS			

Section and Topic	Item #	Checklist item	Location where item is reported		
Study selection	16a	Describe the results of the search and selection process, from the number of records identified in the search to the number of studies included in the review, ideally using a flow diagram.	13		
	16b	Cite studies that might appear to meet the inclusion criteria, but which were excluded, and explain why they were excluded.	SF-10		
Study characteristics	17	Cite each included study and present its characteristics.	SF-9		
Risk of bias in studies	18	Present assessments of risk of bias for each included study.	SF-11		
Results of individual studies	19	For all outcomes, present, for each study: (a) summary statistics for each group (where appropriate) and (b) an effect estimate and its precision (e.g. confidence/credible interval), ideally using structured tables or plots.	SF-7		
Results of	20a	For each synthesis, briefly summarise the characteristics and risk of bias among contributing studies.	SF-11		
syntheses	20b	Present results of all statistical syntheses conducted. If meta-analysis was done, present for each the summary estimate and its precision e.g. confidence/credible interval) and measures of statistical heterogeneity. If comparing groups, describe the direction of the effect.			
	20c	Present results of all investigations of possible causes of heterogeneity among study results.	14		
	20d	Present results of all sensitivity analyses conducted to assess the robustness of the synthesized results.	SF-13		
Reporting biases	21	Present assessments of risk of bias due to missing results (arising from reporting biases) for each synthesis assessed.	NA		
Certainty of evidence	22	Present assessments of certainty (or confidence) in the body of evidence for each outcome assessed.	14		
DISCUSSION					
Discussion	23a	Provide a general interpretation of the results in the context of other evidence.	16		
	23b	Discuss any limitations of the evidence included in the review.	16		
	23c	Discuss any limitations of the review processes used.			
	23d	Discuss implications of the results for practice, policy, and future research.	20		
OTHER INFORMA	1				
Registration and protocol	24a	Provide registration information for the review, including register name and registration number, or state that the review was not registered.	6		
protocor	24b	Indicate where the review protocol can be accessed, or state that a protocol was not prepared.	6		
	24c	Describe and explain any amendments to information provided at registration or in the protocol.	6		
Support	25	Describe sources of financial or non-financial support for the review, and the role of the funders or sponsors in the review.	21		
Competing interests	26	Declare any competing interests of review authors.	21		
Availability of data, code and other materials	27	Report which of the following are publicly available and where they can be found: template data collection forms; data extracted from included studies; data used for all analyses; analytic code; any other materials used in the review.	Supplementary files		

From: Page MJ, McKenzie JE, Bossuyt PM, Boutron I, Hoffmann TC, Mulrow CD, et al. The PRISMA 2020 statement: an updated guideline for reporting systematic reviews. BMJ 2021;372:n71. doi: 10.1136/bmj.n71

For more information, visit: http://www.prisma-statement.org/

Section and Topic	Item #	Checklist item	Reported (Yes/No)
TITLE			
Title	1	Identify the report as a systematic review.	Yes
BACKGROUND	•		
Objectives	2	Provide an explicit statement of the main objective(s) or question(s) the review addresses.	Yes
METHODS			
Eligibility criteria	3	Specify the inclusion and exclusion criteria for the review.	Yes
Information sources	4	Specify the information sources (e.g. databases, registers) used to identify studies and the date when each was last searched.	Yes
Risk of bias	5	Specify the methods used to assess risk of bias in the included studies.	Yes
Synthesis of results	6	Specify the methods used to present and synthesise results.	Yes
RESULTS			
Included studies	7	Give the total number of included studies and participants and summarise relevant characteristics of studies.	Yes
Synthesis of results	8	Present results for main outcomes, preferably indicating the number of included studies and participants for each. If meta-analysis was done, report the summary estimate and confidence/credible interval. If comparing groups, indicate the direction of the effect (i.e. which group is favoured).	Yes
DISCUSSION			
Limitations of evidence	9	Provide a brief summary of the limitations of the evidence included in the review (e.g. study risk of bias, inconsistency and imprecision).	Yes
Interpretation	10	Provide a general interpretation of the results and important implications.	Yes
OTHER			
Funding	11	Specify the primary source of funding for the review.	Yes
Registration	12	Provide the register name and registration number.	Yes

From: Page MJ, McKenzie JE, Bossuyt PM, Boutron I, Hoffmann TC, Mulrow CD, et al. The PRISMA 2020 statement: an updated guideline for reporting systematic reviews. BMJ 2021;372:n71. doi: 10.1136/bmj.n71

For more information, visit: http://www.prisma-statement.org/

Supplementary File 2: Exercise therapy definitions

Therapy Class	Definition	Therapy Treatment	Definition
		Concentric Only	Includes movements where force produced overcomes the resistance such that muscle shortening occurs.
	Exercise designed primarily to increase strength of muscles by causing them to produce substantive force against an applied resistance which can take several forms including the mass of the body or its segments, isoinertial resistance, elastic resistance, or strength training equipment such as isokinetic devices. In tendinopathy, the stimulus may also be intended to provoke tendon remodelling, reduce pain and improve function.	Eccentric Only	Includes movements where force produced is less than the resistance such that controlled muscle lengthening occurs.
Resistance		Concentric and eccentric	Includes movements where force produced exceeds the resistance in one phase and is less than the resistance in another such that controlled muscle lengthening and shortening occurs.
		Isokinetic	Uses specialised exercise equipment such that the resistance is adjusted in real-time to ensure joint angular velocity remains constant.
		Isometric	Includes muscular actions against a resistance such that joint angle remains constant.
Flexibility	Exercise designed to increase joint range of motion and extensibility of muscles and/or associated	Static	Joint range of motion actions where the movement is held at or near the end range of motion.

Therapy Class	Definition	Therapy Treatment	Definition
	tissues. Also referred to as range-of-motion exercises or stretching.	Dynamic	Joint range of motion actions where the movement is performed continuously into and out of the end range of motion.
		PNF	Proprioceptive neuromuscular facilitation is a technique combining passive stretching and isometric action to achieve maximum range of motion.
		Ballistic	Uses the momentum of a moving body or a limb to increase joint range of motion, bouncing into (or out of) a stretched position.

	Exercise designed to enhance the sensation of the joint relative to body position and movement,	Sense of joint position and force	Exercise aimed at enhancing the ability to perceive joint position and force with minimal external cues.
Proprioception	sense of force, and to encourage muscular stabilisation of the joint in the absence of external stabilising devices e.g. ankle brace.	Balance	Includes exercise that require the person to keep or return the displacement of centre of gravity over the base of support through various environmental conditions and changes in body position.

Therapy Class	Definition	Therapy Treatment	Definition
		Movement pattern retraining	Exercise aimed at reeducation of motor control and movement patterns that may involve specific retraining of under- or overactive muscles and alteration of kinematic rotation +- translation timing between body segments. May also be termed motor control or stabilisation.
Plyometric	Exercise where a resistance is overcome by a muscle rapidly stretching then shortening	Plyometric	Exercise where a resistance is overcome by a muscle rapidly stretching then shortening.
Vibration	Exercise where body segments are held stationary or actively displaced as per definitions for other treatment classes whilst applying a rapid oscillating resistance	Vibration	Exercise where body segments are held stationary or actively displaced as per definitions for other treatment classes whilst applying a rapid oscillating resistance

Supplementary File 3: Outcome domain definitions and example tools

Domain	ICON Definition	Example Tools
Disability	Composite scores of a mix of patient-rated pain & disability due to the pain, usually relating to tendon-specific activities/tasks	VISA scales; DASH; quick DASH; SPADI; Patient-rated tennis-elbow evaluation questionnaire; Constant Murley Score; WORC (Western Ontario Rotator Cuff Index); AOFAS (American Orthopaedic Foot & Ankle Society); Roles and Maudsley score; ASES (American Shoulder & Elbow Surgeons Index; Tegner activity score; Lysholm knee scale; Pain free function questionnaire; Ankle activity score; Subjective elbow Value (SEV); Placzek score; Shoulder disability questionnaire; International Knee Documentation Committee form (IKDC); Penn Shoulder score (university of Pennsylvania shoulder score) (PSS); Brief pain inventory (BPI); UCLA Shoulder Rating Scale; FILLA - functional index of leg and lower limb; Neer Shoulder Score; Nirschl phase rating scale; American Shoulder and Elbow Surgeon's (MASES) questionnaire; Mayo Elbow Performance Score (MEPS); Shoulder rating questionnaire (SRQ)
Function:	Participant/patient rated level of function (and not referring to the intensity of their pain; eg, Patient Specific Function Scale on a VAS or NRS).	Patient-specific functional scale
Pain: Pain on loading/activity	Patient reported intensity of pain performing a task that loads the tendon	VAS; NRS; Pain experience scale
Pain: Pain over a specified time	Patient-reported pain intensity over period of time e.g. morning/night/24-hours/1-week	VAS; NRS Painful days in 3 months
Pain: Pain without further specification	Patient asked about pain levels without reference to activity or timeframe	VAS; NRS; Borg CR10 Scale; Pain status
Physical function capacity	Quantitative measures of physical tasks (e.g. hops, times walk, single leg squat) includes muscle strength	Counter movement jump; One-leg triple hop; Single-leg decline squat; Muscle strength measured by dynamometry (hand-held, isokinetic); Repetition maximum; Manual muscle testing. EQ5D; EQ3D; SF-36 or SF-12; Assessment of
Quality of Life	General wellbeing	Quality of Life (AQoL); Nottingham Health Profile; Gothenburg QoL Instrument
Range of Motion (Shoulder only)	Active or passive range of motion in specified plane, measured in degrees.	Hand-held goniometer; inclinometer

Supplementary File 4: Definitions used to define broad and more specific treatment classes for exclusion

Broad treatment	Definition	More specific treatment class	Definition
Exercise only	Exercise therapy is defined as a regimen or program of physical activities specifically designed and prescribed to correct impairments, restore musculoskeletal function, and/or maintain a state of wellbeing.	Same as broad treatment class	Same as broad treatment class
Non-active (placebo, sham, wait and see)	Includes any appropriate inactive treatment such as waiting list control, sham shockwave, sham laser, sham taping or true placebo.	Same as broad treatment class	Same as broad treatment class
		Electrotherapy	Modality that delivers therapeutic levels of physical energy into a biologic system e.g. soft tissue. Includes shockwave, laser and other systems.
Non-exercise	Active treatments used to treat tendinopathy that do not meet the criteria to be considered	Biomechanics	Treatment using external devices that immobilises (e.g. splinting) or alters the kinematics/kinetics of the limb (e.g. taping, bracing and orthotics).
only	exercise.	Manual-therapy	Manual therapy is the skilled application of "hands-on" techniques to treat soft tissues and joint structures for the purpose of improving pain, increasing range of motion, stimulating tissue repair response, and/or improving function.
		Injection therapy	Injection therapy for tendinopathy typically involves direct

administration of a pharmacologically active drug, or combination of drugs using a syringe and needle or equivalent. It may or may not be imageguided. Includes Autologous, drug, and volumetric types.

Surgery

Any relevant surgical intervention for tendinopathy including minimally invasive peritendinous and open intra-tendinous.

Exercise and non-exercise

Treatment comprising multiple components which collectively meet both exercise and non-exercise criteria

Same as broad treatment class

Same as broad treatment class

Supplementary File 5: Search databases and terms

Search last updated 19/01/2021

Embase (Ovid)	(exercise OR exercise*.mp OR "isometric exercise" OR kinesiotherapy OR
Ellibase (Ovid)	
	Eccentric.mp OR concentric.mp OR "heavy slow resistance".mp OR "isokinetic
	exercise" OR plyometrics OR "muscle stretching" OR "muscle training") AND
	(tendinitis OR Tendinopathy.mp OR "tendon injury" OR "shoulder injury" OR
	"rotator cuff injury" OR "tennis elbow" OR tendin.mp OR tendon.mp OR bursitis
	OR "shoulder impingement syndrome" OR 2posterior tibial tendon dysfunction"
	OR "Greater trochanteric pain syndrome".mp)
CINAHL (EBSCO-host)	(MH Exercise OR AB exercise* OR MH "muscle strengthening" OR MH
	"rehabilitation" OR MH "eccentric contraction" OR TX "heavy slow resistance
	exercis*" OR AB eccentric OR AB concentric OR AB isokinetic OR MH
	"therapeutic exercise") AND (MH tendinopathy OR MH "arm injuries" OR
	"tendon injuries" OR MH tendons OR TX tendin* OR TX tendon* OR AB bursitis
	OR MH Bursitis OR MH "Posterior tibial tendon dysfunction" OR MH "shoulder
	impingement syndrome" OR AB "Greater trochanteric pain syndrome")
Medline (EBSCO-host)	(MH exercise OR AB exercise* OR MH "isometric contraction" OR MH
	rehabilitation OR TX eccentric OR TX concentric OR TX "heavy slow resistance"
	OR TX isokinetic) AND (MH tendinopathy OR MH "shoulder injuries" OR MH
	tendons OR MH "tendon injuries OR TX tendin* OR tendon* OR MH bursitis OR
	AB bursitis OR MH "posterior tibial tendon dysfunction" OR MH "shoulder
	impingement syndrome" OR AB "greater trochanteric pain syndrome")
SPORTDiscus (EBSCO-host)	(DE exercise OR DE "exercise therapy" OR AB exercise* OR TX eccentric OR TX
	concentric OR TX "heavy slow resistance" OR DE "isokinetic exercise" OR DE
	plyometrics OR DE "strength training" OR DE "stretch (physiology)" OR DE
	"isometric exercise" OR DE rehabilitation) AND (DE tendinitis OR DE tendinosis
	OR AB tendinopathy OR DE "tendon injuries" OR "shoulder injuries" OR DE
	"tennis elbow" OR AB tendin* OR AB tendon* OR DE bursitis OR AB "shoulder
	impingement syndrome" OR AB "posterior tibial tendon dysfunction" OR AB
	"greater trochanteric pain syndrome")
Amed (EBSCO-host)	(ZU exercise OR ZU "exercise therapy" OR AB exercise OR ZU "muscle stretching
,	exercises" OR ZU "isometric contraction" OR ZU rehabilitation OR TZ eccentric
	OR TZ concentric OR TX "heavy slow resistance" OR TX isokinetic OR AB
	plyometric) AND (ZU tendinopathy OR ZU "tendon injuries" OR ZU tendons OR
	ZU "shoulder injuries" OR ZU "tennis elbow" OR TX tendin* OR TX tendon* OR
	ZU bursitis OR AB bursitis OR ZU "shoulder impingement syndrome" OR ZU
	"posterior tibial tendon dysfunction" OR AB "greater trochanteric pain
	· · · · · · · · · · · · · · · · · · ·
	syndrome")
Open Grey	Tendinopathy AND exercise
	Tendin* AND exercise
	Tendon AND exercise
Mednar	Tendinopathy AND exercise
	Tendin* AND exercise
	Tendon AND exercise
New York Academy Grey	Tendinopathy AND exercise
Literature Report	Tendin* AND exercise
	Tendon AND exercise
EThOS	Tendinopathy AND exercise
	Tendin* AND exercise
	Tendon AND exercise
	ובוועטוו הויוט בגבו עושב

Google Scholar	Tendinopathy AND exercise
	Tendin* AND exercise
	Tendon AND exercise
JBI Evidence Synthesis	Tendinopathy AND exercise
Cochrane Library	Tendinopathy AND exercise
	Tendin* AND exercise
	Tendon AND exercise
PEDro	Tendinopathy AND exercise
	Tendin* AND exercise
	Tendon AND exercise
Epistemonikos	(tendinopathy OR tendon* OR tendin*) AND exercise
CORE	Tendinopathy AND exercise
	Tendin* AND exercise
	Tendon AND exercise
Clinicialtrials.gov	Tendinopathy AND exercise
	Tendin* AND exercise
	Tendon AND exercise
ISRCTN	Tendinopathy AND exercise
	Tendin* AND exercise
	Tendon AND exercise
EU CTR	Tendinopathy AND exercise
	Tendin* AND exercise
	Tendon AND exercise
ANZCTR	Tendinopathy AND exercise
	Tendin* AND exercise
	Tendon AND exercise

ISRCTN – the Research Registry; EU CTN – European Clinical Trials Registry; ANZCTR – Australia and New Zealand Clinical Trials Registry

Supplementary File 6: Extraction codebook

	Column	Heading	Description				
	Olulliii	Heading	Description				
	A	Initials Reviewer	Identification of individual extracting information				
	В	Covidence	Reference number for Covidence				
	Ъ	Identifier					
	C	Author	First author surname et al.,				
	D	Year	Year of publication				
	E	Title	Study title				
	F	Country	Country where study was conducted				
	G	Journal	Journal name				
	Н	Aims/Purpose	Study aims/purpose				
	I	Tendinopathy	1=Achilles; 2= Lateral elbow (tennis); 3 = Patellar; 4 =				
	т	type	Rotator cuff (SI)				
	J K	Study Design	RCT = 1; Quasi-experimental = 2				
	L L	Age Mean Age SD	Mean age of study sample as a whole				
	1.1	Baseline Total N	Standard deviation age of study sample as a whole Total sample across all interventions measured at baseline				
150	2 1V1	Training Status	Brief description of training status of study sample as a				
4	N	Description	whole				
Study Details	-	Training Status	1 = Performance; 2 = Sporting; 3 = Other				
É	j O	Code	1 – 1 chormanee, 2 – Sporting, 3 – Other				
J.	5 P	Sex	Percentage female of study sample as a whole				
	Q	BMI Mean	Mean BMI of study sample as a whole				
	R	BMI SD	Standard deviation of BMI of study sample as a whole				
	C	Symptom	Mean severity measure at baseline of study sample as a				
	S	Severity Mean	whole				
	Т	Symptom	Standard deviation of severity measure at baseline of study				
	1	Severity SD	sample as a whole				
		Symptom	Mean symptom duration reported in months				
	U	Duration Mean					
		(Months)					
		Symptom	Standard deviation symptom duration reported in months				
	V	Duration SD					
		(Months)					
	W	Population	Any additional information relevant to the participants				
		Comments	investigated including diagnostic criteria				
	X	Outcome	1 = Disability; 2 = Pain on loading/activity; 3 = Pain over a				
	Λ	Category	specified time; 4 = Pain without further specification; 5 = Physical function capacity				
ý	3	Outcome Tool	Description of outcome tool				
Ĕ	Y	Outcome 1001	Description of outcome tool				
Ontcomes	3	Reflection	1 = Increase in outcome indicates positive treatment; -1 =				
ē	\mathbf{z}	Reflection	Decrease in outcome indicates positive treatment				
		Magazamamamt					
	A	Measurement	Time of measurement in weeks				
	A	Time (Weeks)					
<u>.</u>	=	Dominant	Only one dominant theme to be selected				
Inter	5 AB	Treatment Class	1 = Resistance; 2 = Plyometric; 3 = Vibration; 4 =				
			Flexibility; 5 = Movement pattern retraining				

	AC	Total Treatment class	Multiple themes to be selected as required 1 = Resistance; 2 = Plyometric; 3 = Vibration; 4 =
	AC	Class	Flexibility; 5 = Movement pattern retraining
	A D	Intervention N	Intervention sample size at specified time
	AE Intervention Total Duration		Total duration of exercise intervention in weeks
	AF	Intervention Adherence %	Reporting of adherence to exercise (reported as a percentage) if applicable
	A G	Intervention Location	Location exercise was performed 1 = Home; 2 = Clinic; 3 = Fitness facility; 4 = NR; 5 = NA
	A H	Intervention Volume	Numerical value describing volume
	AI	Intervention Volume	1 = Duration of session (mins); 2 = sets * repetitions; 3 = number of repetitions; 4 = number of sets
	AJ	Category Intervention Volume	Any additional information relevant.
	A K	Comments Intervention Intensity	Numerical value describing intensity
	AL	Intervention Intensity	1 = Absolute; 2 = Relative
	A	Category Intervention	Number of sessions per week. Where there is progression,
	M	Frequency	average value is to be entered.
	A N	Intervention Frequency Comments	Any additional information relevant.
		Intervention Progression	Multiple themes to be selected as required 1 = No progression; 2 = NR; 3 = Progression volume; 4 =
	A O		Progression intensity; 5 = Progression frequency; 6 = Progression specificity; 7 = Progression capacity; 8 =
		Intervention	Other Any additional information relevant.
	AP	Progression Comments	
7	A	Control	1 = Placebo; 2 = No treatment
Contro	Q	Comparator Control	Any additional information relevant.
ටී	AR	Comparator Comments	·
	AS	Intervention Baseline Mean	Baseline mean for exercise therapy
Ę	AT	Intervention Baseline SD	Baseline standard deviation for exercise therapy
Data	A U	Intervention Measurement	Mean of outcome for exercise therapy at stated time point
	A	Mean Intervention	Standard deviation of outcome for exercise therapy at

V	Measurement SD	stated time point
A	Control Baseline	Baseline mean for control
W	Mean	
A	Control Baseline	Baseline standard deviation for control
X	SD	
A	Control	Mean of outcome for control at stated time point
Y	Measurement	
1	Mean	
AZ	Control	Standard deviation of outcome for control at stated time
AL	Measurement SD	point
	Measurement	State if a different value has been entered for means (e.g.
	Comments	median), a different value for standard deviations (e.g.
BA		standard error, IQR, percentiles, distance from mean to
bА		upper bound). Provide the relevant statistic (width of CI's,
		width of percentiles). Also state if data has been extracted
		by digitization
* O4	C	

^{*} Outcome Specific

Supplementary file 8: Analysis R Code

Variables:

- 1) NT: Number of treatment arms
- 2) ID: Study ID according to included list
- 3) Study Type: Binary RCT or quasi-experimental
- 4) Tendinopathy. Type: Category variable of the tendinopathy locations
- 5) Outcome. Domain: Category variable of the outcome domain
- 6) Outcome. Tool: Description of the outcome measure fitting within the outcome domain
- 7) ExerciseHierarchy: Binary classification of exercise therapy as exercise only or multiple exercise types
- 8) ClassD: Category variable of dominant exercise class in therapy
- 9) ClassAll: Category variable of all exercise classes in therapy
- 10) TreatmentD: Category variable of dominant exercise treatment in therapy
- 11) Treatment All: Category variable of all exercise treatments in therapy
- 12) N: Number of participants data are collected from
- 13) AgeMean: Mean age of participants
- 14) AgeSD: Standard deviation of age of participants
- 15) BMIMean: Mean BMI of participants
- 16) BMISD: Standard deviation of BMI of participants
- 17) Supervision: Category variable of whether exercise therapy was supervised or unsupervised
- 18) Location: Category variable of location exercise therapy was performed
- 19) Random.sequence.generation: Risk of bias variable
- 20) Allocation.concealment: Risk of bias variable
- 21) Blinding.of.participants.personnel: Risk of bias variable
- 22) Blinding.of.outcome.assessment: Risk of bias variable
- 23) Incomplete.outcome.bias: Risk of bias variable
- 24) Selective.reporting: Risk of bias variable
- 25) Other.bias: Risk of bias variable
- 26) Time: Time in weeks of measurement from baseline
- 27) TimeC: Category variable identifying time in weeks of measurement from baseline
- 28) OutcomeLevel: Variable for nesting structure at outcome level
- 29) MesureLevel: Variable for nesting structure at measurement level
- 30) ES: Standardised mean difference effect size
- 31) SEES: Standard error of standardised mean difference effect size

NA: refers to missing data

Four level model with t-distribution: (All Outcomes)

```
\label{eq:modoALL.prior} $$\operatorname{get\_prior}(ES \mid se(SEES,sigma=TRUE) \sim 1 + (1 \mid ID/OutcomeLevel/MeasureLevel), family = student(), data=Data)$$ $\operatorname{ModoALL.prior\$prior}[10] = "student_t(3, 0, 1.5)" $$ $\operatorname{set.seed}(123)$$ $\operatorname{modoALL} = \operatorname{brm}(ES \mid se(SEES,sigma=TRUE) \sim 1 + (1 \mid ID/OutcomeLevel/MeasureLevel), family = student(),
```

```
data = Data, prior = Mod0ALL.prior, chains = 4, iter = 20000, warmup = 10000)
mod0ALLPS= posterior_samples(mod0ALL)
# Quantile function
QuantileFunction = function(data,Model,PosteriorDF){
Preds = data[c('ES', 'SEES', 'ID', 'OutcomeLevel', 'MeasureLevel')] %>%
  add_predicted_draws(Model)
 PredsDF = as.data.frame(Preds)
 PredsL = length(data[,1])
 PredsFitL = length(PosteriorDF[,1])
 PredsDF2 = matrix(data = NA, nrow=PredsL,ncol=PredsFitL+1)
 PredsDF2[,1] =
  PredsDF[(seq(1,PredsL*PredsFitL,
         PredsFitL)),6]
 for(i in 1:PredsFitL){
  PredsDF2[,(i+1)] =
   PredsDF[(seq(1,PredsL*PredsFitL,
          PredsFitL)+(i-1),10
 }
 q025 = c(NULL)
 for(i in 1:PredsFitL){
  q025[i] = quantile(PredsDF2[,i+1],0.25)[[1]]
 q025q = quantile(q025,c(0.025,0.5,0.975))
 q05 = c(NULL)
 for(i in 1:PredsFitL){
  q05[i] = quantile(PredsDF2[,i+1],0.5)[[1]]
 q05q = quantile(q05,c(0.025,0.5,0.975))
 q075 = c(NULL)
 for(i in 1:PredsFitL){
  q075[i] = quantile(PredsDF2[,i+1],0.75)[[1]]
 q075q = quantile(q075,c(0.025,0.5,0.975))
return(rbind(q025q,q05q,q075q))
mod0ALLQ = QuantileFunction(Data, mod0ALL, mod0ALLPS)
# Repeat for different Outcome
```

```
\text{mod}0\underline{\text{Outcome}} = \text{brm(ES } | \text{se(SEES,sigma=TRUE)} \sim 1 + (1 | \text{ID/OutcomeLevel/MeasureLevel}),
family = student(),
      data = Data[Data$Outcome.Domain=="<u>Outcome</u>",], prior = mod0<u>Outcome</u>.prior, chains = 4,
iter = 20000, warmup = 10000)
mod0<u>Outcome</u>PS= posterior_samples(mod0<u>Outcome</u>)
mod<u>Outcome</u>Q = QuantileFunction(Data[Data$Outcome.Domain=="<u>Outcome</u>",], mod0<u>Outcome</u>,
mod00utcomePS)
# Plot
ggplot(Data,
   aes(x=ES)) + geom_density(adjust = 2) +
annotate("text",x=-0.68, y=0, label="All Outcomes", size=3) +
coord\_cartesian(xlim = c(-1, 3.5), ylim = c(0, 0.7), expand = TRUE) +
theme_classic() + theme(axis.line.y = element_blank()) + ylab("") +
theme(axis.text.y = element_blank()) + theme(axis.ticks.y = element_blank()) +
scale_x_continuous("Effect Size", breaks = seq(-1,3.5,0.5)) +
theme(axis.text.x = element_text(size=8)) + theme(axis.title.x = element_text(size=9)) +
annotate("pointrange", y=0, x =mod0N4ALLQ[[1,2]],xmin = mod0N4ALLQ[[1,1]], xmax =
mod0N4ALLQ[[1,3]],colour = "black", size = 0.6, shape = 18) +
annotate("pointrange", y=0, x=mod0N4ALLQ[[2,2]],xmin = mod0N4ALLQ[[2,1]], xmax =
mod0N4ALLQ[[2,3]],colour = "black", size = 0.6, shape = 18) +
annotate("pointrange", y=0, x = mod0N4ALLQ[[3,2]], xmin = mod0N4ALLQ[[3,1]], xmax = mod0N4ALLQ[[3,1]]
mod0N4ALLQ[[3,3]],colour = "black", size = 0.6, shape = 18) +
annotate("text",x=-0.6, y=0.5, label="Disability", size=3) +
annotate("pointrange", y=0.5, x =mod0N4DISQ[[1,2]],xmin = mod0N4DISQ[[1,1]], xmax =
mod0N4DISQ[[1,3]],colour = "red", size = 0.4) +
annotate("pointrange", y=0.5, x =mod0N4DISQ[[2,2]],xmin = mod0N4DISQ[[2,1]], xmax =
mod0N4DISQ[[2,3]], colour = "red", size = 0.4) +
annotate("pointrange", y=0.5, x =mod0N4DISQ[[3,2]],xmin = mod0N4DISQ[[3,1]], xmax =
mod0N4DISQ[[3,3]],colour = "red", size = 0.4) +
annotate("text",x=-0.6, y=0.6, label="Function", size=3) +
annotate("pointrange", y=0.6, x = mod0N4FUNQ[[1,2]], xmin = mod0N4FUNQ[[1,1]], xmax = mod0N4FUNQ[[1,1]]
mod0N4FUNQ[[1,3]],colour = "red", size = 0.4) +
 annotate("pointrange", y=0.6, x =mod0N4FUNQ[[2,2]],xmin = mod0N4FUNQ[[2,1]], xmax =
mod0N4FUNQ[[2,3]],colour = "red", size = 0.4) +
 annotate("pointrange", y=0.6, x = mod0N4FUNQ[[3,2]], xmin = mod0N4FUNQ[[3,1]], xmax = mod0N4FUNQ[[3,1]]
mod0N4FUNQ[[3,3]],colour = "red", size = 0.4) +
annotate("text",x=-0.6, y=0.4, label="Pain", size=3) +
 annotate("pointrange", y=0.4, x = mod0N4PAINQ[[1,2]], xmin = mod0N4PAINQ[[1,1]], xmax = mod0N4PAINQ[[1,2]]
mod0N4PAINQ[[1,3]],colour = "red", size = 0.4) +
 annotate("pointrange", y=0.4, x = mod0N4PAINQ[[2,2]], xmin = mod0N4PAINQ[[2,1]], xmax = mod0N4PAINQ[[2,1]]
mod0N4PAINQ[[2,3]], colour = "red", size = 0.4) +
 annotate("pointrange", y=0.4, x =mod0N4PAINQ[[3,2]],xmin = mod0N4PAINQ[[3,1]], xmax =
mod0N4PAINQ[[3,3]],colour = "red", size = 0.4) +
annotate("text",x=-0.6, y=0.2, label="PFC", size=3) +
annotate("pointrange", y=0.2, x =mod0N4PFCQ[[1,2]],xmin = mod0N4PFCQ[[1,1]], xmax =
mod0N4PFCQ[[1,3]],colour = "red", size = 0.4) +
```

```
annotate("pointrange", y=0.2, x =mod0N4PFCQ[[2,2]],xmin = mod0N4PFCQ[[2,1]], xmax =
mod0N4PFCQ[[2,3]],colour = "red", size = 0.4) +
  annotate("pointrange", y=0.2, x=mod0N4PFCQ[[3,2]],xmin = mod0N4PFCQ[[3,1]], xmax =
mod0N4PFCQ[[3,3]],colour = "red", size = 0.4) +
  annotate("text",x=-0.6, y=0.1, label="QoL", size=3) +
  annotate("pointrange", y=0.1, x=mod0N4QOLQ[[1,2]],xmin = mod0N4QOLQ[[1,1]], xmax =
mod0N4QOLQ[[1,3]],colour = "red", size = 0.4) +
  annotate("pointrange", y=0.1, x=mod0N4QOLQ[[2,2]],xmin = mod0N4QOLQ[[2,1]], xmax =
mod0N4QOLQ[[2,3]], colour = "red", size = 0.4) +
  annotate("pointrange", y=0.1, x =mod0N4QOLQ[[3,2]],xmin = mod0N4QOLQ[[3,1]], xmax =
mod0N4QOLQ[[3,3]],colour = "red", size = 0.4) +
  annotate("text",x=-0.6, y=0.3, label="ROM", size=3) +
  annotate("pointrange", y=0.3, x=mod0N4ROMQ[[1,2]], xmin=mod0N4ROMQ[[1,1]], xmax=mod0N4ROMQ[[1,1]], xmax=mod0N4ROMQ[[1,2]], xmin=mod0N4ROMQ[[1,1]], xmax=mod0N4ROMQ[[1,1]], x
mod0N4ROMQ[[1,3]],colour = "red", size = 0.4) +
  annotate("pointrange", y=0.3, x =mod0N4ROMQ[[2,2]],xmin = mod0N4ROMQ[[2,1]], xmax =
mod0N4ROMQ[[2,3]],colour = "red", size = 0.4) +
  annotate("pointrange", y=0.3, x =mod0N4ROMQ[[3,2]],xmin = mod0N4ROMQ[[3,1]], xmax =
mod0N4ROMQ[[3,3]],colour = "red", size = 0.4)
```

Supplementary File 9: Table of included studies and reference list

Study (first author, year, country)	Design	Tendinopathy Location	(number (n); sex (%female); mean (sd) age; mean (sd) symptom duration in months)	Exercise Treatment arms	Exercise Treatment classes	Findings
Aceituno- Gómez 2019 Spain ¹	Quasi- experimental	Rotator cuff - subacromial impingement	N=43 % female 60.9 Age 59 (8.9) Symptoms NR Training status Other	1	(Flexibility, Resistance)	High-intensity laser therapy plus exercise did not give greater improvements in pain and functionality in patients with subacromial syndrome than exercise alone.
Akkaya 2016 Turkey ²	RCT	Rotator cuff - subacromial impingement	N=34 % female 67.6 Age 41.7 (8.9) Symptoms 6.9 (4.1) Training status Other	2	2*(Flexibility)	Weighted and un-weighted solo pendulum exercises achieved significant clinical improvements but showed no differences in ultrasonographic acrohumeral distance measurements between groups.
Alfredson 1998 Sweden ³	Quasi- experimental	Achilles	N= 30 % female 20.0 Age 44.0 (7.0) Symptoms 25.9 (3- 100)** Training status Recreational	1	(Resistance)	Our treatment model with heavy-load eccentric calf muscle training has a very good short-term effect on athletes in their early forties.
Alfredson 1999 Sweden ⁴	Quasi- experimental	Achilles	N= 24 % female 14.3 Age 42.6 (9.0) Symptoms 23.7 (3- 100)** Training status Recreational	1	(Resistance)	Heavy-loaded, eccentric calf-muscle training seems to be a good treatment mode for chronic Achilles tendinosis.
Arias-Buría 2017 Spain ⁵	RCT	Rotator cuff - subacromial impingement	N= 50 % female 26.0 Age 48.5 (5.5) Symptoms 71.9 (21.6) Training status Other	1	(Resistance)	This study found that the inclusion of 2 sessions of TrP-DN into an exercise program was effective for improving shoulder pain-related disability at short-, medium-, and long-term; however, no greater improvement in shoulder pain was observed.
AriasBuría 2015 Spain ⁶	RCT	Rotator cuff - subacromial impingement	N= 36 % female 75.0	1	(Resistance)	Ultrasound-guided percutaneous electrolysis combined with eccentric

			Age 57.5 (6.4) Symptoms 10.9 (2.6) Training status Other			exercises resulted in better short-term outcomes compared to eccentric exercises alone.
Bae 2011 Korea (Republic of) ⁷	Quasi- experimental	Rotator cuff - subacromial impingement	N=35 % female 65.7 Age 49.1 (4.9) Symptoms Training status Other	1	(Proprioception, Resistance)	The motor control and strengthening programme improved pain, function, strength and ROM.
Bahr 2006 Norway ⁸	RCT	Patellar	N= 40 % female 12.5 Age 30.5 (7.9) Symptoms 34 (28.7) Training status Other	1	(Resistance)	No added benefit was observed for surgical treatment to eccentric strength training. Eccentric training should be offered for 12 weeks before tenotomy is considered for the treatment of patellar tendinopathy.
Balius 2016 Spain ⁹	RCT	Achilles	N=37 % female 20.4 Age 41.4 (11.7) Symptoms NR Training status Other	6	4*(Resistance);2*(Flexibility)	Findings confirmed the therapeutic potential of eccentric exercise at reactive and degenerative stages of tendinopathy. MCVC supplementation decreased pain more than eccentric exercise alone (reactive tendinopathy) Personalized stretching regime supplemented with MCVC may be appropriate for some patients
Bang 2000 United States ¹⁰	RCT	Rotator cuff - subacromial impingement	N=50 % female 42.3 Age 43 (9.1) Symptoms 5 (3.3) Training status Other	1	(Flexibility, Resistance)	Manual physical therapy applied by experienced physical therapists combined with supervised exercise in a brief clinical trial is better than exercise alone for increasing strength, decreasing pain, and improving function in patients with shoulder impingement syndrome
Başkurt 2011 Turkey ¹¹	Quasi- experimental	Rotator cuff - subacromial impingement	N= 40 % female 67.5 Age 51.4 (10.0) Symptoms NR Training status Other	2	1*(Flexibility, Resistance);1*(Flexibility, Proprioception)	Scapular stabilisation combine with stretching and strengthening exercises can be more effective in the short-term for SIS.
Beyer 2015 Denmark ¹²	RCT	Achilles	N= 58 % female 31.9 Age 48.0 (2.0) Symptoms 18.1 (4.3) Training	2	2*(Resistance)	Both traditional eccentric exercise and HSR yield positive, equally good and lasting clinical results in patients with Achilles tendinopathy. HSR is associated with greater patient satisfaction after 12

			status Other			weeks but not after 52 weeks.
Blume 2015 United States ¹³	RCT	Rotator cuff - subacromial impingement	N= 34 % female 58.0 Age 49.4 (15.6) Symptoms 22.7 (24.3) Training status Other	2	2*(Flexibility, Resistance)	Both eccentric and concentric PRE programs resulted in improved function, AROM, and strength in patients with SAIS. However, no difference was found between the two exercise modes, suggesting that therapists may use exercises that utilize either exercise mode in their treatment of SAIS.
Boudreau 2019 Canada ¹⁴	RCT	Rotator cuff - subacromial impingement	N= 42 % female 52.4 Age 42.9 (12.0) Symptoms 43.0 (46.6) Training status Other	2	2*(Resistance)	No additional benefit was found to adding coactivation to regular rotator cuff strengthening exercises at 6-weeks.
Breda 2020 Netherlands ¹⁵	RCT	Patellar	N=76 % female 23.7 Age 24 (3.9) Symptoms 98.5 (NR) Training status Performance	2	1*(Plyometric, Resistance);1*(Flexibility, Resistance)	In patients with patellar tendinopathy, progressive tendon-loading exercises resulted in a significantly better clinical outcome after 24 weeks than eccentric exercise therapy. Progressive tendon-loading exercises are superior to eccentric exercise therapy and are therefore recommended as initial conservative treatment for patellar tendinopathy.
Brox 1999 Norway ¹⁶	RCT	Rotator cuff - subacromial impingement	N= 125 % female 44.0 Age 47.6 (23-66)** Symptoms NR Training status Other	1	(Proprioception, Resistance)	At 2.5 years follow-up, both arthroscopic surgery and supervised exercises are better treatments than placebo with no significant difference between the 2 active treatments.
Calis 2011 Turkey ¹⁷	RCT	Rotator cuff - subacromial impingement	N= 52 % female 67.3 Age 49.2 (12.6) Symptoms 3.0 (1-24)** Training status Other	1	(Flexibility)	Ultrasound and laser treatments were not superior to each other in the treatment of SIS
Chaconas 2017 United States ¹⁸	RCT	Rotator cuff - subacromial impingement	N=46 % female 41.7 Age 45.9 (17.4) Symptoms 49.1 (80) Training status Other	2	2*(Flexibility, Resistance)	An eccentric program targeting the external rotators was superior to a general exercise program for strength, pain, and function after six months. The findings suggest eccentric training may be efficacious to improve self-

						report function and strength for those with SAPS.
Cheng 2007 Hong Kong, China (SAR) ¹⁹	RCT	Rotator cuff - subacromial impingement	N=94 % female Age 32.4 (10.2) Symptoms 23.4 Training status Other	2	1*(Flexibility, Proprioception);1*(Resistance, Flexibility)	An eccentric program targeting the external rotators was superior to a general exercise program for strength, pain, and function after six months. The findings suggest eccentric training may be efficacious to improve self-report function and strength for those with subacromial pain syndrome.
Cho 2017 Korea (Republic of) ²⁰	Quasi- experimental	Patellar	N= 30 % female 46.7 Age 33.1 (29.1) Symptoms 15.1 (16.1) Training status Other	1	(Flexibility, Proprioception, Resistance)	A rehabilitation exercise programme was more effective at improving pain, strength and function in patellar tendinopathy that injection therapy alone.
de Jonge 2008 Netherlands ²¹	RCT	Achilles	N= 70 % female NR Age 44.6 (26-59) ** Symptoms 30.7 (2-204) ** Training status Other	1	(Resistance)	Eccentric exercises with or without a night splint improved functional outcome at one year follow-up. At follow-up there was no significant difference in clinical outcome when a night splint was used in addition to an eccentric exercise.
de Oliveira 2020 Canada ²²	RCT	Rotator cuff - subacromial impingement	N= 52 % female 42.3 Age 30.2 (8.3) Symptoms 22.6 (26.7) Training status Other	1	(Flexibility, Proprioception, Resistance)	Whereas symptoms, functional limitations, ROM, and AHD improved in both groups, the addition of KT did not lead to superior outcomes compared with exercise-based treatment alone, in the mid and long term, for individuals with RCRSP.
de Vos 2007 Netherlands ²³	RCT	Achilles	N= 63 % female 41.3 Age 44.6 (8) Symptoms 30.6 (50.6) Training status Recreational	1	(Resistance)	A night splint has no added benefit to eccentric exercises in the treatment of chronic midportion Achilles tendinopathy. There was no significant difference between the two groups in VISA-A score and patient satisfaction.
Dejaco 2017 Netherlands ²⁴	RCT	Rotator cuff - subacromial impingement	N=36 % female 47.3 Age 49.5 (11.3) Symptoms 19.7 (20.1) Training status Other	2	2*(Flexibility, Resistance)	12-week-isolated eccentric training programme of the RC is beneficial for shoulder function and pain after 26 weeks in patients with RC tendinopathy. However, it is no more beneficial than a conventional exercise programme for the RC and scapular muscles.
Devereaux 2016 Canada ²⁵	RCT	Rotator cuff - subacromial impingement	N= 100 % female 37.9	2	1*(Flexibility, Proprioception);1*(Resistance, Flexibility)	The improvements in pain and function observed with an NSAID or precut

			Age 48.0 (11.9) Symptoms NR Training status Other			kinesiology tape as adjuvant treatments were no greater than with rehabilitation exercise alone.
Dimitrios 2013 Cyprus ²⁶	Quasi- experimental	Patellar	N= 60 % female 36.7 Age 47.57 (5.9) Symptoms 4.5 (NR) Training status Other	2	2*(Flexibility, Resistance)	A specific supervised exercise programme is superior to a specific home exercise programme in reducing pain and improving function in patients with LET at the end of the treatment and at the 3 month follow-up.
Dimitrios 2012 Greece ²⁷	Quasi- experimental	Patellar	N= 60 % female 36.7 Age 47.6 (5.9) Symptoms 4.5 (NR) Training status Other	2	1*(Flexibility, Resistance);1*(Resistance)	Eccentric training and static stretching exercises is superior to eccentric training alone to reduce pain and improve function in patients with patellar tendinopathy at the end of the treatment and at follow-up.
Dupuis 2018 Canada ²⁸	RCT	Rotator cuff - subacromial impingement	N=43 % female 55.8 Age 33.3 (11.7) Symptoms 0.9 (0.3) Training status Other	2	1*(Flexibility);1*(Flexibility, Resistance)	Both groups showed statistically significant improvements on symptoms and function at 2 weeks and 6 weeks but there was no difference between the short-term effect of cryotherapy and a gradual reloading exercise programme.
Engebretsen 2009 Norway ²⁹	RCT	Rotator cuff - subacromial impingement	N= 104 % female 50.0 Age 48.0 (10.6) Symptoms 12.5 (NR) Training status Other	1	(Plyometric, Proprioception, Resistance)	Supervised exercises are superior to ESWT in terms of shoulder pain, disability and some work-related outcomes.
Engebretsen 2011 Norway ³⁰	RCT	Rotator cuff - subacromial impingement	N= 104 % female 50.0 Age 48.0 (10.6) Symptoms 12.5 (NR) Training status Other	1	(Proprioception)	Both radial ESWT and the supervised exercise regime devised by Bohmer (1998) provided similar benefits in pain and function-related outcomes. However, exercise may be superior for work-related outcomes.
Gatz 2020 Germany ³¹	RCT	Achilles	N= 42 % female 35.7 Age 50.0 (12.0) Symptoms 27.5 (23.8) Training status Other	2	2*(Resistance)	No additional clinical benefits of adding ISOs to a basic EE program could be found in this preliminary randomized controlled trial study over a period of 3 months. SWE was able to differentiate between insertional and midportion tendon tissue and localize reported symptoms to sublocations but this did not correlate with better clinical scores (VISA-A) over a 3-

Giray 2019 Turkey ³²	RCT	Lateral elbow/tennis elbow	N= 30 % female 86.7 Age 44.46 (9.92) Symptoms 1.69 (NR) Training status Other	1	(Flexibility, Resistance)	month follow-up period. Kinesiotaping in addition to exercises is more effective than sham taping and exercises alone in improving pain in daily activities and arm disability due to lateral epicondylitis.
Granviken 2015 Norway ³³	RCT	Rotator cuff - subacromial impingement	N=44 % female 48 Age 47.9 (9.9) Symptoms 14.5 Training status Other	2	1*(Flexibility, Proprioception);1*(Resistance, Flexibility)	No significant differences in pain and disability were found between home exercises and supervised exercises of more than the first session of a 6-week exercise regime for people with subacromial impingement.
Hallgren 2014 Sweden ³⁴	RCT	Rotator cuff - subacromial impingement	N= 50 % female 37.0 Age 52 (30-65)** Symptoms 18 (6-186)* Training status Other	2	1*(Resistance);1*(Flexibility)	Specific exercises produced positive short-term improvements at 1-year follow-up and reduces the need for surgery. Full-thickness tear and a low CMS score appear to be predictors of poor outcome.
Hallgren 2017 Sweden ³⁵	RCT	Rotator cuff - subacromial impingement	N= 108 % female 34.1 Age 58 (NR) Symptoms NR Training status Other	2	1*(Flexibility, Resistance);1*(Flexibility)	More patients in the specific exercise group managed to avoid surgery compared to the unspecific exercise group at 5-year follow-up supporting it's prescription as an initial treatment for patients with subacromial pain.
Heron 2017 United Kingdom ³⁶	RCT	Rotator cuff - subacromial impingement	N= 120 % female 41.0 Age 49.9 (NR) Symptoms NR Training status Other	3	2*(Flexibility, Resistance);1*(Flexibility)	Open chain, closed chain, and range of movement exercises all seem to be effective in bringing about short-term changes in pain and disability in patients with rotator cuff tendinopathy.
Hotta 2020 Brazil ³⁷	RCT	Rotator cuff - subacromial impingement	N=60 % female 70 Age 49 (9) Symptoms 28.5 (24) Training status Other	2	1*(Resistance, Proprioception);1*(Resistance)	The inclusion of the isolated scapular stabilization exercises, emphasizing retraction and depression of the scapula, to a progressive general periscapular strengthening protocol did not add benefits to self-reported shoulder pain and disability, muscle strength, and ROM in patients with subacromial pain syndrome.
Johansson 2005 Sweden ³⁸	RCT	Rotator cuff - subacromial impingement	N=85 % female 69.4 Age 49 (7.5) Symptoms NR Training	1	(Flexibility, Resistance)	Acupuncture was more effective than ultrasound when applied in addition to home exercises.

			status Other			
Johnson 2005 Sweden ³⁹	RCT	Patellar	N=15 % female 13.3 Age 24.9 (8.2) Symptoms 17.5 (13.2) Training status Performance	2	2*(Resistance)	Eccentric, but not concentric, quadriceps training on a decline board, seems to reduce pain in jumper's knee.
Jonsson 2009 Sweden ⁴⁰	Quasi- experimental	Achilles	N= 15 % female 13.3 Age 25.0 (NR) Symptoms 17.5 (13.2) Training status Other	2	2*(Resistance)	Treatment with painful eccentric calf-muscle training showed good clinical results based on VAS scores, patient satisfaction, and return to pre-injury activity levels in patients with chronic painful mid-portion Achilles tendinosis, but not in patients with chronic insertional Achilles tendon pain.
Juul- Kristensen 2019 Denmark ⁴¹	RCT	Rotator cuff - subacromial impingement	N= 58 % female 51 Age 42.9 (12.4) Symptoms NR Training status Other	2	2*(Proprioception, Flexibility)	Electromyography- biofeedback neuromuscular shoulder exercises and neuromuscular shoulder exercises were both effective in reducing pain to a clinically relevant level, while electromyography biofeedback did not make a difference. The current neuromuscular shoulder exercise protocol is recommended
Ketola 2009 Finland ⁴²	RCT	Rotator cuff - subacromial impingement	N=134 % female 62.9 Age 47.1(23.3- 60.0)** Symptoms 2.6 (NR) Training status Other	1	(Resistance, Proprioception)	Arthroscopic acromioplasty provides no clinically important effects over a structured and supervised exercise programme alone in terms of subjective outcome or costeffectiveness when measured at 24 months.
Ketola 2013 Finland ⁴³	RCT	Rotator cuff - subacromial impingement	N=140 % female 62.9 Age 41.7 Symptoms Training status Other	1	(Flexibility, Resistance)	Differences in the patient- centred primary and secondary parameters between the two treatment groups were not statistically significant, suggesting that acromioplasty is not cost- effective.
Kim 2017 Korea (Republic of) ⁴⁴	RCT	Rotator cuff - subacromial impingement	N= 40 % female 72.5 Age 51.1 (10.6) Symptoms NR Training status Other	2	2*(Proprioception))	The use of visual feedback and 3D motion images can improve pain and function in SIS.
Kim	RCT	Rotator cuff -	N= 40	1	(Proprioception, Vibration)	Both the Neurac modality

2020 Korea (Republic of) ⁴⁵		subacromial impingement	% female 100 Age 46.2 (4.6) Symptoms NR Training status Other			and manual therapy induced pain relief, improved function, and increased ROM. The Neurac intervention also resulted in a significant enhancement of shoulder muscle strength indicating its superiority as an effective therapeutic modality for this particular
Knobloch 2008 Italy ⁴⁶	RCT	Achilles	N= 92 % female 35.0 Age 47.5 (11.0) Symptoms NR Training status Recreational	1	(Resistance)	patient group. Patients with tendinopathy of the main body of the AT experienced improved clinical outcome with both management options. Although tendon microcirculation was optimized in the combined group of eccentric training and AirHeel Brace, these micro-vascular advantages do not translate into superior clinical performance when compared with eccentric training alone.
Knobloch 2007 Germany ⁴⁷	RCT	Achilles	N= 20 % female 45.0 Age 32.5 (11.0) Symptoms NR Training status	1	(Resistance)	training alone. An eccentric-training program performed daily over 12 weeks reduced the increased paratendinous capillary blood flow in Achilles tendinopathy by as much as 45% and decreased pain level based on a visual analog scale. Local paratendon oxygenation was preserved while paratendinous postcapillary venous filling pressures were reduced after 12 weeks of eccentric training, which appears to be beneficial from the perspective of microcirculation.
Knobloch 2007 Germany ⁴⁸	RCT	Achilles	N= 118 % female 40 Age 48.5 (12) Symptoms NR Training status Other	1	(Resistance)	Achilles tendon oxygen saturation is increased, and capillary venous clearance facilitated using an Achilles wrap in addition to daily 12-week eccentric training
Kongsgaard 2009 Denmark ⁴⁹	RCT	Patellar	N= 37 % female 0 Age 32.4 (8.8) Symptoms 18.7 (12.3) Training status Recreational	2	2*(Resistance)	Corticosteroid injection has good short-term but poor long-term clinical effects, in patellar tendinopathy. Heavy-slow resistance exercise has good short-and long-term clinical effects accompanied by pathology improvement and increased collagen turnover.
Kromer 2014	RCT	Rotator cuff - subacromial	N= 90 % female	1	(Flexibility, Proprioception, Resistance)	The use of MT including Physiotherapy provides no

Germany ⁵⁰		impingement	51.1 Age 51.8 (11.2) Symptoms 24.1 (35.1) Training status Other			additional benefits and is more expensive in comparison to exercise only interventions.
Kromer 2013 Germany ⁵¹	RCT	Rotator cuff - subacromial impingement	N= 90 % female 51.1 Age 51.8 (11.2) Symptoms 7.8 (9.8) Training status Other	1	(Flexibility, Proprioception, Resistance)	Individually adapted exercises were effective in the treatment of patients with shoulder impingement syndrome. Individualized manual Physiotherapy contributed only a minor amount to the improvement in pain intensity.
Littlewood 2016 United Kingdom ⁵²	RCT	Rotator cuff - subacromial impingement	N= 60 % female 50.3 Age 54.7 (NR) Symptoms 14.6 (NR) Training status Other	1	(Resistance)	Self-management programme based on a single exercise were comparable to usual Physiotherapy in the short-, mid- and long-term.
Ludewig 2003 United States ⁵³	RCT	Rotator cuff - subacromial impingement	N= 85 % female 0.0 Age 48.8 (2.1) Symptoms NR Training status Other	1	(Flexibility, Resistance)	Home exercise programme are more effective in reducing symptoms and improving function (Shoulder Rating Questionnaire, shoulder satisfaction score) than the control group in construction workers with shoulder pain.
Luginbuhl 2008 Switzerland ⁵⁴	RCT	Lateral elbow/tennis elbow	N= 30 % female 72.7 Age 47 (9) Symptoms 10 (11) Training status Other	1	(Resistance)	No beneficial effect of neither the forearm support band nor the strengthening exercises could be found.
Maenhout 2013 Belgium ⁵⁵	RCT	Rotator cuff - subacromial impingement	N= 61 % female 59.0 Age 39.8 (13.0) Symptoms NR Training status Other	2	2*(Resistance)	Adding heavy load eccentric training resulted in a higher gain in isometric strength at 90 degree of scapular abduction but was not superior for decreasing pain and improving shoulder function. The addition of a limited amount of Physiotherapy sessions combined with a daily home exercise programme is highly effective in patients with impingement.
Mafi 2001 Sweden ⁵⁶	RCT	Achilles	N= 44 % female 45.5 Age 48.3 (8.8) Symptoms 20.5 (3-	2	2*(Resistance)	Eccentric calf muscle training showed superior results to concentric training in the treatment of chronic Achilles tendinosis based on patient satisfaction and return to activity level.

			120)** Training status Other			
Manias 2006 United Kingdom ⁵⁷	RCT	Lateral elbow/tennis elbow	N= 40 % female 67.5 Age 42.86 (6.23) Symptoms NR Training status Other	2	2*(Resistance)	An exercise programme consisting of eccentric and static stretching exercises had reduced the pain in patients with lateral epicondyle tendinopathy at the end of the treatment and at the follow up whether or not ice was included.
Martinez- Silvestrini 2005 United States ⁵⁸	Quasi- experimental	Lateral elbow/tennis elbow	N= 81 % female 46.8 Age 45.5 (7.7) Symptoms NR Training status Other	3	2*(Flexibility, Resistance);1*(Flexibility)	Eccentric strengthening for the wrist extensors in subjects with lateral epicondylitis demonstrated improvement at six weeks but was not statistically different from that achieved with a conservative program with stretching or a concentric strengthening program.
Marzetti 2014 Italy ⁵⁹	RCT	Rotator cuff - subacromial impingement	N= 48 % female 61.4 Age 62.1 (12.5) Symptoms NR Training status Other	2	1*(Flexibility, Resistance);1*(Proprioception)	Neurocognitive rehabilitation is effective in reducing pain and improving function in patients with shoulder impingement syndrome, with benefits maintained for at least 24 weeks.
McCormack 2016 United States ⁶⁰	RCT	Achilles	N= 15 % female 68.8 Age 53.6 (38-69)** Symptoms 9.9 (NR) Training status Other	1	(Resistance)	Soft tissue treatment (Astym) plus eccentric exercise was more effective than eccentric exercise alone at improving function during both short- (26 weeks) and long-term (52 weeks) follow-up periods.
Melegati 2000 Italy ⁶¹	RCT	Rotator cuff - subacromial impingement	N= 90 % female 65.5 Age 54.4 (3.0) Symptoms NR Training status Other	1	(Flexibility, Resistance)	Groups A (kinesitherapy) and B (ESWT + kinesitherapy) achieved a significant constant score improvement, whereas the increase in group C (control) was not significant.
Mulligan 2016 United States ⁶²	RCT	Rotator cuff - subacromial impingement	N=50 % female 65 Age 50.1 (10.7) Symptoms 7.9 (7.4) Training status Other	1	(Proprioception, Resistance)	Patients with SAIS demonstrate improvement in pain and function with a standardized program of physical therapy regardless of group exercise sequencing.
Nishizuka 2017 Japan ⁶³	RCT	Lateral elbow/tennis elbow	N=110 % female 39.1 Age 53.6 (11.8) Symptoms 2.04 (1.77)	1	(Flexibility)	A forearm band may have no more than a placebo effect and is not recommended based on its effectiveness.

			Training status Other			
Nørregaard 2007 Denmark ⁶⁴	RCT	Achilles	N= 35 % female 49.0 Age 42.0 (2.0)*** Symptoms 28.4 (8.8)*** Training status Other	2	1*(Resistance);1*(Flexibility)	Symptoms gradually improved during the 1-year follow-up period and were significantly better assessed by pain and symptoms after 3 weeks and all later visits. However, no significant differences could be observed between the two groups.
Nowotny 2018 Germany ⁶⁵	RCT	Lateral elbow/tennis elbow	N= 31 % female 57 Age 46 (NR) Symptoms NR Training status Other	1	(Resistance)	The use of an elbow orthosis appears to reduce pain and improve other subjective outcome measures. However, the long-term results do not appear to be any greater than those received through Physiotherapy alone.
Østerås 2010 Norway ⁶⁶	RCT	Rotator cuff - subacromial impingement	N=61 % female 20.5 Age 43.9 (13) Symptoms 40.2 (56.3) Training status Other	2	2*(Flexibility, Resistance)	In long-term subacromial pain syndrome, high dosage medical exercise therapy is superior to a conventional low dosage exercise programme
Paavola 2018 Finland ⁶⁷	RCT	Rotator cuff - subacromial impingement	N= 186 % female 69.8 Age 50.6 (5.0) Symptoms 19.5 (18.9) Training status NR	1	(Flexibility, Proprioception, Resistance)	Arthroscopic subacromial decompression provided no benefit over diagnostic arthroscopy in patients with shoulder impingement syndrome.
Park 2010 Korea (Republic of) ⁶⁸	RCT	Lateral elbow/tennis elbow	N=31 % female 61.3 Age 50.2 (34-63)** Symptoms 6.3 (2-17)** Training status NR	1	(Resistance)	Isometric strengthening exercises done early in the course of LE (within 4 weeks) provides a clinically significant improvement.
Pearson 2012 New Zealand ⁶⁹	RCT	Patellar	N= 40 % female 62.5 Age 50.0 (8.2) Symptoms 11.0 (10.0) Training status Other	1	(Resistance)	There is some evidence for small short-term symptomatic improvements with the addition of autologous blood injection to standard treatment for Achilles tendinopathy.
Pearson 2018 Australia ⁷⁰	RCT	Achilles	N= 16 % female 0 Age 28 (4.25) Symptoms 34.17 (1.95) Training	2	2*(Resistance)	Pain was significantly reduced after isometric loading on both SLDS and hop tests. Pain and quadriceps function improved over the 4 weeks. Short-duration isometric

			status Performance			contractions are found to be as effective as longer duration contractions for relieving patellar tendon pain when total time under tension is equalized.
Pekyavas 2016 Turkey ⁷¹	RCT	Rotator cuff - subacromial impingement	N=70 % female NR Age 47.1 (13.8) Symptoms NR Training status Other	1	(Flexibility, Resistance)	HILT and MT were found to be more effective in reducing pain and disability and improving ROM in patient with SAIS.
Petersen 2007 Germany ⁷²	RCT	Achilles	N= 86 % female 40.0 Age 42.5 (11.1) Symptoms 7.4 (2.3) Training status Recreational	1	(Resistance)	The AirHeel brace is as effective as eccentric training in the treatment of chronic Achilles tendinopathy. There is no added benefit to combining both treatments.
Peterson 2011 Sweden ⁷³	RCT	Lateral elbow/tennis elbow	N= 81 % female 42 Age 48.25 (8.35) Symptoms 23.3 (35.9) Training status Other	2	2*(Resistance)	Exercise appears to be superior to the control group in reducing pain in chronic lateral epicondylosis.
Peterson 2014 Sweden ⁷⁴	RCT	Lateral elbow/tennis elbow	N= 120 % female 47.5 Age 47.9 (8.1) Symptoms NR Training status Other	1	(Resistance)	Eccentric graded exercise reduced pain and increased muscle strength in chronic tennis elbow more effectively than concentric graded exercise at follow-up. However, there were no significant differences in function or quality of life measures between the two groups.
Polimeni 2003 Italy ⁷⁵	RCT	Rotator cuff - subacromial impingement	N= 50 % female 72.0 Age 56 (16) Symptoms NR Training status Other	1	(Flexibility)	All patients experienced improvement with treatment, but the association of physical therapy and functional rehabilitation did not seem to lead to added benefit for the patient.
Praet 2019 Australia ⁷⁶	RCT	Achilles	N= 20 % female 35.0 Age 43.7 (7.9) Symptoms 54 (90) Training status Recreational	1	(Resistance)	Oral supplementation of specific collagen peptides may accelerate the clinical benefits of a well-structured calf-strengthening and return-to-running programme in patients with chronic Achilles tendinopathy.
Rabusin 2020 Australia ⁷⁷	RCT	Achilles	N= 100 % female 52.0	1	(Resistance)	In adults with mid-portion Achilles tendinopathy, heel lifts were more effective

			Age 45.85 (9.4) Symptoms 20.25 (NR) Training status Other			than calf muscle eccentric exercise in reducing pain and improving function at 12 weeks.
Reyhan 2020 Turkey ⁷⁸	RCT	Lateral elbow/tennis elbow	N= 40 % female 82.5 Age 42.4 (9.9) Symptoms 4 (0.78) Training status Other	1	(Flexibility, Resistance)	MWM plus exercise and cold therapy is safe and effective at improving elbow pain, functional capacity, and grip strength.
Rio 2017 Australia ⁷⁹	RCT	Patellar	N= 20 % female 10.0 Age 22.5 (4.7) Symptoms NR Training status Performance	2	2*(Resistance)	Both isometric and isotonic contraction protocols appear efficacious for inseason athletes to reduce pain, however, isometric contractions demonstrated significantly greater immediate analgesia throughout the 4-week trial.
Romero- Morales 2020 Spain ⁸⁰	RCT	Achilles	N= 61 % female 26 Age 41.6 (8.7) Symptoms 4.25 (3.5) Training status Other	2	1*(Resistance, Vibration);1*(Resistance)	Authors encourage the use of vibration with respect to cryotherapy added to eccentric exercise programs in order to enhance multifidus cross-sectional area in addition to lower limb functionality in individuals who suffer from chronic non-insertional AT.
Rompe 2007 Germany ⁸¹	RCT	Achilles	N=75 % female 61.3 Age 48.5 (10.6) Symptoms 10.8 (8.5) Training status Other	1	(Flexibility, Resistance)	At 4-month follow-up, eccentric loading and low-energy shock-wave therapy showed comparable results. The wait-and-see strategy was ineffective for the management of chronic recalcitrant Achilles tendinopathy.
Rompe 2009 Germany ⁸²	RCT	Achilles	N= 68 % female 55.9 Age 49.7 (9.9) Symptoms 14.5 (6.0) Training status Other	1	(Resistance)	The likelihood of recovery after 4 months was higher after a combined approach of both eccentric loading and shock-wave therapy compared to eccentric loading alone.
Rompe 2008 Germany ⁸³	RCT	Achilles	N= 50 % female 60.0 Age 39.8 (11) Symptoms 25.55 (9.45) Training status Other	I	(Resistance)	Eccentric loading as applied in the present study showed inferior results to low-energy shock wave therapy as applied in patients with chronic recalcitrant tendinopathy of the insertion of the Achilles tendon at four months follow-up.

Roos 2004 Sweden ⁸⁴	RCT	Achilles	N= 44 % female 52.3	1	(Resistance)	Eccentric exercises reduce pain and improve function in patients with Achilles
			Age 45 (26-60)** Symptoms 5.5 (1-180)* Training status Recreational			tendinopathy.
Şenbursa 2011 Turkey ⁸⁵	RCT	Rotator cuff - subacromial impingement	N= 47 % female NR Age 49.0 (9.3) Symptoms NR Training status Other	2	2*(Flexibility, Resistance)	Supervised exercise, supervised and MT, and home-based exercise are all effective and promising treatments for patients with subacromial impingement syndrome. The addition of an initial MT may improve outcomes with exercise.
Seven 2017 Turkey ⁸⁶	RCT	Rotator cuff - subacromial impingement	N= 101 % female 45.5 Age 48.5 (11.6) Symptoms 19.5 (12.4) Training status Other	1	(Proprioception)	Prolotherapy is an easily applicable treatment which may be superior in enhancing pain and function outcomes in comparison to exercise alone.
Sevier 2015 United States ⁸⁷	RCT	Lateral elbow/tennis elbow	N= 90 % female 57.9 Age 46.95 (6.55) Symptoms NR Training status Other	1	(Flexibility, Resistance)	Astym therapy is an effective treatment option for patients with LE tendinopathy, as an initial treatment, and after an eccentric exercise program has failed.
Silbernagel 2007 Sweden ⁸⁸	RCT	Achilles	N= 38 % female 47.4 Age 46.0 (8.0) Symptoms 36.2 (66.5) Training status Other	2	1*(Flexibility, Plyometric, Resistance);1*(Flexibility, Plyometric)	Our treatment protocol which gradually increases the load on the Achilles tendon and calf muscle, demonstrated significant improvements. Continuing tendon loading activity such as running and jumping with the use of a painmonitoring model did not have any adverse effect.
Silbernagel 2001 Sweden ⁸⁹	RCT	Achilles	N= 47 % female 22.5 Age 44.0 (12.5) Symptoms 30.5 (40.7) Training status Recreational	2	1*(Flexibility, Proprioception, Resistance);1*(Flexibility)	The eccentric overload protocol used in the present study can be recommended for patients with chronic pain from the Achilles tendon. More patients achieved full recovery, improved pain and ROM in the Exp group compared to the control group.
Şimşek, 2013 Turkey ⁹⁰	RCT	Rotator cuff - subacromial impingement	N= 38 % female 65.8 Age 51.0 (18-69)** Symptoms NR	1	(Proprioception, Resistance)	Findings were inconclusive and require further research.

			Training status Other			
Stasinopoulos 2017 Cyprus ⁹¹	RCT	Lateral elbow/tennis elbow	N= 34 % female 55.8 Age 43.7 (4.6) Symptoms 6 (NR) Training status Recreational	3	2*(Flexibility, Resistance);1*(Resistance)	Eccentric training, eccentric-concentric training, and eccentric-concentric training combined with isometric contraction reduced pain and improved function at the end of the treatment and follow-up. The eccentric-concentric training combined with isometric contraction produced the largest effect at the end of the treatment and follow-up.
Stasinopoulos 2006 Greece ⁹²	Quasi- experimental	Lateral elbow/tennis elbow	N= 75 % female 38.6% Age 40.3 (5.8) Symptoms 5 (NR) Training status Other	1	(Flexibility, Resistance)	Cyriax Physiotherapy, a supervised exercise programme, and polarized polychromatic non-coherent light reduced pain and improved function at the end of the treatment and at any of the follow-up time points. The supervised exercise programme produced the largest effect in the short, intermediate and long term.
Stasinopoulos 2010 Greece ⁹³	Quasi- experimental	Lateral elbow/tennis elbow	N= 70 % female 52.9 Age 45.1 (5.8) Symptoms 5 (NR) Training status NR	2	2*(Flexibility, Resistance)	Supervised exercise programme is superior to home exercise programme to reduce pain and improve function in patients with LET at the end of the treatment and at the follow-up.
Stefansson 2019 Iceland ⁹⁴	RCT	Achilles	N= 58 % female 20.0 Age NR Symptoms NR Training status Other	1	(Resistance)	Similar results for pressure massage and eccentric exercise. Combining pressure massage and eccentric exercise did not improve outcomes
Steunebrink 2013 Netherlands ⁹⁵	RCT	Patellar	N= 33 % female 24.2 Age 32.9 (10) Symptoms 11 (8) Training status Recreational	1	(Resistance)	Continuous topical GTN treatment in addition to an eccentric exercise programme does not improve clinical outcome compared to placebo patches and an eccentric exercise programme in patients with chronic patellar tendinopathy.
Stevens 2014 United Kingdom ⁹⁶	RCT	Achilles	N= 28 % female 60.7 Age 48.7 (10.8) Symptoms 7.4 (4.0) Training status Other	2	2*(Resistance)	Performing a 6-week do-as- tolerated program of eccentric heel-drop exercises compared to the recommended 180 repetitions per day, did not lead to lesser improvement for individuals with midportion Achilles

						tendinopathy, based on VISA-A and VAS scores.
Svernlov 2001 Sweden ⁹⁷	Quasi- experimental	Lateral elbow/tennis elbow	N= 57 % female 61.3 Age 50.15 (NR) Symptoms 6.3 (NR) Training status Other	1	(Flexibility, Resistance)	Significant improvements observed for VAS and grip strength warrants clinical use of this regime.
Tahran 2020 Turkey ⁹⁸	RCT	Rotator cuff - subacromial impingement	N= 67 % female 30.5 Age 52.9 (11.0) Symptoms NR Training status Other	2	2*(Flexibility)	All treatments improved pain, shoulder mobility, function, and disability in patients with SIS. However, modified posterior shoulder stretching exercises in addition to a treatment program was superior to the treatment program alone in improving pain with activity, internal rotation ROM, and dysfunction. Moreover, stretching provided clinically significant improvements.
Tonks 2007 United Kingdom ⁹⁹	RCT	Lateral elbow/tennis elbow	N= 34 % female NR Age 44.3 (7.1) Symptoms NR Training status Other	1	(Flexibility, Resistance)	Patients who received steroid injection were statistically significantly better for all outcome measures at follow up. No statistically significant effect of Physiotherapy nor interaction between Physiotherapy and injection was found.
Turgut 2017 Turkey ¹⁰⁰	RCT	Rotator cuff - subacromial impingement	N= 30 % female 46.7 Age 36.45 (17.5) Symptoms 6.28 (5.4) Training status Other	2	1*(Flexibility, Proprioception, Resistance);1*(Flexibility)	Progressive exercise training independent from specific scapular stabilization exercises provides decreased disability and pain severity in impingement syndrome. All groups showed improvement, however, there were no significant differences between the groups.
Vallés- Carrascosa 2018 Spain ¹⁰¹	RCT	Rotator cuff - subacromial impingement	N= 22 % female 54 Age 59.0 (58.5-70.0)* Symptoms Training status Other	2	2*(Flexibility, Resistance)	Both rotator cuff eccentric exercise protocols with scapular stabilising and stretching of upper trapezius were equally effective in improving pain, function, and active ROM in the short-term in patients with subacromial
vanArk 2016 Australia ¹⁰²	RCT	Patellar	N= 19 % female 6.9 Age 23 (4.7) Symptoms 35.8 (33.8) Training status Recreational	2	2*(Resistance)	syndrome. This study found favourable results for athletes with patellar tendinopathy without modification of the training. Both isometric and isotonic exercise programs reduced pain and improve function in athletes with patellar tendinopathy during a season.

Vinuesa- Montoya 2017 Spain ¹⁰³	RCT	Rotator cuff - subacromial impingement	N= 40 % female 26.8 Age 47.0 (9.0) Symptoms 6.2 (3.8) Training status Other	1	(Flexibility, Resistance)	Cervicothoracic manipulative treatment with mobilisation plus exercise therapy may improve intensity of pain and ROM compared with home exercise alone.
Visnes 2005 Norway ¹⁰⁴	RCT	Patellar	N= 29 % female 38.5 Age 26.58 (NR) Symptoms 73.6 (62.3) Training status Performance	1	(Resistance)	There was no effect on knee function (VISA) from a 12-week program with eccentric training among a group of volleyball players with patellar tendinopathy who continued to train and compete during the treatment period. Whether the training would be effective if the patients did not participate in sports activity is not known.
Vuvan 2019 Australia ¹⁰⁵	RCT	Lateral elbow/tennis elbow	N= 39 % female 28 Age 48.5 (9) Symptoms 4 (NR) Training status Other	2	2*(Flexibility, Resistance)	Unsupervised isometric exercise was effective in improving pain and disability, but not perceived rating of change and painfree grip strength when compared with wait-and-see at 8 wk. With only one of the three primary outcomes being significantly improved, it is doubtful if isometric exercises can be an efficacious standalone treatment.
Walther 2004 Germany ¹⁰⁶	RCT	Rotator cuff - subacromial impingement	N= 60 % female 43.3 Age 50.7 (NR) Symptoms 27.3 (NR) Training status Other	2	1*(Flexibility, Resistance);1*(Flexibility, Proprioception)	There were no statistically significant differences among the groups. Guided self-training can lead to results similar to those of conventional Physiotherapy.
Wegener 2016 Australia ¹⁰⁷	RCT	Lateral elbow/tennis elbow	N= 40 % female 70 Age 49.52 (8.09) Symptoms NR Training status NR	1	(Flexibility, Resistance)	Whilst all groups improved on key outcomes, it is possible that exercise alone and/or natural recovery were responsible for improvements.
Wen 2011 United States ¹⁰⁸	RCT	Lateral elbow/tennis elbow	N= 28 % female 46.4 Age 46 (7.3) Symptoms 3.3 (2.2) Training status Other	1	(Resistance)	The authors were unable to show any statistical advantage to eccentric exercises for lateral epicondylosis compared with local modalities and stretching exercises.
Werner 2002 Germany ¹⁰⁹	RCT	Rotator cuff - subacromial impingement	N=20 % female 50 Age 51.75	2	1*(Flexibility, Resistance);1*(Proprioception, Resistance)	Strengthening of the centering muscles around the humeral head lead to

			(NR) Symptoms 27.5 Training status Other			positive outcomes for subacromial impingement. Self-training after instruction showed no difference to physiotherapist-supervised exercises.
Wiedmann 2017 Germany ¹¹⁰	RCT	Achilles	N= 20 % female 65.0 Age 43.0 (6.0) Symptoms NR Training status Other	1	(Resistance)	Eccentric training improved the VISA-A and VAS scores after 12 weeks more than Physiotherapy treatment.
Yelland 2011 Australia ¹¹¹	RCT	Achilles	N= 43 % female NR Age 46.7 (NR) Symptoms 17 (NR) Training status Other	1	(Resistance)	Prolotherapy and particularly eccentric loading exercises combined with prolotherapy gave more rapid improvements in Achilles tendinosis symptoms than eccentric loading exercises alone. Long term VISA-A scores were similar.
Yerlikaya 2018 Turkey ¹¹²	Quasi- experimental	Lateral elbow/tennis elbow	N= 90 % female 71.1 Age 48.6 (8.8) Symptoms NR Training status Other	1	(Flexibility, Resistance)	Lateral epicondylitis does not seem to be affected by either leukocyte-rich-PRP or leukocyte-poor-PRP on pain and function in the short term.
Young 2005 Australia ¹¹³	RCT	Patellar	N= 17 % female 23.5 Age 27.3 (1.8) Symptoms NR Training status Performance	2	2*(Resistance)	Both exercise protocols improved pain and sporting function in volleyball players over 12 months. The decline squat protocol offers greater clinical gains during a rehabilitation programme for patellar tendinopathy in athletes who continue to train and play with pain.
Yu 2013 Korea (Republic of) ¹¹⁴	Quasi- experimental	Achilles	N= 32 % female 0.0 Age 30.3 (1.6) Symptoms 11.7 (2.1) Training status Other	2	1*(Resistance, Flexibility)1*(Resistance)	Eccentric strengthening was more effective than concentric strengthening in reducing pain and improving function in patients with Achilles tendinopathy.

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Supplementary file 10: List of excluded studies with reasons

Citation	Exclusion reason
Walther M, Werner A, Stahlschmidt T, <i>et al.</i> The subacromial impingement syndrome of the shoulder treated by conventional physiotherapy, self-training, and a shoulder brace: results of a prospective, randomized study. <i>J Shoulder Elbow Surg</i> 2004;1:417-23.	Duplicate
van Ark M. Patellar tendinopathy: Physical therapy and injection treatments (Doctoral dissertation, University of Groningen).2015:1-136.	Duplicate
Jensen B, Bliddal H, Danneskiold-Samsøe B. Comparison of two different treatments of lateral humeral epicondylitis" tennis elbow". A randomized controlled trial. <i>Ugeskr Laeg</i> 2001;1:1427-31.	Duplicate
Frohm A, Saartok T, Halvorsen K, <i>et al.</i> Eccentric treatment for patellar tendinopathy: a prospective randomised short-term pilot study of two rehabilitation protocols. <i>Br J Sports Med</i> 2007;41:e7.	Duplicate
Cannell LJ, Taunton JE, Clement DB, <i>et al.</i> A randomised clinical trial of the efficacy of drop squats or leg extension/leg curl exercises to treat clinically diagnosed jumper's knee in athletes: pilot study. <i>Br J Sports Med</i> 2001;35:60-64.	Duplicate
Stasinopoulos D, Manias P. Comparing two eccentric exercise programmes for the management of Achilles tendinopathy. A pilot trial. <i>J Bodyw Mov Ther</i> 2013;17:309-115.	Duplicate
Manias P, Stasinopoulos D. A controlled clinical pilot trial to study the effectiveness of ice as a supplement to the exercise programme for the management of lateral elbow tendinopathy. <i>Br J Sports Med</i> 2006;40:81-85.	Duplicate
Stergioulas A, Stergioula M, Aarskog R, <i>et al.</i> Effects of low-level laser therapy and eccentric exercises in the treatment of recreational athletes with chronic achilles tendinopathy. <i>Am J Sports Med</i> 2008;36:881–7.	Duplicate
Jonsson P, Alfredson H. Superior results with eccentric compared to concentric quadriceps training in patients with jumper's knee: a prospective randomised study. <i>Br J Sports Med</i> 2005;39:847–50	Duplicate
Heron SR, Woby SR, Thompson DP. Comparison of three types of exercise in the treatment of rotator cuff tendinopathy/shoulder impingement syndrome: A randomized controlled trial. <i>Physiotherapy</i> 2017;103:167–73.	Duplicate
Ganderton C, Semciw A, Cook J, <i>et al.</i> Gluteal loading versus sham exercises to improve pain and dysfunction in postmenopausal women with greater trochanteric pain syndrome: a randomized controlled trial. <i>J Women's Heal</i> 2018;27:815–29.	Duplicate
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Balius R, Álvarez G, Baró F, <i>et al.</i> A 3-Arm Randomized Trial for Achilles Tendinopathy: Eccentric Training, Eccentric Training Plus a Dietary Supplement Containing Mucopolysaccharides, or Passive Stretching Plus a Dietary Supplement Containing Mucopolysaccharides. <i>Curr Ther Res</i> 2016;78:1–7.	Duplicate
Tumilty S, Mani R, Baxter GD. Photobiomodulation and eccentric exercise for Achilles tendinopathy: a randomized controlled trial. <i>Lasers Med Sci</i> 2016;31:127–135.	Duplicate

van Ark M, Rio E, Cook J, et al. Clinical improvements are not explained by changes in tendon structure on UTC following an exercise program for patellar tendinopathy. <i>Am J Phys Med</i> 2018;97:708-714.	Duplicate
Blume CL. Comparison of an eccentric exercise intervention to a concentric exercise intervention in adults with subacromial impingement syndrome (Doctoral dissertation, Texas Woman's University). 2014:1-218.	Duplicate
Coombes BK, Bisset L, Brooks P, Khan A, Vicenzino B. Effect of corticosteroid injection, physiotherapy, or both on clinical outcomes in patients with unilateral lateral epicondylalgia: a randomized controlled trial. <i>JAMA</i> . 2013;309:461-469.	Duplicate
Berg OK, Paulsberg F, Brabant C, <i>et al.</i> High-Intensity Shoulder Abduction Exercise in Subacromial Pain Syndrome. <i>Med Sci Sports Exerc</i> 2021;53:1-9.	Insufficient exercise data
Jensen B, Bliddal H, Danneskiold-Samsøe B. Comparison of two different treatments of lateral humeral epicondylitis" tennis elbow". A randomized controlled trial. <i>Ugeskr Laeg</i> 2001;163:1427-1431.	Insufficient exercise data
Chapman-Jones D, Hill D. Novel microcurrent treatment is more effective than conventional therapy for chronic Achilles tendinopathy: randomised comparative trial. <i>Physiotherapy</i> 2002;1:471-80.	Insufficient exercise data
Kumar N, Nehru A, Rajalakshmi D. Effect of taping as a component of conservative treatment for subacromial impingement syndrome. <i>Health</i> 2012;26:237-241.	Insufficient exercise data
Schmitt J, Haake M, Tosch A, <i>et al.</i> Low-energy extracorporeal shock-wave treatment (ESWT) for tendinitis of the supraspinatus: a prospective, randomised study. <i>J Bone Surg Joint Am</i> 2001;83:873-876.	Insufficient exercise data
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Tetschke E, Rudolf M, Lohmann CH, <i>et al.</i> Autologous proliferative therapies in recalcitrant lateral epicondylitis. <i>Am J Phys Med Rehabil</i> 2015;1:696-706.	Insufficient exercise data
Coff L, Massy-Westropp N, Caragianis S. Randomized controlled trial of a new electrical modality (InterX) and soft tissue massage, stretching, ultrasound and exercise for treating lateral epicondylitis. <i>Hand Ther</i> 2009;14:46-52.	Insufficient exercise data
Furia JP. High-energy extracorporeal shock wave therapy as a treatment for insertional Achilles tendinopathy. <i>Am J Sports Med</i> 2006;34:733-740.	Insufficient exercise data
Cloke DJ, Watson H, Purdy S, <i>et al.</i> A pilot randomized, controlled trial of treatment for painful arc of the shoulder. <i>J Shoulder Elbow Surg</i> 2008;17:S17-21.	Insufficient exercise data
Krogh TP, Ellingsen T, Christensen R, <i>et al.</i> Ultrasound-guided injection therapy of Achilles tendinopathy with platelet-rich plasma or saline: a randomized, blinded, placebo-controlled trial. <i>Am J Sports Med</i> 2016;44:1990-1997.	Insufficient exercise data
Rasmussen S, Christensen M, Mathiesen I, <i>et al.</i> Shockwave therapy for chronic Achilles tendinopathy: a double-blind, randomized clinical trial of efficacy. <i>Acta Orthop</i> 2008;79:249-256.	Insufficient exercise data
Branson R, Naidu K, du Toit C, <i>et al.</i> Comparison of corticosteroid, autologous blood or sclerosant injections for chronic tennis elbow. <i>J Sci Med Sport</i> 2017;20:528-533.	Insufficient exercise data
Yuksel E, Yesilyaprak SS. The Effectiveness of Scapular Stabilization Exercises in Patients with Subacromial Impingement Syndrome and Scapular Dyskinesis. <i>Ann Rheum Dis</i> 2015;74:1316.	Insufficient exercise data

Dragoo JL, Braun HJ, Wasterlain AS. Platelet-Rich Plasma as a Treatment for Patellar Tendinopathy: A Double-Blind Randomized Controlled Trial. <i>Am J Sports Med</i> 2014;42:610-618.	Insufficient exercise data
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extracorporeal shockwave therapy alone for subacromial impingement	
syndrome? A randomized clinical trial. Journal of Orthopaedic & Sports Physical Not including exerc	ise
Therapy. 2016 Sep;46(9):714-25.	.150
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impingement syndrome: randomised controlled study. Bmj. 2012 Feb 20;344. only treatment arm	
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of motion and function in individuals with subacromial impingement syndrome.	
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in chronic partial supraspinatus tears. Iranian Red Crescent Medical Journal. Not including exerc	ica
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Thijs KM, Zwerver J, Backx FJ, Steeneken V, Rayer S, Groenenboom P, Moen	
MH. Effectiveness of shockwave treatment combined with eccentric training for	
patellar tendinopathy: a double-blinded randomized study. Clinical journal of Not including exerc	ise
sport medicine. 2017 Mar 1;27(2):89-96. only treatment arm	
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in conservative treatment of subacromial impingement syndrome. Clinical Not including exerc	ise
rheumatology. 2007 Aug;26(8):1234-9. only treatment arm	
Gedrimė DJ, Gedrimas D, Karpavičienė A, Skurvydas A. Effect of Visual and	
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	150
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physical therapy and physical therapy alone on the management of patients with	
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Lee WC, Ng GY, Zhang ZJ, Malliaras P, Masci L, Fu SN. Changes on tendon stiffness and clinical outcomes in athletes are associated with patellar tendinopathy after eccentric exercise. Clinical Journal of Sport Medicine. 2020 Jan 1;30(1):25-32.	Not including exercise only treatment arm
Bağcıer F, Yılmaz N. The Impact of Extracorporeal Shock Wave Therapy and Dry Needling Combination on the Pain, Grip Strength and Functionality in Patients Diagnosed with Lateral Epicondylitis. Turkish Journal of Osteoporosis/Turk Osteoporoz Dergisi. 2019 Aug 1;25(2).	Not including exercise only treatment arm
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Tyler TF, Thomas GC, Nicholas SJ, McHugh MP. Addition of isolated wrist extensor eccentric exercise to standard treatment for chronic lateral epicondylosis: a prospective randomized trial. Journal of Shoulder and Elbow surgery. 2010 Sep 1;19(6):917-22.	Not including exercise only treatment arm
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Rodríguez-Huguet M, Góngora-Rodríguez J, Lomas-Vega R, Martín-Valero R, Díaz-Fernández Á, Obrero-Gaitán E, Ibáñez-Vera AJ, Rodríguez-Almagro D. Percutaneous electrolysis in the treatment of lateral epicondylalgia: A single-blind randomized controlled trial. Journal of Clinical Medicine. 2020 Jul 1;9(7):2068.	Not including exercise only treatment arm
Ramon S, Russo S, Santoboni F, Lucenteforte G, Di Luise C, de Unzurrunzaga R, Vetrano M, Albano M, Baldini R, Cugat R, Stella G. Focused shockwave treatment for greater trochanteric pain syndrome: a multicenter, randomized, controlled clinical trial. JBJS. 2020 Aug 5;102(15):1305-11.	Not including exercise only treatment arm
Buyuksireci DE, Turk AC. Evaluation of the effectiveness of dexamethasone iontophoresis in patients with subacromial impingement syndrome. Journal of Orthopaedic Science. 2021 Sep 1;26(5):786-91.	Not including exercise only treatment arm
Beaudreuil J, Lasbleiz S, Yelnik A, Bardin T, Orcel P. Effect of dynamic humeral centering on painful active elevation of the arm in subacromial impingement syndrome: A randomized trial. Annals of Physical and Rehabilitation Medicine. 2012(55):e161.	Not including exercise only treatment arm
Pekgöz F, Taşkıran H, Mutlu EK, Atalay A, Çeliker R. Comparison of mobilization with supervised exercise for patients with subacromial impingement	Not including exercise only treatment arm

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injuries in Swedish elite sprinters and jumpers: a prospective randomised controlled clinical trial comparing two rehabilitation protocols. British journal of sports medicine. 2014 Apr 1;48(7):532-9. Jeong TH, Oh JK, Lee HJ, Yang YJ, Nha KW, Suh JS. The effect of the combined stretching and strengthening exercise on the clinical symptoms in posterior tibial tendon dysfunction patient. Journal of Korean Foot and Ankle Society. 2008;12(1):47-54. Genç E, Duymaz T. Effectiveness of kinesio taping in bicipital tendinitis treatment: A randomized controlled trial. Annals of Clinical and Analytical Medicine. 2020. Ginn K, Cohen M. Exercise therapy for shoulder pain aimed at restoring neuromuscular control: a randomized comparative clinical trial. Journal of Rehabilitation Medicine. 2005 Mar 1;37(2):115-22. Østerås H, Arild Torstensen T, Arntzen G, S Østerås B. A comparison of work absence periods and the associated costs for two different modes of exercise therapies for patients with longstanding subacromial pain. Journal of Medical Economics. 2008 Jan 1;11(3):371-81. Wrong outcomes
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Grävare Silbernagel K, Crossley KM. A proposed return-to-sport program for patients with midportion Achilles tendinopathy: rationale and implementation. journal of orthopaedic & sports physical therapy. 2015 Nov;45(11):876-86. Insufficient data
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Tonks JH. Evaluation of short-term conservative treatment in patients with tennis elbow (lateral epicondylitis): A prospective randomised, assessor-blinded trial (Doctoral dissertation, University of Central Lancashire). Insufficient data

Supplementary File 11: Risk of bias assessment for individual studies

Author, Year	Random sequence generation	Allocation concealment	Blinding of participants/pe rsonnel	Blinding of outcome assessment	Incomplete outcome bias	Selective reporting	Other bias
1. Aceituno-Gómez et al 2019	Low risk	Unclear	Low risk	Low risk	Low risk	Low risk	High risk
2. Akkaya et al 2016	Low risk	Unclear	High risk	High risk	Low risk	Unclear	Low risk
3. Alfredson et al 1998	High risk	Unclear	High risk	Unclear	Low risk	Unclear	High risk
4. Alfredson et al 1999	Not applicable (quasi)	Not applicable (quasi)	Not applicable (quasi)	Not applicable (quasi)	Low risk	Unclear	High risk
5. Arias-Buría et al 2015	Low risk	Low risk	High risk	Low risk	Low risk	Unclear	Low risk
6. AriasBuría et al 2017	Low risk	Low risk	Unclear	Low risk	Unclear	Low risk	High risk
7. Bae et al 2011	Not applicable (quasi)	Not applicable (quasi)	Unclear	Unclear	Unclear	Unclear	High risk
8. Bahr et al 2006	Low risk	Low risk	High risk	Low risk	Low risk	Unclear	Low risk
9. Balius et al 2016	Low risk	Low risk	Unclear	Low risk	Low risk	Unclear	Low risk
10. Bang et al 2000	Low risk	Unclear	Unclear	Low risk	Low risk	Unclear	Low risk
11. Başkurt et al 2011	Low risk	Unclear	High risk	Unclear	Low risk	Unclear	Low risk
12. Beyer et al 2015	Low risk	Low risk	Low risk	Low risk	Low risk	Low risk	High risk
13. Blume et al 2015	Unclear	Low risk	Low risk	Low risk	Low risk	Unclear	Low risk
14. Boudreau et al 2019	Low risk	Low risk	Low risk	Low risk	Low risk	Low risk	Low risk
15. Breda et al 2020	Low risk	Low risk	High risk	Low risk	Low risk	High risk	High risk
16. Brox et al 1999	High risk	High risk	High risk	High risk	No Data	No Data	No Data
17. Calis et al 2011	Low risk	Low risk	Unclear	Unclear	Low risk	Unclear	Unclear
18. Chaconas et al 2017	Low risk	Unclear	Unclear	Low risk	High risk	Unclear	High risk
19. Cheng et al 2007	High risk	High risk	Unclear	Unclear	Unclear	Unclear	High risk
20. Cho et al 2017	High risk	High risk	Unclear	Unclear	Low risk	Low risk	Unclear
21. De Jonge et al 2008	Unclear	Low risk	Low risk	Low risk	Low risk	Low risk	Low risk
22. De Oliveira 2021	Low risk	Low risk	Low risk	Low risk	Low risk	High risk	High risk
23. De Vos et al 2007	Low risk	Unclear	Low risk	Low risk	Unclear	Low risk	High risk

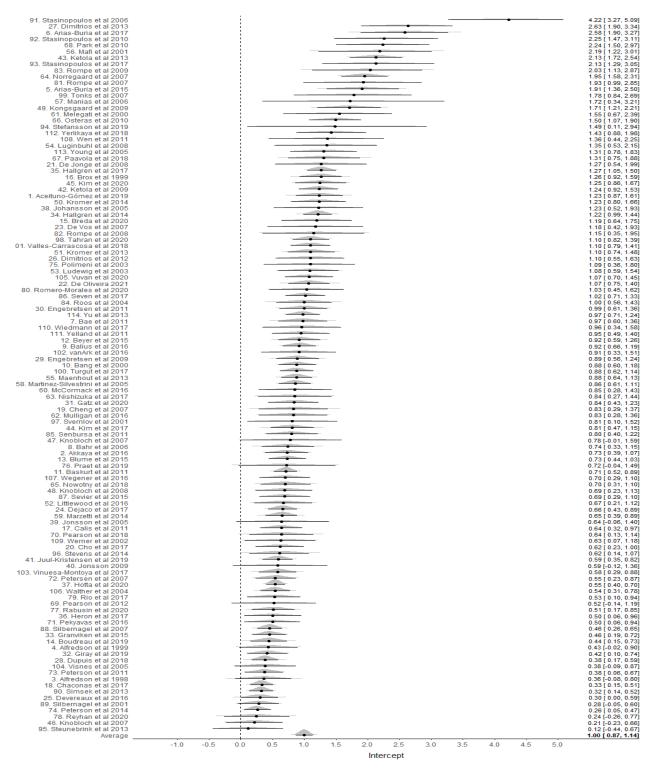
Author, Year	Random sequence generation	Allocation concealment	Blinding of participants/pe rsonnel	Blinding of outcome assessment	Incomplete outcome bias	Selective reporting	Other bias
24. Dejaco et al 2017	Low risk	Low risk	Low risk	Low risk	Low risk	High risk	Low risk
25. Devereaux et al 2016	Low risk	High risk	High risk	High risk	High risk	Unclear	Low risk
26. Dimitrios et al 2012	Not applicable (quasi)	Not applicable (quasi)	Low risk	Low risk	Low risk	Unclear	High risk
27. Dimitrios et al 2013	Not applicable (quasi)	Not applicable (quasi)	Low risk	Low risk	Low risk	Unclear	High risk
28. Dupuis et al 2018	Low risk	Low risk	High risk	Low risk	Low risk	Low risk	High risk
29. Engebretsen et al 2009	Low risk	Low risk	High risk	Low risk	Low risk	Unclear	Low risk
30. Engebretsen et al 2011	Low risk	Low risk	High risk	Low risk	Low risk	Unclear	Low risk
31. Gatz et al 2020	Low risk	Low risk	Low risk	Low risk	Unclear	Unclear	High risk
32. Giray et al 2019	Low risk	Low risk	High risk	Low risk	Low risk	Low risk	High risk
33. Granviken et al 2015	Low risk	Low risk	Low risk	Low risk	Low risk	Low risk	Low risk
34. Hallgren et al 2014	High risk	Low risk	High risk	Low risk	Unclear	Low risk	Low risk
35. Hallgren et al 2017	Unclear	Unclear	Low risk	Low risk	Low risk	Low risk	Low risk
36. Heron et al 2017	Low risk	Low risk	Low risk	Low risk	High risk	High risk	Low risk
37. Hotta et al 2020	Low risk	Low risk	High risk	Low risk	Low risk	Low risk	High risk
38. Johansson et al 2005	Low risk	Unclear	Unclear	Low risk	Low risk	Unclear	High risk
39. Jonsson et al 2005	Unclear	Unclear	Low risk	Unclear	High risk	Unclear	High risk
40. Jonsson 2009	Unclear	Unclear	Unclear	Unclear	Unclear	Unclear	Unclear
41. Juul-Kristensen et al 2019	Low risk	Low risk	Low risk	Low risk	Low risk	Unclear	Low risk
42. Ketola et al 2009	Low risk	Low risk	Unclear	Low risk	Low risk	Unclear	High risk
43. Ketola et al 2013	Low risk	Low risk	High risk	Low risk	Low risk	Unclear	Low risk
44. Kim et al 2017	Low risk	Unclear	Unclear	Low risk	Unclear	Low risk	Low risk
45. Kim et al 2020	Low risk	Low risk	High risk	High risk	Low risk	Unclear	High risk
46. Knobloch et al 2007	Low risk	Low risk	Unclear	Unclear	High risk	Unclear	High risk

Author, Year	Random sequence generation	Allocation concealment	Blinding of participants/pe rsonnel	Blinding of outcome assessment	Incomplete outcome bias	Selective reporting	Other bias
47. Knobloch et al 2007	Unclear	Unclear	High risk	Low risk	Unclear	Unclear	High risk
48. Knobloch et al 2008	Unclear	Low risk	Unclear	Unclear	Unclear	Unclear	High risk
49. Kongsgaard et al 2009	Low risk	Low risk	High risk	Low risk	Low risk	Unclear	Low risk
50. Kromer et al 2014	Low risk	Low risk	High risk	High risk	Low risk	Low risk	Low risk
51. Kromer et al 2013	Low risk	Low risk	Low risk	Low risk	Low risk	Low risk	Low risk
52. Littlewood et al 2016	Low risk	Low risk	High risk	High risk	Unclear	Unclear	High risk
53. Ludewig et al 2003	Low risk	Unclear	High risk	Unclear	Low risk	Unclear	Low risk
54. Luginbuhl et al 2008	Unclear	Unclear	Unclear	Unclear	Unclear	Unclear	High risk
55. Maenhout et al 2013	Unclear	High risk	High risk	High risk	Low risk	Unclear	Low risk
56. Mafi et al 2001	Low risk	Unclear	Unclear	Unclear	Unclear	Unclear	High risk
57. Manias et al 2006	High risk	High risk	High risk	High risk	Low risk	Unclear	Unclear
58. Martinez-Silvestrini et al 2005	Unclear	Unclear	Unclear	Unclear	Low risk	Unclear	High risk
59. Marzetti et al 2014	Low risk	Low risk	Low risk	Low risk	Low risk	Low risk	Low risk
60. McCormack et al 2016	Low risk	Low risk	Unclear	Unclear	Low risk	Low risk	Low risk
61. Melegati et al 2000	Unclear	Unclear	Unclear	Unclear	Unclear	Unclear	High risk
62. Mulligan et al 2016	Low risk	Low risk	Low risk	Low risk	Low risk	Unclear	High risk
63. Nishizuka et al 2017	Low risk	Low risk	High risk	Unclear	Low risk	Unclear	High risk
64. Nørregaard et al 2007	Low risk	Low risk	Unclear	Unclear	Unclear	Unclear	High risk
65. Nowotny et al 2018	Low risk	Unclear	Low risk	Low risk	High risk	Unclear	High risk
66. Østerås et al 2010	Low risk	Low risk	High risk	High risk	Low risk	Unclear	High risk
67. Paavola et al 2018	Low risk	Low risk	Low risk	Low risk	Low risk	Low risk	Low risk
68. Park et al 2010	Low risk	Unclear	Unclear	Unclear	Unclear	Unclear	High risk
69. Pearson et al 2012	Unclear	Unclear	High risk	Unclear	Low risk	Unclear	High risk

Author, Year	Random sequence generation	Allocation concealment	Blinding of participants/pe rsonnel	Blinding of outcome assessment	Incomplete outcome bias	Selective reporting	Other bias
70. Pearson et al 2018	Low risk	Low risk	High risk	Low risk	Low risk	Low risk	High risk
71. Pekyavas et al 2016	Low risk	Low risk	High risk	Low risk	Unclear	Unclear	Low risk
72. Petersen et al 2007	Low risk	Unclear	Unclear	Unclear	Unclear	Unclear	High risk
73. Peterson et al 2011	Low risk	Low risk	Unclear	High risk	Low risk	Low risk	Low risk
74. Peterson et al 2014	Low risk	Unclear	Low risk	High risk	Low risk	Low risk	Low risk
75. Polimeni et al 2003	Unclear	Unclear	High risk	Low risk	Unclear	Unclear	Unclear
76. Praet et al 2019	Low risk	Low risk	Low risk	Low risk	Low risk	Unclear	Low risk
77. Rabusin et al 2020	Low risk	Low risk	High risk	High risk	Low risk	Low risk	High risk
78. Reyhan et al 2020	Low risk	Low risk	Unclear	Unclear	Low risk	Unclear	High risk
79. Rio et al 2017	Low risk	Low risk	Low risk	Low risk	Low risk	Low risk	High risk
80. Romero-Morales et al 2020	Unclear	Unclear	Unclear	Unclear	Low risk	Low risk	High risk
81. Rompe et al 2007	Low risk	Low risk	Unclear	Low risk	Low risk	Unclear	Low risk
82. Rompe et al 2008	Low risk	Low risk	Unclear	Low risk	Low risk	Unclear	Unclear
83. Rompe et al 2009	Low risk	Low risk	High risk	Low risk	Low risk	Unclear	Low risk
84. Roos et al 2004	Low risk	Unclear	Unclear	Low risk	Low risk	Unclear	Low risk
85. Şenbursa et al 2011	Low risk	Unclear	Unclear	Unclear	Low risk	Unclear	Low risk
86. Seven et al 2017	Low risk	Low risk	High risk	Low risk	High risk	Unclear	Low risk
87. Sevier et al 2015	Low risk	Unclear	High risk	High risk	High risk	Unclear	High risk
88. Silbernagel et al 2007	Low risk	Low risk	High risk	High risk	Low risk	Unclear	Low risk
89. Silbernagel et al 2001	Unclear	Unclear	Unclear	Unclear	Unclear	Unclear	High risk
90. Şimşek et al 2013	Unclear	Unclear	Unclear	Low risk	Unclear	Unclear	Unclear
91. Stasinopoulos et al 2006	Not applicable (quasi)	Not applicable (quasi)	Unclear	Low risk	Low risk	Unclear	High risk
92. Stasinopoulos et al 2010	Not applicable (quasi)	Not applicable (quasi)	Low risk	Low risk	Low risk	Unclear	High risk
93. Stasinopoulos et al 2017	Low risk	Unclear	Low risk	Low risk	Low risk	Unclear	High risk
94. Stefansson et al	Low risk	Unclear	High risk	Low risk	High risk	Unclear	Low risk

Author, Year	Random sequence generation	Allocation concealment	Blinding of participants/pe rsonnel	Blinding of outcome assessment	Incomplete outcome bias	Selective reporting	Other bias
2019							
95. Steunebrink et al 2013	Low risk	Low risk	Low risk	Low risk	Low risk	Unclear	Low risk
96. Stevens et al 2014	Unclear	Unclear	High risk	High risk	Unclear	Unclear	High risk
97. Svernlov et al 2001	Not applicable (quasi)	Not applicable (quasi)	Unclear	Unclear	Unclear	Unclear	High risk
98. Tahran et al 2020	Low risk	Unclear	Low risk	Low risk	Low risk	Unclear	Low risk
99. Tonks et al 2007	Low risk	Low risk	Low risk	High risk	High risk	Low risk	Low risk
100. Turgut et al 2017	Low risk	Unclear	Unclear	Unclear	High risk	Unclear	Low risk
101. Vallés-Carrascosa et al 2018	Low risk	Low risk	Low risk	High risk	Low risk	Low risk	High risk
102. vanArk et al 2016	Low risk	Low risk	Low risk	Unclear	Unclear	Low risk	Low risk
103. Vinuesa-Montoya et al 2017	Low risk	Low risk	High risk	Low risk	Low risk	Low risk	Low risk
104. Visnes et al 2005	Low risk	Low risk	High risk	Low risk	Unclear	Unclear	Unclear
105. Vuvan et al 2020	Low risk	Low risk	High risk	High risk	Low risk	Low risk	Low risk
106. Walther et al 2004	Unclear	Unclear	Unclear	Unclear	Low risk	Unclear	Unclear
107. Wegener et al 2016	Low risk	Low risk	High risk	Low risk	Low risk	Unclear	High risk
108. Wen et al 2011	Unclear	Unclear	Low risk	Unclear	High risk	Unclear	High risk
109. Werner et al 2002	Low risk	Unclear	Unclear	Unclear	Unclear	Unclear	High risk
110. Wiedmann et al 2017	Low risk	Low risk	Unclear	Unclear	Unclear	Unclear	High risk
111. Yelland et al 2011	Low risk	Low risk	High risk	Low risk	Low risk	Unclear	Low risk
112. Yerlikaya et al 2018	Low risk	Unclear	Low risk	Low risk	Unclear	Unclear	High risk
113. Young et al 2005	Unclear	Unclear	High risk	Low risk	High risk	Unclear	High risk
114. Yu et al 2013	Low risk	Low risk	Low risk	Unclear	Low risk	Unclear	Unclear

Supplementary file 12: Forest plot of effect sizes illustrated across studies



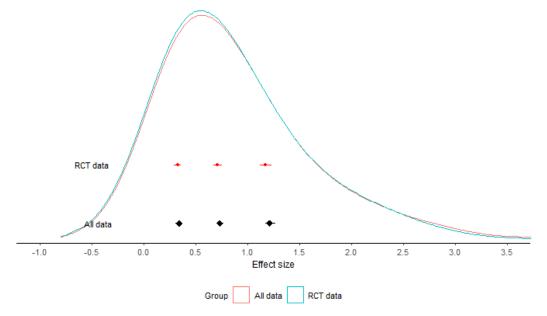
Distributions represent "shrunken estimates" based on all relevant effect sizes, the random effects model fitted, and borrowing of information across studies to reduce uncertainty. Black circles and connected intervals represent the median value and 95% credible intervals for the shrunken estimates.

Supplementary file 13: Sensitivity analyses

Sensitivity analysis checking the influence of study type was conducted by comparing the distribution of effect sizes and the small, medium, and large threshold obtained when analysing all data (114 studies) and data from randomised control studies only (110 studies). The plot below illustrates the empirical distributions using a density plot of the directly calculated effect sizes and the small medium and large thresholds:

All data - Small: $(0.25\text{-quantile}_{0.5} = 0.34 \text{ [95\%CrI: } 0.31 \text{ to } 0.37])$; Medium: $(0.5\text{-quantile}_{0.5} = 0.73 \text{ [95\%CrI: } 0.70 \text{ to } 0.77])$; and Large: $(0.75\text{-quantile}_{0.5} = 1.21 \text{ [95\%CrI: } 1.17 \text{ to } 1.27])$. Randomised control trials only - Small: $(0.25\text{-quantile}_{0.5} = 0.33 \text{ [95\%CrI: } 0.29 \text{ to } 0.36])$; Medium: $(0.5\text{-quantile}_{0.5} = 0.71 \text{ [95\%CrI: } 0.67 \text{ to } 0.75])$; and Large: $(0.75\text{-quantile}_{0.5} = 1.17 \text{ [95\%CrI: } 1.12 \text{ to } 1.24])$.

Effect size distributions across the whole data set and randomised controlled trials only with identification of small, medium, and large thresholds.



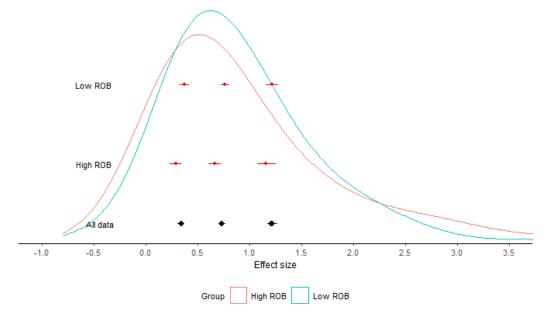
Curve represents density plot of empirical effect size distribution. Diamonds with intervals represent small, medium, and large thresholds with credible intervals (black: all data; red: RCT only).

Sensitivity analysis checking the influence of study quality was conducted by comparing the distribution of effect sizes and the small, medium, and large threshold obtained when analysing data from studies identified as low risk of bias (60 studies) and from studies identified as high risk of bias only (54 studies). The plot below illustrates the empirical distributions using a density plot of the directly calculated effect sizes and the small medium and large thresholds:

Low risk of bias - Small: (0.25-quantile_{0.5} = 0.37 [95%CrI: 0.33 to 0.41]); Medium: (0.5-quantile_{0.5} = 0.76 [95%CrI: 0.72 to 0.80]); and Large: (0.75-quantile_{0.5} = 1.22 [95%CrI: 1.16 to 1.28]). High risk of bias - Small: (0.25-quantile_{0.5} = 0.30 [95%CrI: 0.24 to 0.34]); Medium: (0.5-quantile_{0.5} = 0.68

[95%CrI: 0.61 to 0.73]); and Large: (0.75-quantile_{0.5} = 1.16 [95%CrI: 1.09 to 1.25]).

Effect size distributions across studies identified as low or high risk of bias with identification of small, medium, and large thresholds.



Curve represents density plot of empirical effect size distribution. Diamonds with intervals represent small, medium, and large thresholds with credible intervals (black: all data; red: different study quality).