Partial Meniscectomy Versus Physical Therapy

Arthroscopic partial meniscectomy versus physical therapy for degenerative meniscus lesions. How robust is the current evidence? A critical systematic review and qualitative synthesis.

Running Title: Partial meniscectomy versus physical therapy

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Abstract

Purpose:
The purpose of this systematic review was to investigate study quality and risk of bias for randomised trials comparing partial meniscectomy versus physical therapy in middle-aged patients with degenerative meniscus tears.

Methods:
A systematic review of Medline, Embase, Scopus, and Google Scholar was performed from 1990 through 2017. The inclusion criteria were: at least one validated outcome score, and middle-aged patients (40 years and older) with a degenerative meniscus tear. Studies with a sham arm, acute and concomitant injuries were excluded. Risk of bias was assessed with the Cochrane Risk of Bias Tool. The quality of studies was assessed with the Cochrane GRADE tool and quality assessment tool (EPHPP). Publication bias was assessed by funnel plot and Egger’s test. The $I^2$ statistics was calculated a measure of statistical heterogeneity.

Results:
Six studies were included and all were assessed as having a high risk of bias. There was no publication bias ($p=0.23$). All studies were downgraded (low, $n=5$; very low, $n=1$). EPHPP assessed one study as strong, two as moderate, and three as weak. The overall results demonstrated moderate to low quality of the included studies. The $I^2$ statistic was 96.2%, demonstrating substantial heterogeneity between studies.

Conclusion:
The results of this systematic review strongly suggest there is currently no compelling evidence to support arthroscopic partial meniscectomy versus physical therapy. The studies evaluated here exhibited a high risk of bias, and the weak to moderate quality of the available studies, the small sample sizes, and the diverse study characteristics
do not allow any meaningful conclusions to be drawn. Therefore, the validity of the results and conclusions of prior systematic reviews and meta-analyses must be viewed with extreme caution. The quality of the available published literature is not robust enough at this time to support allegations of superiority for either alternative, and both arthroscopic partial meniscectomy or physical therapy could be considered reasonable treatment options for this condition.

The results of this systematic review strongly suggest that the high risk of bias, the weak to moderate quality of the available studies, the small sample sizes, and the diverse study characteristics do not allow any meaningful conclusions to be drawn concerning treatment with either of these alternatives. The validity of the results and conclusions of prior systematic reviews and meta-analyses must be viewed with extreme caution. The quality of the currently available published literature is not robust enough to support allegations of superiority for either arthroscopic partial meniscectomy or physical therapy as treatment options for this condition.

**Keywords:**
Degenerative meniscus; knee arthroscopy; meniscectomy; physical therapy, physiotherapy; robustness of evidence

**Level of evidence**
Level II, systematic review of Level I and II studies
Introduction

Degenerative meniscal lesions are a common source of knee pain, and are frequently diagnosed in middle-aged and elderly patients using magnetic resonance imaging.\(^1\)\(^2\) These lesions are commonly associated with aging and osteoarthritis.\(^3\)\(^4\) Englund et al. demonstrated that the prevalence of meniscal damage in the 50-59 year age group was 32\%, and 56\% in persons aged between 70-90 years.\(^3\) However, it is still debated whether the associated symptoms occur as a consequence of the osteoarthritis, the meniscal tear, or the combination of both factors.\(^5\)\(^-\)\(^7\)

The treatment of these lesions, if they are or become symptomatic, is currently a matter of considerable controversy.\(^8\)\(^-\)\(^11\) The 2013 guidelines from the American Academy of Orthopedic Surgeons do not recommend arthroscopy with lavage and/or debridement in patients with primary symptomatic osteoarthritis of the knee.\(^12\) However, the guidelines make no recommendation either for or against partial meniscectomy in osteoarthritic patients with a torn meniscus.\(^12\) Currently accepted indications for surgery are a clear history of mechanical symptoms, such as locking and catching, with joint line pain and/or acute onset of symptoms that have failed non-surgical treatment.\(^8\)\(^,\)\(^13\) Moreover, partial arthroscopic meniscectomy and debridement is also performed with the belief that partial resection treats the underlying cause, rather than producing a placebo effect.\(^14\)

Recently, the indication for arthroscopic surgery has been challenged by several randomized studies and meta-analyses, which were unable to demonstrate any clinical benefit from surgical treatment.\(^14\)\(^-\)\(^21\) For example, Thorlund et al. reported that knee arthroscopy is associated with harm and is not recommended for middle-aged or
older patients with or without signs of osteoarthritis. The results of this study have been criticized for the inclusion of non-relevant studies and other related biases. Other studies demonstrated a superior outcome of arthroscopic partial meniscectomy in patients with symptomatic meniscal tears. Recently, Ha et al. critically reviewed the published literature and determined that valid conclusions cannot be drawn with regards to the optimal treatment for meniscal tears. Buchbinder noted that despite purportedly ‘high quality’ randomised trials suggesting that arthroscopy is no more effective than placebo or non-operative treatment, convincing evidence in support of non-operative treatment for degenerative meniscal tears is also lacking.

The purpose of this systematic review was to investigate the study quality and risk of bias of randomised trials comparing partial meniscectomy versus physical therapy in middle-aged patients with degenerative meniscus tears. We hypothesized that the quality of the currently available published literature would not robust enough to allow valid conclusions of superiority of arthroscopic treatment versus physical therapy.

Methods

The research was conducted according to the methods described in the Cochrane Handbook. The results of this study are reported according to the Preferred Reporting Items for Systematic Reviews and Meta-Analysis (PRISMA) guidelines statement.
Eligibility criteria

All randomized level I and level II studies comparing arthroscopic partial meniscectomy with physical therapy published between 1990 and 2017 were identified. Studies were included if at least one validated outcome score was utilized, treated patients were of middle-age (defined here as 40 years and older), and had a degenerative meniscus tear. Patients with osteoarthritis were not specifically excluded. The studies were excluded if the protocol included sham treatment, level III, IV, and V evidence, case reports, reviews, and letters to the editor. Concomitant injuries, such as acute and chronic cruciate or collateral ligament injuries, were also excluded. No specific restrictions were used for age in order to capture all published literature, and a final decision for inclusion or exclusion was based on a full text review.

Literature research

A systematic review of the literature was performed to identify all relevant publications in the English and German literature on June 14, 2017. Medline, Embase, Scopus, and Google Scholar were searched using the terms and Boolean operators: “meniscus tear” AND “degenerative” AND/OR “knee arthroscopy”; “partial meniscectomy” AND “physical therapy” AND/OR “physiotherapy”. Two reviewers conducted independent title and abstract screening. All eligible articles were manually cross-referenced to ensure that all potential studies were included. Disagreements regarding the included studies were resolved by consensus; if no consensus was reached, studies were subjected to a full text review.
Data extraction and quality assessment

For studies that met the inclusion criteria, an electronic data extraction form was used to obtain the following information from each article: author, journal and year of publication, any conflicts of interest, surgical technique and type, duration and number of physical therapy interventions (if available), sample size, study duration and length of follow-up, and demographic data of the study population. Two authors independently completed data extraction, and the third reviewer and senior author verified the data.

Risk of bias was assessed adapting the Cochrane Collaboration’s Risk of Bias Tool. The use of this tool allowed for the assessment of the adequacy of patient allocation, allocation concealment, blinding, clarity of outcome data, and the potential for selective reporting. Low risk of bias was allocated to studies that had low risk of bias assessments for all key domains; unclear risk of bias if one or more key domains were found to have an unclear risk of bias, and high risk of bias if one or more of the domains were assessed as high risk (Table 8.7a Cochrane Handbook).

The GRADE system was used by two reviewers to assess the quality of the evidence for each outcome measure; the third reviewer verified these assessments. The recommendations from the Cochrane Handbook were followed, and studies were downgraded if there were limitations in the design, indirectness of evidence, unexplained heterogeneity, imprecision of results, and high probability of publication bias. All institutional and author information was concealed to the third reviewer, who independently reviewed the included studies. Any disagreement between reviewers was resolved by consensus and/or arbitration of the two senior authors. The GRADE
assessments served as the main outcome measure to determine whether the conclusions of the included studies were valid or inconclusive. ‘Inconclusive’ was defined as any double or triple downgrade resulting in a low or very low quality rating. ‘Possible inconclusiveness’ was defined as a single downgrade resulting in moderate quality. Only a high quality rating was defined as ‘conclusive’. The factors resulting in a downgrade are outlined in Table 12.2.b in the Cochrane handbook: limitations in the design, indirectness of evidence, unexplained study heterogeneity or inconsistency of results, imprecision of results and high probability of publication bias. Factors that may increase the quality of the body of evidence are outlined in Table 12.2.c in the Cochrane handbook: large magnitude of effect, all plausible confounding would reduce a demonstrated effect or suggest a spurious effect when the results show no effect, dose-response gradient.

To include risk of bias assessment into the GRADE quality assessment, the following procedure was followed: studies with an unclear risk of bias were downgraded one level, whereas studies with a high risk of bias were downgraded two levels. Although this approach is somewhat arbitrary, the authors felt that it was important to incorporate risk of bias for the quality assessment. The results here will therefore be presented both with and without the risk of bias assessment.

In addition to GRADE, the Effective Public Health Practice Project (EPHPP) was used as a quality assessment tool. EPHPP was initially developed for the assessment of public health studies, and allows a comprehensive assessment of the overall quality of a quantitative study. The tool rates selection bias, study design, confounders, blinding, data collection methods, withdrawals and dropouts, intervention strategy,
and analysis. Three different quality ratings can be allocated: strong, moderate or weak. 28

**Statistical analysis**

Inter-observer differences for risk of bias and study quality were measured using Cohen’s kappa coefficient. Meta-analysis was not performed because the included studies used different outcome measures, methods of calculation, or did not consistently report measures of variability (standard deviation), thereby prohibiting pooling of the data to determine a common treatment effect. For the purposes of creating funnel plots to assess publication bias, the KOOS pain scores, 2,29,30 Lysholm scores 18,31 and WOMAC scores 16 were used to establish treatment effects, Hedge’s g, difference of means, and standard difference of means. For these ‘pooled outcomes’ a random effects model was selected. If the authors did not report standard deviations, the standard deviation was calculated using the following formula: SD= max-min/4. Hozo, et al. demonstrated that this formula provides a good estimate for standard deviation. 32 If publication bias was present based upon visual inspection of the funnel plot, Egger’s test of intercept was conducted to test for asymmetry. The I-squared statistic as a measure of statistical heterogeneity was calculated as a further measure of clinical and methodological diversity. 33 The degree of heterogeneity was defined as suggested by Higgins et al.: 0-25% low, 26-50% moderate, 51-75% moderately large and >75% high. 34 Funnel plots, as well as all statistical analyses, were performed using STATA SE (Version 12.0; StataCorp, College Station, Texas, USA) for Windows and the comprehensive meta-analysis software package (CMA), version 3 (Biostat Inc, Englewood, NJ, USA).
Figure 1. Of the initial 166 only 6 studies were then included in the analysis.
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### Table 1: Studies Included in the Analysis: Demographic and Treatment Details

<table>
<thead>
<tr>
<th>Study</th>
<th>Mean Age</th>
<th>Male Sex (%)</th>
<th>Physical Therapy</th>
<th>Surgical</th>
<th>Outcome measures</th>
<th>OA inclusion</th>
<th>Length of Follow-up</th>
<th>Loss to Follow-Up</th>
<th>Outcome</th>
</tr>
</thead>
<tbody>
<tr>
<td>Osteras 2012</td>
<td>47 – 52.7</td>
<td>88.9 – 62.5</td>
<td>N=9</td>
<td>N=8</td>
<td>Pilot study VAS, KOOS</td>
<td>Kellgren Grade 0-2</td>
<td>Not reported</td>
<td>3 mts</td>
<td>No loss</td>
</tr>
<tr>
<td>[LOE: 2]</td>
<td>PT/ASC**</td>
<td>PT/ASC**</td>
<td>'standard' partial meniscectomy</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>VAS 2 – 2.6 KOOS 39.7 – 40.9 Compliance Cons group: 84%</td>
</tr>
<tr>
<td>Herrlin 2013</td>
<td>56 - 54</td>
<td>63.3% - 67.8%</td>
<td>N=49</td>
<td>N=47</td>
<td>KOOS, Lysholm, Tegner</td>
<td>Kellgren Grade 0-3</td>
<td>0: 67.7% - 78% 1: 32.7% - 22%</td>
<td>60 mts</td>
<td>4.1% - 4.2% Lysholm 95 – 99 Tegner 3 – 3 VAS 0 - 0</td>
</tr>
<tr>
<td>[LOE: 1]</td>
<td>PT/ASC**</td>
<td>PT/ASC**</td>
<td>'standard protocol', details not mentioned</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Yim 2013</td>
<td>57.6 – 54.9</td>
<td>80 - 82</td>
<td>N=52</td>
<td>N=50</td>
<td>VAS,Lysholm, Tegner</td>
<td>Kellgren Grade 0-3</td>
<td>0: 67.7% - 78% 1: 32.7% - 22%</td>
<td>24 mts</td>
<td>Not reported VAS 1.7 – 1.8 Lysholm 84.3 – 83.2 Pain relief Complete 67% - 68% Improved 23% - 26% Persistent 10% - 6%</td>
</tr>
<tr>
<td>[LOE: 1]</td>
<td>PT/ASC**</td>
<td>PT/ASC**</td>
<td>Physical exercise 60 min 3xweekly under PT guidance x 3 weeks then 8 weeks home exercise program x 8 weeks</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Katz 2013</td>
<td>57.8 - 59</td>
<td>42.6 – 44.1</td>
<td>N=169</td>
<td>N=161</td>
<td>WOMAC, KOOS, SF36</td>
<td>Kellgren Grade 0-3</td>
<td>0: 21.3% - 21.1% 1: 20.7% - 16.1% 2: 23.1% - 23% 3: 23.1% - 28%</td>
<td>12 mts</td>
<td>Not reported WOMAC physical function improvement 18.5 – 20.9 @ 6 mts</td>
</tr>
<tr>
<td>[LOE: 1]</td>
<td>PT/ASC**</td>
<td>PT/ASC**</td>
<td>Home based exercise</td>
<td></td>
<td></td>
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</tr>
<tr>
<td>Gauffin 2014</td>
<td>74.7 – 70.1</td>
<td>3 months twice weekly, gym and home based exercise program</td>
<td>N=75</td>
<td>N=75</td>
<td>KOOS, EQ5D, VAS, PAS</td>
<td>Kellgren Grade 0-2</td>
<td>0: 43% - 49% 1: 48% - 45% 2: 9% - 5%</td>
<td>12 mts</td>
<td>13% KOOS Pain: 13.9 – 16.6 KOOS Symp: 9.8 – 15 KOOS ADL: 10.8 – 11.7 KOOS Sport: 9.2 – 21.1 KOOS QOL: 10.5 – 21.9 EQ5D: 0.06 – 0.16 VAS: 7 – 9.1</td>
</tr>
<tr>
<td>[LOE:1]</td>
<td>PT/ASC**</td>
<td>PT/ASC**</td>
<td>Partial meniscectomy</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Kise 2016</td>
<td>50.2 – 48.9</td>
<td>61% - 61%</td>
<td>N=70</td>
<td>N=70</td>
<td>KOOS, SF 36, one-leg-hop, 6m timed-hop</td>
<td>Kellgren 0-3</td>
<td>0: 70% - 73% 1: 26% - 23% 2: 3% - 4% 3: 1% -</td>
<td>24 mts</td>
<td>14.3% - 8.6% KOOS: 25.3 – 24.4 improvement @ 24 mts</td>
</tr>
<tr>
<td>[LOE: 1]</td>
<td>PT/ASC**</td>
<td>PT/ASC**</td>
<td>Neuromuscular and strength exercise over 12 weeks, 2-3 weekly session</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

*LOE: Level of Evidence; ** PT/ASC: Physical Therapy/Arthroscopy; left column figures for PT, right column data for ASC
Results

Study selection and characteristics

The literature search initially identified 166 studies. Following removal of duplicates, the abstracts of 76 studies were screened. After reviewing these 76 articles, 37 publications were excluded, and an examination of the remaining 39 full text manuscripts was conducted. The eligibility criteria were met in only 6 out of 39 articles, and these 6 studies were then included in the analysis. (Figure 1) 2,16,18,29-31

Overall, agreement between the two reviewers regarding final eligibility was excellent (kappa value 0.93, 95% CI 0.91-0.95). All six studies were published in the English language within the last 5 years, which included a total of 905 patients. The study characteristics and results are summarized in Table 1.

Risk of bias

The findings of the risk of bias assessment are summarized in Table 2. Using the criteria from the Cochrane Handbook, all six studies were assessed as having a high risk of bias. The domain ‘blinding of participants and personnel’ was assessed as a high risk for all studies. The best performing study design was that of Kise et al., yet even in this publication, in addition to the ‘blinding’ domain concerns, the domain ‘other bias’ was assessed as high risk because 38% of those eligible refused participation, and ‘reporting bias’ was assessed as unclear. 30 Visual inspection of the funnel plot did not imply asymmetry, but three studies were outside of the projected triangle, suggesting the possibility of publication bias (Figure 2). However, Egger’s intercept value was not significant (p=0.23 two-tailed) and was calculated to be -4.75 (95% CI: -14.28-4.77, t=1.38), refuting the possibility of publication bias.
Table 2: Risk of Bias Assessment

<table>
<thead>
<tr>
<th>Study</th>
<th>Random Sequence Generation (Selection Bias)</th>
<th>Allocation Concealment (Selection Bias)</th>
<th>Blinding of Participants and Personnel (Performance Bias)</th>
<th>Blinding of Outcome Assessment (Detection Bias)</th>
<th>Incomplete Outcome Data (Attrition Bias)</th>
<th>Selective reporting (Reporting Bias)</th>
<th>(Other Bias)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Osteras, 2012</td>
<td>?</td>
<td>?</td>
<td>-</td>
<td>-</td>
<td>+</td>
<td>+</td>
<td>-</td>
</tr>
<tr>
<td>Herrlin, 2013</td>
<td>+</td>
<td>+</td>
<td>-</td>
<td>-</td>
<td>+</td>
<td>?</td>
<td>+</td>
</tr>
<tr>
<td>Yim, 2013</td>
<td>?</td>
<td>+</td>
<td>-</td>
<td>-</td>
<td>?</td>
<td>+</td>
<td>?</td>
</tr>
<tr>
<td>Katz, 2013</td>
<td>+</td>
<td>?</td>
<td>-</td>
<td>-</td>
<td>+</td>
<td>?</td>
<td>-</td>
</tr>
<tr>
<td>Gauffin, 2014</td>
<td>+</td>
<td>+</td>
<td>-</td>
<td>-</td>
<td>+</td>
<td>?</td>
<td>-</td>
</tr>
<tr>
<td>Kise, 2016</td>
<td>+</td>
<td>+</td>
<td>-</td>
<td>-</td>
<td>+</td>
<td>+</td>
<td>?</td>
</tr>
</tbody>
</table>

+ low risk of bias; ? unclear risk of bias; - high risk of bias

**Figure 2:** On visual inspection of the funnel plot appeared symmetric but three studies were outside of the projected triangle, suggesting the possibility of publication bias. However, Egger’s intercept value was not significant.
Study Quality Assessment and Heterogeneity

Using the GRADE quality assessment criteria, the quality of evidence was double downgraded for five studies\textsuperscript{2,16,18,29,30} using the following two factors: limitations in the design, and imprecision of results with wide confidence intervals. The study by Herrlin et al. was triple downgraded based on the indirectness of evidence.\textsuperscript{31} These authors based inclusion on positive MRI findings and clinical history. Patients were then contacted by telephone, and following written consent were randomised. However, it appeared that the clinical examination was only performed following randomisation, and this resulted in at least one exclusion [Table 3].\textsuperscript{17,31} When incorporating both the risk of bias assessment and the GRADE quality assessment, all six studies were downgraded two levels for their high risk of bias. Furthermore, when evaluating the studies as a whole the differences in eligibility criteria, interventions, indirectness of evidence, and imprecision of results, the cumulative body of evidence was triple downgraded to a very low quality.

<table>
<thead>
<tr>
<th>Study</th>
<th>Grade</th>
<th>EPHPP</th>
</tr>
</thead>
<tbody>
<tr>
<td>Osteras, 2012</td>
<td>Low (1,4)</td>
<td>weak</td>
</tr>
<tr>
<td>Herrlin, 2013</td>
<td>Very low (1,2,4)</td>
<td>weak</td>
</tr>
<tr>
<td>Yim, 2013</td>
<td>Low (1,4)</td>
<td>moderate</td>
</tr>
<tr>
<td>Katz, 2013</td>
<td>Low (1,4)</td>
<td>weak</td>
</tr>
<tr>
<td>Gauffin, 2014</td>
<td>Low (1,4)</td>
<td>moderate</td>
</tr>
<tr>
<td>Kise, 2016</td>
<td>Low (1,4)</td>
<td>strong</td>
</tr>
</tbody>
</table>

(1) Limitations in study design or execution
(2) Indirectness of evidence
(3) Unexplained heterogeneity
(4) Imprecision of results
(5) Publication bias
The results of the quality assessment tool for quantitative studies (EPHPP) are shown in Table 3 and 4. The study by Kise et al. scored the highest on all of the items and was assessed as a strong study. The studies by Gauffin et al. and Yim et al. had only one weak score and were assessed as moderate quality studies, whereas the articles by Herrlin et al., Osteras et al., and Katz et al. were all assessed as weak, and had two or more weak item scores. The I-squared statistic was 96.2%, demonstrating substantial heterogeneity between studies.

Table 4: The Results of the Quality Assessment using the Effective Public Health Practice Project (EPHPP)

<table>
<thead>
<tr>
<th></th>
<th>Selection Bias</th>
<th>Study Design</th>
<th>Confounders</th>
<th>Blinding</th>
<th>Data Collection Methods</th>
<th>Withdrawals</th>
<th>Dropouts</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Osteras</td>
<td>3</td>
<td>1</td>
<td>2</td>
<td>3</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>weak</td>
</tr>
<tr>
<td>Herrlin</td>
<td>3</td>
<td>1</td>
<td>1</td>
<td>3</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>weak</td>
</tr>
<tr>
<td>Yim</td>
<td>2</td>
<td>1</td>
<td>1</td>
<td>3</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>moderate</td>
</tr>
<tr>
<td>Katz</td>
<td>3</td>
<td>1</td>
<td>1</td>
<td>3</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>weak</td>
</tr>
<tr>
<td>Gauffin</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>3</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>moderate</td>
</tr>
<tr>
<td>Kise</td>
<td>2</td>
<td>1</td>
<td>1</td>
<td>2</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>strong</td>
</tr>
</tbody>
</table>


**Discussion**

The results of this systematic review demonstrated high and varying risk of bias, moderate to low methodological quality, and substantial statistical heterogeneity among the six eligible randomized trials comparing arthroscopic partial meniscectomy versus physical therapy treatment in middle-aged patients with a degenerative meniscus tear. This study found that differences in eligibility criteria, outcome measures, and the nature of non-operative interventions, coupled with generally small samples contributed to a diverse but not generalizable group of studies that lacked precision. Consequently, it is clear the evidence from these studies.
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remains insufficient to reach meaningful conclusions regarding the superiority of one treatment over the other.

Recent systematic reviews and meta-analyses have questioned the benefit of surgical interventions in middle-aged patients with a symptomatic degenerative meniscus lesion. 19,20,23 Thorlund et al. performed a systematic review comparing arthroscopic partial meniscectomy and/or debridement to various control treatment ranging from placebo surgery to exercise and concluded that the benefit of surgery was small and absent only one to two years following surgery. 20 They also suggested arthroscopy might be associated with harm, such as deep vein thrombosis, pulmonary embolism, infection, or death. 20 However, eight of the nine included studies were considered to have a high risk of bias, which considerably weakened the confidence of the results. 24 Although Thorlund et al. did not assess methodological quality, four of the nine studies in their analysis were included in the present review and were deemed to be of only moderate to low quality. 2,16,18,29 Given the high risk of bias and moderate study quality, conclusions drawn from the synthesis of these studies would be considered dubious, at best.

The authors have also concluded that knee arthroscopy was associated with harm. They have included separate studies for this analysis and the scientific validity of this approach must be questioned. Hetsroni et al. investigated the incidence of symptomatic pulmonary embolism in 413,323 outpatient procedures and reported 117 adverse events (0.03%), identifying age and operating time as the only variables that significantly increased the risk. 33 Maletis et al. investigated 20,770 cases and documented a 0.25% incidence of symptomatic venous thromboembolism and 0.17% for pulmonary embolism after elective knee arthroscopy, with only one post-operative
death attributable to an embolism. A study by Hame identified the incidence of pulmonary embolus (PE) and deep vein thrombosis (DVT) in 314,578 patients undergoing arthroscopic meniscectomy over the age of 65 from Medicare data. In their study cohort 982 patients developed PE (0.3%) and 2507 patients developed DVT (0.8%). In contrast, Katz et al. directly compared adverse events between arthroscopic partial meniscectomy and physical therapy. They did not observe any differences between the two groups, suggesting that these adverse events might not be related to the surgical intervention, and instead reflect the normal incidence and prevalence of these phenomena. Katz et al. performed an a priori sample size calculation which was based on a 10 point between group difference. The difference is representative of the MCID of the WOMAC functional scale and was also observed in their preliminary observational pilot data. The target sample size was a total of 340 patients but only 334 were included in the final analysis and 48 patients crossed from physical therapy to the operative arm of the study. Applying strict scientific criteria a type II error can therefore not entirely excluded.

In a systematic review by Khan et al. seven studies were included. Five studies compared physical exercise versus arthroscopic meniscectomy, one study sham versus meniscectomy, and one study intra-articular steroid injection versus meniscectomy. Similar to Thorlund et al., the risk of bias of the included studies was unclear due to the small and non-significant effect size, and, therefore, study quality was not assessed. The conclusions by Khan et al. are cautiously worded, suggesting no benefit from arthroscopic surgery in patients with degenerative meniscal tears. However, it is possible that unidentified limitations of the included studies did not even allow them to reach valid conclusions.
For the practising physician, it is challenging to remain current with the huge volume of contemporary medical literature and the rapidly changing state of knowledge. It has been suggested that systematic reviews and meta-analyses are the highest forms of evidence, and should be considered a guideline to stay up to date on new clinical advances. However, it is essential to understand that the quality of these reviews are heavily dependent on the quality of the primary studies, and their limitations may not allow any valid conclusions to be reached. For example, Ioannidis demonstrated that approximately 20% of the currently produced meta-analyses were inherently flawed, 13% were misleading, 17% were acceptable but not useful, 27% were redundant and unnecessary, and only 3% were of good quality and useful. Therefore, the application of these evidence-based recommendations may not be beneficial, and may even be considered harmful.

Examining potential differences in the outcomes between operative or non-operative treatment can be challenging. Blinding of the subject and the provider is generally not practical. While this reality creates an inevitable assessment of “high” in the blinding domain on the Risk of Bias tool, additional measures can be included to ensure that all other aspects of the study are conducted in a manner to reduce the threat of bias. In the present review, four of the six studies did not blind the study groups to the personnel making outcome assessments. Furthermore, knowledge of the group assignment introduces the potential for detection bias, and there were various domains where the bias assessment was rated “unclear.” Three of six studies did not adequately describe methods of allocation concealment. This may reflect a reporting problem, as the allocation of group assignments may have been concealed but not reported in the
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manuscripts. However, having the knowledge of group assignment would have introduced the potential for selection bias.

In addition to the defined domains of the Risk of Bias tool, all studies included in this review had further concerns of additional biases. Four studies reported a lack of participation by eligible subjects, creating concern over selection bias. Both ‘crossover’ and ‘intention-to-treat’ analysis potentially introduce additional risk of bias. Cross-over refers to patients who were allocated to one group and switched to another against protocol. It likely contributed to the heterogeneity and certainly contributed to the studies being lower quality. One-way crossover could introduce bias due to non-adherence to the randomization protocol. With two-way crossover the likelihood of bias is decreased; however, if more patients switch from the ‘failed’ non-surgical group into the surgical treatment group, similar conclusions would apply. In some cases, the biases introduced may result in conclusions that are wrong. Four studies reported crossover ranging from 10-30%, one study did not specifically report crossover, and the pilot study by Osteras et al. had no crossover between the two small groups. With the addition of crossover bias to the other methodological flaws, the validity of the conclusions made by the individual authors of systematic reviews and meta-analyses, including this review, must be interpreted with caution.

In an effort to fully evaluate the existing evidence, this systematic review employed two different quality assessment tools in addition to the Cochrane Risk of Bias Tool. Using the GRADE quality assessment, five of the six studies were of low quality, and one study was of very low quality. When applying GRADE across the
six included studies, the overall quality had to be further downgraded due to the
differences in eligibility criteria, interventions, and outcomes from study to study. In
addition, the high risk of bias and heterogeneity further weakened the quality and
confidence of the results. 24 In contrast to GRADE, EPHPP assigned strong quality to
the study by Kise et al. 30 Despite the strong rating, the components selection bias and
blinding were assessed as only moderate, thereby introducing potential bias.

Bias and methodological quality aside, the included studies varied substantially in
terms of eligibility criteria, outcomes, and interventions. Five of the six studies
identified knee pain or daily symptoms as the main factors for inclusion, and used the
Kellgren Lawrence scale as an inclusion criteria however the distribution of the
grades varied substantially across studies. 2,16,18,29,31 In addition, only Yim et al.
specifically mentioned mechanical symptoms affecting daily living as a mandatory
requirement to determine the treatment outcomes, yet they failed to elaborate on what
consstituted ‘mechanical symptoms’. 18 The nature of non-operative intervention was
generally described and varied widely across studies. The lack of a well described and
structured rehabilitation programs makes it difficult to determine exactly what was
being compared in and amongst the studies.

Sample size was also consistently problematic among the eligible studies reviewed
here. All of the included studies performed an a-priori sample size calculation, but
only the study by Kise et al. recruited the required patients. 30 Furthermore, when
performing a post-hoc calculation the calculated power was only 32%, and in the
studies by Osteras et al. and Yim et al. the calculated post-hoc power was only 11%.
18,29 Post hoc power analysis can be considered a futile exercise, confirming studies
not reaching significance are not adequately powered. However, the wide confidence intervals among the studies included here indicate a lack of precision in estimating differences in treatment effects between arthroscopic partial meniscectomy versus physical therapy treatment. Studies with larger sample sizes are absolutely necessary to definitively determine if meaningful differences in outcomes exist between these two groups. Randomized clinical trials in orthopaedic surgery have several additional weaknesses such as validity limited to a specific study population reducing external validity, outcome measures not correlating with outcomes of interest, resource intensity and that completion may not occur until after the introduction of new treatment methods. Consensus statements based on pooling expert opinions may be a very reasonable alternative to the current evidence based approach. 

**Limitations**

Quality assessment of published research is dependent upon the subjective assessment of the investigators, and may unfortunately reflect their own biases. Even though GRADE and EPHPP allow for a certain amount of subjectivity, the high kappa value of 0.93 indicated excellent agreement and strongly suggests that the subjectivity in this review was low and almost certainly did not influence the outcome. However, this only mitigates but does not eliminate the risk of bias, as the two authors can theoretically have bias in their assessment but agree with each other. The principle limitations of systematic reviews and meta-analyses are always directly related to the limitations of the included studies. However, these limitations are not applicable here as the main purpose of this systematic review was to investigate these limitations, and to assess the rigor and strength of the currently available evidence.
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Conclusion

The results of this systematic review strongly suggest there is currently no compelling evidence to support arthroscopic partial meniscectomy versus physical therapy. The studies evaluated here exhibited a high risk of bias, and the weak to moderate quality of the available studies, the small sample sizes, and the diverse study characteristics do not allow any meaningful conclusions to be drawn. Therefore, the validity of the results and conclusions of prior systematic reviews and meta-analyses must be viewed with extreme caution. The quality of the available published literature is not robust enough at this time to support allegations of superiority for either alternative, and both arthroscopic partial meniscectomy or physical therapy could be considered reasonable treatment options for this condition.

References


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