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Effectiveness of manual thoracic therapy in treating impingement syndrome: a systematic review

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Abstract

Background: Impingement syndrome, a common cause of shoulder pain, often leads to functional limitations. Manual thoracic therapy is frequently employed as a non-surgical intervention, but its efficacy remains debated. This review assesses the impact of manual thoracic therapy on pain reduction and functional improvement in impingement syndrome.

Methods: A systematic review of randomized clinical trials was conducted, focusing on studies that applied manual thoracic therapy to patients with impingement syndrome. The primary outcomes were pain reduction and functional improvement. Studies were evaluated for methodological quality using the PEDro scale, with scores \geq 6 indicating high quality.

Results: Nine studies met the inclusion criteria. All studies demonstrated high methodological quality (PEDro score \geq 6). Pain reduction was consistent across studies, with an NPRS score reduction of 0.6 to 1.5 points immediately after treatment and up to 3.2 points at follow-up. Functionality improvements were statistically significant in some studies. However, the results showed limited homogeneity, and the majority of studies did not report substantial differences between intervention and placebo groups.

Conclusion: This review suggests that manual thoracic therapy may lead to pain reduction in impingement syndrome, with some evidence of functional improvement. However, the variability in manual therapy techniques and the limitations in research methodologies indicate a need for further controlled studies. These findings underscore the potential of manual therapy as a supplementary treatment but also highlight the necessity for more robust clinical trials to fully ascertain its effectiveness in clinical practice.

Keywords

 $Impingement\ syndrome\cdot Manual\ thoracic\ therapy\cdot Pain\ reduction\cdot Shoulder\ dysfunction\cdot Musculoskeletal\ rehabilitation$

Protocol registration PROSPERO (CRD42023491212)



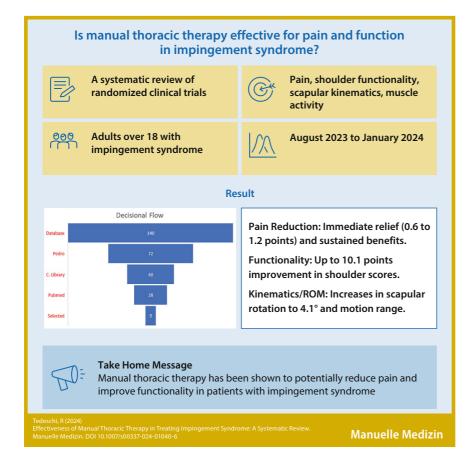
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Introduction

Shoulder impingement syndrome is emerging as a complex and multidimensional pathology, involving a variety of etiological and biomechanical factors [2, 13]. Characterized by compression

of subacromial structures, the syndrome leads to a clinical picture of pain, weakness, and paresthesia in the upper limb. Originally described by Adam in 1852 [34], the pathology was widely disseminated and characterized by the work of Charles Neer in 1972 [17], highlighting the reduc-

Graphic abstract



tion of subacromial space as a key element in the pathogenesis of the syndrome [13]. The classification of subacromial conflict syndrome, developed by Neer in 1983 [18], divides the condition into three stages based on the patient's age, clinical signs, and the evolution of the pathology. These stages range from edema and inflammation, typical in individuals under 25 years, to fibrosis and tendinitis in patients aged 25 to 40 years, up to the development of bone spurs and tendon ruptures in individuals over 40 years. It is important to note that impingement syndrome can be further classified based on the nature of the conflict, distinguishing between primary conflict, secondary conflict, and internal impingement [14, 37]. These subclassifications are based on differences in etiology and clinical presentation, such as the abnormal relationship between the rotator cuff and the coracoacromial arch in primary conflict, glenohumeral or scapulothoracic instability in secondary conflict, and contact between the greater

tuberosity and the posterior-superior face of the glenoid in internal impingement [12, 33]. In addition, a significant link has been identified between impingement syndrome and various biomechanical factors such as alterations in scapular kinematics, abnormal muscle activation, and incorrect posture [1, 4, 19, 21, 23, 25, 26]. Recent studies have emphasized how increased thoracic kyphosis can contribute to the pathology, altering scapular mechanics and reducing the subacromial space [33]. This systematic review focuses on exploring the efficacy of manual therapy techniques, particularly those applied to the thoracic district, in patients with subacromial conflict syndrome [27, 29, 32]. The objective is to assess how these interventions influence pain, functionality, scapular kinematics, and muscle strength compared to other therapeutic methods or the absence of treatment [27, 28, 31]. Considering the importance of regional interdependence and the influence of scapular kinematics in the syndrome,

this review offers a critical and updated evaluation of therapeutic options, with a particular focus on the short- and long-term outcomes of different therapeutic approaches [5–11, 22, 24, 30, 35].

Materials and methods

This systematic review was carried out following the methodological guidance contained in the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) checklist [20]. The protocol was published in the International Prospective Register of Systematic Reviews (PROSPERO) under registration number CRD42023491212.

Research method

Search strategy

An electronic bibliographic search was conducted in three databases: PubMed, Cochrane Central Register of Controlled Trials, PEDro database. The P.I.C.O.(M.) strategy was used to formulate the research question of the review. The search was conducted up to January 20, 2024, with no date restriction and no linguistic limits. The search terms used were as follows:

((shoulder pain [MeSH terms]) OR (shoulder impingement) OR (rotator cuff related shoulder pain) OR (shoulder impingement syndrome [MeSH terms])) AND ((thoracic) OR (spine)) AND ((manipulation) OR (manual therapy) OR (joint mobilization) OR (musculoskeletal manipulations [MeSH terms])). PubMed

((shoulder pain) OR (shoulder impingement) OR (rotator cuff related shoulder pain) OR (shoulder impingement syndrome)) AND ((thoracic) OR (spine)) AND ((manipulation) OR (manual therapy) OR (joint mobilization) OR (musculoskeletal manipulations)). Cochrane Central

(shoulder pain, shoulder impingement, thoracic, spine, manual therapy, mobilization, joint mobilization, manipulation). PEDro

Study selection criteria

Inclusion criteria

Inclusion criteria population: adult subjects > 18 years old, both males and

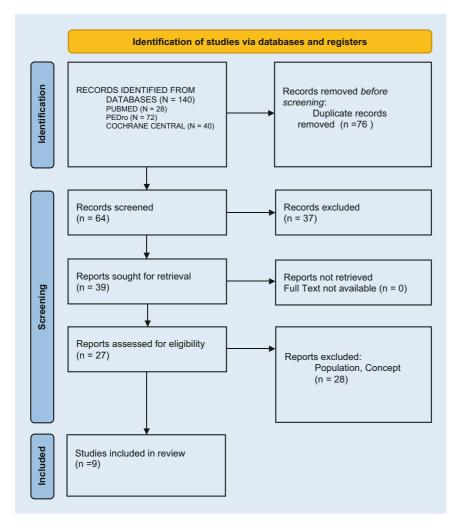


Fig. 1 ▲ Flowdiagram. (Adapted from Page et al. [38], PRISMA [39]; CCBY 4.0, https://creativecommons.org/licenses/by/4.0/deed.de)

females, with a diagnosis of impingement syndrome after recognized clinical tests. Specific shoulder pathologies (adhesive capsulitis, surgical operations, dislocations).

Intervention: mandatory: any manual therapy technique applied to the thoracic district. Optional: manual therapy to the shoulder, cervical spine, massage therapy, ortherapeutic exercise. Exclusively manual therapy to the shoulder, cervical spine, and use of instrumental physical therapies.

Outcomes: pain, shoulder functionality, scapular kinematics, muscle activity.

Studies: randomized clinical trials (RCTs).

Exclusion criteria Observational studies, secondary studies, pilot studies.

Study selection process

The records retrieved from database searching were collected and imported into EndNote V.X9 (Clarivate Analytics, Philadelphia, PA, USA). Duplicates were removed through the Endnote deduplicator tool. In the screening phase, two reviewers independently read all titles and abstracts, excluding articles that did not answer the research question. A third reviewer intervened to reach a final decision on the list of articles to be read in full text. The study selection process and the reasons for exclusion were recorded and presented in the PRISMA flow diagram.

Data extraction and assessment

The methodological quality of the intervention studies included in the review

was assessed by the researchers using the PEDro scale tool [3, 15]. The results of the assessment were entered in a table, the fulfilment of the criterion was indicated with "yes," and the absence of the specific item in the analyzed study with "no." Two independent reviewers, both of whom were experts in the field, were involved in the quality assessment. In cases of disagreement, a third reviewer with extensive research and practice experience was called upon to intervene. Training was provided by a third physiotherapist experienced in research methodology. Summary tables and graphs of the extracted data from all included studies and a narrative summary were provided.

Data analysis

The reviewers independently extracted data from the studies and summarized them in a summary table. The following data were extracted: author, year, participants, treatment description, and outcome.

For the final analysis, we considered the "NA" items as items not reported and described by the authors.

Results

From the initial search, 140 articles were retrieved. Specifically, 28 articles were found in PubMed, 72 in PEDro, and 40 in Cochrane. Of these articles, 76 were removed as duplicates, resulting in a total of 64 articles. With the first reading of titles and abstracts, 37 studies were removed. After the full reading of the remaining studies, 18 studies were eliminated for not meeting the inclusion criteria. Thus, 9 articles were included in the review (Table 1). The entire study selection process is outlined in the PRISMA statement flow diagram (Fig. 1), which details the excluded studies and the reasons for their exclusion.

Haik et al. (2014) [7]. Both the thoracic spinal manipulation (TSM) impingement group and the sham TSM impingement group reported statistically significant average reductions (0.6 points) in post-intervention pain, regardless of the treatment received (p = 0.04). 60% of the TSM

group reported pain improvement compared to 36% in the placebo therapy. The TSM impingement group showed an increase in scapular lateral rotation of 1.7° (p < 0.001) during arm elevation and 1.1° during lowering (p = 0.019) after the intervention. The asymptomatic TSM group demonstrated a 2.2° (p < 0.001) increase in scapular lateral rotation in arm elevation post-intervention, where the placebo group improved by only 1° (p = 0.05). The asymptomatic TSM group also reported a 1.9° (p < 0.001) increase in lateral scapular rotation and a 0.9° (p = 0.002) change in scapular tilt during elevation and 1.1° (p = 0.002) during lowering post-intervention. No dropouts were reported.

Kardouni et al. (2014) [10]. Following manipulation, both pain (numeric pain rating scale [NPRS] score) and functionality (Penn Shoulder Score) improved in both groups over time (p < 0.001). The NPRS score decreased by 1.1 points from pre-treatment to post-treatment and by 1.5 points from pre-treatment to followup. The Penn Shoulder Score improved by 10.1 points from pre-intervention to follow-up. No differences were observed in the global rating of change (GROC) scale between the two groups. A correlation was found between the pressure pain threshold (PPT) in the lower trapezius of the unaffected shoulder and the pretreatment Penn Score in the SMT group (p = 0.009).

Kardouni et al. (2015) [9]. No differences were observed between groups in thoracic excursion, pain (NPRS), or functionality (Penn Shoulder Score). The average NPRS scores decreased by 1 point from pre-treatment to post-treatment, and by 1.2 points from pre-treatment to follow-up (24–48 h later). The Penn Shoulder Score improved by 9.1 points from pre-treatment to follow-up. Both groups showed a mean change in scapular internal rotation of 0.9° (p = 0.003) during elevation and 0.8° (p =0.041) during lowering post-intervention. No significant differences in thoracic excursion were noted in either group posttreatment.

Riley et al. (2015) [22]. Immediately postintervention, no statistically significant differences were observed regarding functionality (shoulder pain and disability index [SPADI]) and pain (NPRS), irrespective of the message or intervention. A difference of 0.9 points in NPRS was noted between pre-treatment and follow-up (p < 0.001), but no differences were found immediately post-treatment. For the SPADI scale, an eight-point difference was noted between pre-treatment and follow-up (p < 0.001).

Haik et al. (2016) [6]. No significant differences in pain (NPRS) were observed between the two groups. The TSM group showed a 0.8-point difference on day 2 pre-intervention and 0.9 points on day 2 post-intervention. No significant differences were noted between the two groups for the functionality scales disabilities of the arm, shoulder, and hand (DASH) and Western Ontario rotator cuff index (WORC; p > 0.5). Data collected on scapular kinematics revealed a mean difference in both groups during arm elevation, of 2.6° (p < 0.01) in lateral scapular rotation on day 2 pre-intervention and 4.1° (p < 0.01) postintervention. The sham TSM group showed improvement (p = 0.04) compared to the TSM group in anterior scapular tilt during arm elevation. During arm lowering, the TSM group demonstrated improved lateral rotation (p = 0.01) on day 2 pre-intervention, post-intervention, and at follow-up. Regarding muscle activity, differences in the upper trapezius were observed, with greater activity in the sham TSM group on day 2 post-intervention (p < 0.01) and at follow-up (p < 0.01). Both groups showed decreased activity in the lower trapezius (p < 0.1) on day 2 pre-intervention, with the sham group showing this decrease post-intervention and the TSM group at follow-up. Increased activity in the upper trapezius was noted in the sham TSM group at follow-up (p < 0.01), and a decrease in the lower trapezius in the sham TSM group on day 2 pre-intervention, postintervention, and at follow-up in the TSM group (p < 0.01).

Da Silva et al. (2019) [24]. Following the intervention, a significant increase in range of motion (ROM) in flexion (p < 0.01) and abduction (p < 0.01) of the painful shoulder was observed in the manipulation group (MG), and also in abduction of the non-painful shoulder (p = 0.03). The placebo group demonstrated a significant increase in ROM in flexion of the affected shoulder (p = 0.03) and in abduction (p <0.01). No particular differences in pain were observed between groups, with a time-effect reduction in pain in both groups (-0.53 cm MG; -0.37 cm PG).

Land et al. (2019) [11]. All three groups experienced a significant decrease in pain from the initial assessment to the 12th week (p = 0.04). The upper thoracic intervention group showed improvement from baseline to the 6th week in the SPADI scale (p = 0.007), passive internal rotation (p < 0.001), and posterior shoulder range (p = 0.004). The posterior shoulder intervention group showed improvement from baseline to the 6th week in the SPADI scale (p = 0.03), passive internal rotation (p = 0.005), and posterior shoulder range (p = 0.01). The upper thoracic intervention group maintained improvements until the 12th week regarding the SPADI scale (p =0.006) and passive internal rotation (p =0.02). The posterior shoulder intervention group maintained only passive internal rotation (p = 0.04). A significant improvement in the SPADI scale was maintained until the 6-month follow-up after treatment in both the upper thoracic (p =0.05) and posterior shoulder intervention groups (p = 0.02).

Grimes et al. (2019) [5]. Following manipulation, no significant differences were observed in pain, functionality, or satisfaction (p > 0.05). No differences in scapular kinematics, pectoralis minor length, or isometric force of the middle trapezius, lower trapezius, or serratus anterior (p > 0.05)were observed between the two groups.

Hunter et al. (2022) [22]. The MET-only group showed significant improvements in pain and disability (DASH, SPADI, VAS) compared to the placebo group at discharge (DASH: -8.4, p = 0.003; SPADI: -14.7, p = 0.0; VAS: -15.5, p = 0.001), at 6 months (DASH: -11.1, p = 0.04; SPADI: -14.9, p = 0.010; VAS: -14.1, p = 0.02), and at 12 months (DASH: -13.4, p=0.013; SPADI: -19, p = 0.005; VAS: -17.3, p = 0.01). The MET + STM group showed

	type		Salliple Size, age, M/F			tion	
Haik et al. [7]	2014, Brazil, RCT	Evaluate immediate effects of thoracic manipulations on pain and scapular kinematics	N = 97 Age = 28.7 ± 7.9 F = 45, M = 52	NPRS, DASH, WORC, scapular kinematics	TSM = thrust from seated position, repeated up to 3 times if no cavitation	Sham TSM: placebo manipulations without final thrust	Pain reduction during arm elevation/lowering after a single session
Kardouni et al. [10]	2014, America, RCT	Evaluate immediate pain response to manipulative therapy using PPT	N=48 Age=31.1±12.2 F=23, M=25	NPRS, PENN, GROC, PPT of deltoid and lower trapezius	SMT = each technique applied twice, for a total of 6 manipulations	Sham SMT = placebo manipulations without final thrust	No major effects on pain sensitivity or perception compared to placebo
Kardouni et al. [9]	2015, America, RCT	Determine if SMT alters scapular kinematics, thoracic excursion, and kinematics	N = 52 Age = 32 ± 12.2 F = 24, M = 28	NPRS, PENN, GROC, scapular kinematics	SMT = twice for each thoracic vertebral segment	Sham SMT = placebo manipulations without final thrust	No immediate changes in kinematics or thoracic extension
Riley et al. [22]	2015, America, RCT	Evaluate effects of HVLA manipulations and messages on shoulder pain	N= 97 Age = 48.7 ± 11 F= 54, M= 43	NPRS, SPADI	Thoracic HVLA: thrust in PA direction at levels T1-T2, T3-T4, T6-T7	Scapular HVLA: thrust in anterolateral direction	Statistically significant improvements in pain and functionality
Haik et al. [6]	2016, Brazil, RCT	Investigate short-term effects on pain, function- ality, scapular kinematics, muscle activity	N=61 Age=31.9±11.5 F=23, M=38	NPRS, DASH, WORC, scapular kinematics, muscle activity	TSM = manipulations in the middle segment of the thoracic column	Sham TSM = placebo manipulations without final thrust	No changes in pain, in- creased lateral rotation and activity of upper trapezius
Da Silva et al. [24]	2019, Brazil, RCT	Investigate the influence of thoracic manipulations (SM) on pain and shoulder range of motion	N=60 Age=45.2±14.1 F=41, M=19	VAS, ROM shoulder	High velocity low amplitude (HVLA) manipulation applied twice in the upper segment T4-T5 from prone position	Placebo group: placebo manipulations without final thrust	Thoracic spinal manipulation causes statistically significant but not clinically important changes in reducing pain
Land et al. [11]	2019, Australia, RCT	Compare various interventions targeting the upper thoracic spine and posterior shoulder	N = 69 Age = 51 ± 5.2 F = 34, M = 35	NPRS, SPADI, thoracic ROM, glenohumeral internal rotation, and shoulder range	Upper thoracic: transverse grade 3 mobilizations from T1 to T6, costovertebral T1-T6 grade 3 mobilizations for a total of 20 min + exercise	Posterior shoulder: massage, passive mobilizations, and stretching for a total of 20 min. Control group: ultrasound	Both manual therapy interventions had a positive effect in reducing pain, improving functionality, and increasing posterior shoulder range
Grimes et al. [5]	2019, America, RCT	Compare the immediate effects of upper thoracic spine manipulation	N=60 Age=36.5±15.1 F=23, M=37	PENN, scapular kinematics, isometric force of middle trapezius, lower trapezius, serratus anterior, pectoralis minor length	TSTM supine: manipulation applied in AP direction. TSTM seated: manipulation applied in cephalic direction with traction	Sham-TSTM: placebo manipulations without final thrust	No differences between two manipulative techniques and a placebo technique in terms of pain and functionality
Hunter et al. [8]	2022, Australia, RCT	Verify if MET applied to the thoracic spine de- crease pain and disability associated with shoulder impingement	N=75 Age=60.1±10 F=28, M=47	DASH, SPADI, VAS, PSFS, GROC, Thoracic ROM	MET only: 15 min of MET technique to the thoracic spine from supine position applied up to 4 times then cranially. MET + STM: 10 min of thoracic MET and 5 min of massage to the rotator cuff and triceps	Control group: inactive laser	MET only treatment on the thoracic spine showed improvement in pain and disability compared to a placebo group

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Table 2 Summary of the PEDro scale	he PEDro scale											
Author, Year	Criteria 1*	Criteria 2	Criteria 3	Criteria 4	Criteria 5	Criteria 6	Criteria 7	Criteria 8	Criteria 9	Criteria 10	Criteria 11	Total
Haik, 2014 [7]	Yes	Yes	No	Yes	No	No	Yes	Yes	No	Yes	Yes	9
Kardouni, 2014 [10]	Yes	Yes	Yes	Yes	Yes	No	Yes	Yes	No	Yes	Yes	8
Kardouni, 2015 [9]	Yes	Yes	Yes	Yes	No	No	Yes	Yes	No	Yes	Yes	7
Riley, 2015 [22]	Yes	Yes	No	Yes	No	No	Yes	Yes	No	Yes	Yes	9
Haik, 2016 [6]	Yes	Yes	No	Yes	Yes	No	Yes	Yes	Yes	Yes	Yes	8
Da Silva, 2019 [24]	Yes	Yes	Yes	Yes	No	No	No	Yes	Yes	Yes	Yes	7
Land, 2019 [11]	Yes	Yes	Yes	Yes	No	No	Yes	No	No	Yes	Yes	9
Grimes, 2019 [5]	Yes	Yes	Yes	No	No	No	Yes	Yes	No	Yes	Yes	9
Hunter, 2022 [8]	Yes	Yes	Yes	Yes	No	No	Yes	No	Yes	Yes	Yes	7

improvements in disability but not in pain at discharge (DASH: -8.2, p = 0.006; SPADI: -13.5, p = 0.02) and at 6 months (DASH: -9.0, p = 0.024; SPADI: -12.4, p = 0.041). Regarding the patient-specific functional scale (PSFS), the MET-only group improved compared to placebo at discharge (1.3, p=0.03) and at 12 months (1.8,p = 0.008), while the MET+STM group improved at 12 months (1.7, p = 0.02). No differences were observed between the MET-only and MET+STM groups in terms of thoracic flexion-extension ROM or thoracic posture.

Risk of bias

The methodological quality with which the studies were conducted and the risk of bias to which they are subjected were assessed by means of the PEDro scale, as shown in Table 2. "Yes" was indicated as fulfilment of the criterion and "no" as the absence of the specific item in the study analyzed.

Pedro scale: 1 eligibility criteria were specified; 2 subjects were randomly allocated to groups (in a crossover study, subjects were randomly allocated an order in which treatments were received); 3 allocation was concealed; 4 the groups were similar at baseline regarding the most important prognostic indicators; 5 there was blinding of all subjects; 6 there was blinding of all therapists who administered the therapy; 7 there was blinding of all assessors who measured at least one key outcome; 8 measures of at least one key outcome were obtained from more than 85% of the subjects initially allocated to groups; 9 all subjects for whom outcome measures were available received the treatment or control condition as allocated or, where this was not the case, data for at least one key outcome was analyzed by "intention to treat;" 10 the results of betweengroup statistical comparisons are reported for at least one key outcome; 11 the study provides both point measures and measures of variability for at least one key

The first item was included for completeness with respect to the Delphi list [36], but is not included in the final count, which is based on ten, from requirement number 2 to number 11. If the criterion is

fulfilled, one point is awarded, otherwise it has a value of zero. The authors of the PEDro scale divide clinical studies into four categories, linked to the score obtained: low quality when it varies between 0 and 3; medium quality if it is 4 or 5; high quality if it is 6 to 8; excellent quality with a score of 9 or 10.

Discussion

The objective of this review was to evaluate whether manual therapy techniques applied to the thoracic district can provide benefits in terms of pain or functionality in patients with impingement syndrome. This review also included studies where, in addition to thoracic manual therapy, therapeutic exercise was performed or part of the treatment was directly dedicated to the shoulder, as this is the standard approach for patients with subacromial conflict syndrome. In the majority of the studies [7, 9-11, 22, 24], only manipulative techniques at the thoracic level were applied. Riley's study [22] associated a "row" type reinforcement exercise, Land's study [11] a supine postural exercise, and Hunter's study [8] performed a muscle energy technique (MET). Six studies [5-7, 9, 10, 24] used a sham manipulation as a control group, described in the study by Michener et al. [16] as a valid comparative tool. The four outcomes investigated by this review are pain, functionality, scapular kinematics, and muscle activation. Regarding pain, all authors agree that following treatment, the intervention groups do not show substantial differences compared to placebo groups. However, in all studies, there was a reduction in pain (NPRS) between 0.6 and 1.5 points in the immediate post-treatment period, with a maximum of 3.2 in the follow-up. Specifically, Haik's study [7] reported a difference of 0.6 points NPRS post-intervention, Kardouni's study [10] 1.1 points NPRS post-treatment, and 1.5 at the 24/48 h follow-up. Kardouni's second study [9] had a 1-point NPRS difference post-treatment and 1.2 at the 24/48 h follow-up. Rilev's study [22] found an average difference of 0.9 points NPRS between pre-treatment and 6/9 days follow-up. Haik's second study [6] found a difference of 0.8 points on the second day of intervention and 0.9 immediately after the second intervention. Land's study [11] found a 3.6-point NPRS reduction after 12 weeks of treatment. Da Silva's study [24], which used the VAS scale, achieved a 0.53-cm reduction. Finally, Hunter's study [8] demonstrated a VAS pain reduction of 1.5 cm at discharge, 1.4 at the 6-month followup, and 1.7 at the 12-month follow-up. For some authors, the average reduction in scores describing pain is attributed to the factor of time; this would explain the similarity with the scores of various placebo groups. Authors who conducted trials with analysis of outcomes immediately after treatment believe that the reduction in pain is caused by the placebo effect, patient positioning, interaction with a healthcare professional, or mobilization within the range of motion. For the outcome of functionality, as with pain, the majority of studies agree that intervention groups and placebo groups do not show differences. However, in some studies there have been statistically significant changes in assessment scale scores posttreatment or during follow-up. Specifically, Kardouni's studies [9, 10] show a decreased Penn Shoulder Scale score by 10.1 and 9.1 points, respectively, in the 24/48 h follow-up. Riley's study [22] achieved an 8-point improvement in the SPADI scale between pre-treatment and 6/9 days follow-up. Land's study [11] found a 32-point improvement in the SPADI scale in the thoracic intervention group at 12 weeks, and the results were maintained up to the 12-month follow-up. Hunter's study [8] found an improvement in the DASH and SPADI scales both at discharge with 8.4 and 14.7 points of difference, at the 6-month follow-up with 11.1 and 14.9 points of difference, and at 12 months with 13.4 and 19 points of difference compared to the placebo group, respectively. Regarding the outcome of scapular kinematics, there was heterogeneity in results, as some studies had significant changes, while others found no differences. Haik's study [7] reported improvements following manual therapy in both symptomatic and asymptomatic groups, with the symptomatic group showing an increase of 1.7° in lateral scapular rotation during elevation and 1.1° during lowering. The asymptomatic group showed an increase of 2.2° in

lateral scapular rotation during elevation and 1.9° during lowering. The asymptomatic group also showed an increase of 0.9° in scapular tilt during elevation and 1.1° during lowering. Kardouni's study [9] reported scapular improvements in medial rotation of 0.9° in both groups during elevation and 0.8° during lowering. Haik's second study [6] found an increase in lateral scapular rotation of 2.6° during elevation, improving to 4.6° after the second treatment. This improvement in lateral rotation was also observed during lowering by 4°, improving to 5.3° after the second intervention. Da Silva's study [24] measured shoulder range of motion globally, indicating significant improvements in flexion and abduction movements by 3.5° and 8.0°, respectively, as well as improved abduction by 2.0° in the non-painful shoulder. Land's study [11] reported an improvement in passive internal shoulder rotation of 17.0° at 6 weeks and 19.8° at 12 weeks. The last outcome, muscle activity, was measured in Haik's study [6], where increased activity of the upper trapezius and decreased activity of the lower trapezius were observed following manipulative therapy, in both flexion and lowering arm movements. Grimes's study [5] found no differences in isometric strength of the trapezius or serratus anterior muscles or in the length of the pectoralis minor muscle after performing manipulations. Some results obtained from various studies are described by the authors as "statistically significant but not clinically important," as minimal variations in pain and functionality scales or a few degrees in scapular range of motion can be important statistically but hardly translate into real changes in clinical practice. This review aims to verify the efficacy of thoracic manual therapy not to completely replace the standard treatment but more to integrate it within it and understand whether applying these techniques can speed up the healing process and pain reduction in patients.

Strengths and limitations

This systematic review of randomized clinical trials on manual thoracic therapy for impingement syndrome is characterized by high methodological quality, as all included studies scored ≥6 on the PEDro

scale, indicating a low risk of bias. The review examined diverse outcomes, including pain, functionality, scapular kinematics, and muscle activation. A consistent reduction in pain (NPRS score between 0.6 and 1.5 points immediately posttreatment and up to 3.2 points in followups) was observed across studies, despite no substantial differences between intervention and placebo groups. Statistically significant improvements in functionality scores were noted in some studies during post-treatment and follow-up periods. Several studies reported improvements in scapular kinematics, such as increases in lateral rotation and tilt of the scapula, and changes in muscle activity, particularly in the trapezius muscles, in response to manual therapy. The evidence was synthesized narratively, not quantitatively, limiting the ability to comprehensively measure the effect size. The variability in manual therapy techniques, due to differing levels of training and execution capabilities of therapists, could impact treatment effectiveness. The inability to apply treatments blindly and the experience of the practitioner pose challenges in objectifying research in this field. The reliance on selfadministered questionnaires or self-evaluations for assessing outcomes introduces subjectivity into the results. The review highlighted limited homogeneity in the results, questioning the therapeutic approach to impingement syndrome through thoracic manual therapy. The review aims not to replace but to supplement standard treatment, by assessing whether thoracic manual therapy can accelerate the healing process and pain reduction in patients. Future studies with more treatments and controlled randomized designs are suggested to verify the efficacy in a setting more similar to real clinical practice.

Clinical practice

This review on manual thoracic therapy for impingement syndrome shows high methodological quality and a consistent reduction in pain across studies. However, it faces limitations like variability in treatment application and reliance on self-evaluations for outcomes. These findings suggest potential benefits of manual therapy in clinical practice but highlight the

need for further research with more controlled designs to confirm efficacy.

Conclusion

This review indicates that manual thoracic therapy may reduce pain in impingement syndrome, but due to methodological limitations and variability in treatment application, further research is needed to confirm its efficacy in clinical practice.

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Declarations

Conflict of interest. R. Tedeschi, D. Platano, G. Melotto and D. Danilo declare that they have no competing interests.

For this article no studies with human participants or animals were performed by any of the authors. All studies mentioned were in accordance with the ethical standards indicated in each case.

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Wirksamkeit manueller thorakaler Therapie zur Behandlung des drohenden Impingement-Syndroms: systematische Übersicht

Hintergrund: Ein Impingement-Syndrom, eine weit verbreitete Ursache für Schulterschmerzen, führt oft zu funktionellen Einschränkungen. Häufig wird die manuelle thorakale Therapie als nichtchirurgische Intervention eingesetzt, aber ihre Wirksamkeit ist weiterhin umstritten. In der vorliegenden Übersicht werden die Auswirkungen manueller thorakaler Therapie auf die Schmerzreduktion und funktionelle Verbesserung beim Impingement-Syndrom untersucht.

Methoden: Dazu wurde eine systematische Übersicht über randomisierte klinische Studien erstellt, hierbei lag der Fokus auf Studien, in denen manuelle thorakale Therapie bei Patienten mit Impingement-Syndrom angewendet wurde. Die primären Endpunkte waren Schmerzreduktion und funktionelle Verbesserung. Mithilfe der Skala der Physiotherapy Evidence Database (PEDro-Skala) wurden die Studien in Bezug auf methodische Qualität untersucht, wobei Werte ≥ 6 eine hohe Qualität anzeigen. Ergebnisse: Die Einschlusskriterien wurden von 9 Studien erfüllt. Alle Studien wiesen eine hohe methodische Qualität auf (PEDro-Score ≥ 6). Über alle Studien hinweg war die Schmerzreduktion konsistent − mit einer Reduktion des Werts auf der Numeric Pain Rating Scale (NPRS) von 0,6 auf 1,5 Punkte unmittelbar nach Behandlung und von bis zu 3,2 Punkten bei der Nachuntersuchung. Die funktionelle Verbesserung war in einigen Studien statistisch signifikant. Jedoch wiesen die Ergebnisse nur eine begrenzte Homogenität auf, und in der Mehrzahl der Studien wurden keine wesentlichen Unterschiede zwischen den Interventions- und den Placebogruppen

Schlussfolgerung: Den Ergebnissen der vorliegenden Übersicht zufolge kann manuelle thorakale Therapie zu einer Schmerzreduktion bei einem Impingement-Syndrom führen, auch bestehen einige Hinweise auf eine funktionelle Verbesserung. Allerdings zeigen die Variabilität bei den Techniken der manuellen Therapie und die Limitationen bei der wissenschaftlichen Methodik, dass Bedarf an weiteren kontrollierten Studien besteht. Diese Ergebnisse unterstreichen das Potenzial der manuellen Therapie als einer ergänzenden Behandlung, aber auch die Notwendigkeit robusterer klinischer Studien, um ihre Wirksamkeit in der klinischen Anwendung vollständig zu ermitteln.

Schlüsselwörter

 $Impingement-Syndrom \cdot Manuelle\ thorakale\ Therapie \cdot Schmerzreduktion \cdot Schulterfunktionsstörrung \cdot Muskuloskeletale\ Rehabilitation$