

ORIGINAL ARTICLE

Physical therapy in tension-type headache: a systematic review of randomized controlled trials

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ABSTRACT

Background: Tension-type headache (TTH) is the most prevalent neurological disorder globally, affecting over 2 billion individuals and posing a substantial public health and socioeconomic burden. Although pharmacological treatments, such as non-steroidal anti-inflammatory drugs, offer symptomatic relief, concerns regarding long-term efficacy and medication overuse have driven interest in physical therapy as an alternative or adjunctive intervention.

Methods: In line with the Preferred Reporting Items for Systematic Reviews and Meta-Analyses guidelines, a systematic search from inception to December 2024 was conducted in the following electronic databases: PubMed, Google Scholar, Web of Science, the Cochrane Library, and DOAJ. Randomized controlled trials investigating the effect of physiotherapy on TTH were included.

Results: Fifty-one randomized controlled trials involving 3,404 participants were included in this review. The studies investigated various physical therapy interventions, including manual therapy, therapeutic exercise, and multimodal approaches. The majority of the included studies demonstrated statistically significant improvements in headache intensity, frequency, and duration compared to control groups. Adverse events, when reported, were minor and transient. No serious complications were associated with the physical therapy interventions.

Conclusion: Physical therapy interventions can be safe with minimal adverse effects in managing patients with TTH. Various physical therapy interventions are effective in reducing pain intensity, frequency, and duration of TTH, both chronic and episodic. However, the lack of long-term outcome data highlights the need for future high-quality randomized controlled trials with extended follow-up.

Keywords: Headache, tension-type headache, physical therapy, physiotherapy, manual therapy.

Introduction

Tension-type headache (TTH) is the most prevalent neurological disorder worldwide, affecting over 2 billion individuals [1]. It ranks as the second leading cause of chronic disease and disability, underscoring its significant public health burden [2,3]. Beyond its clinical impact, TTH imposes considerable socioeconomic consequences, as it contributes to reducing work efficiency due to absenteeism and presenteeism, ultimately diminishing productivity [2].

Epidemiological data from the Global Burden of Disease study confirm the widespread prevalence of headache

disorders. However, methodological discrepancies across studies have resulted in substantial variability

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Received: 03 April 2026 | **Revised (1):** 31 May 2026 |

Accepted: 03 June 2026



in prevalence estimates, creating uncertainty regarding regional differences and trends over time [3]. Given the high global burden of TTH, understanding its underlying pathophysiology and optimizing treatment approaches are critical to improving patient outcomes.

TTH is a multifactorial disorder involving cranial and cervical muscle tension, psychological stress, and central sensitization [4]. Among individuals with chronic headache, sleep disturbances such as insomnia are frequently reported, with studies suggesting a bidirectional relationship between sleep dysfunction and headache severity [5]. The complex interplay of musculoskeletal, neurological, and behavioral factors in TTH necessitates a comprehensive approach to treatment that goes beyond symptomatic relief.

The primary pharmacological treatment for TTH involves simple analgesics and non-steroidal anti-inflammatory drugs, with ibuprofen (400 mg) and aspirin (1,000 mg) being widely recommended due to their efficacy and safety profile. However, reliance on pharmacotherapy raises concerns regarding medication overuse headache, potential side effects, and limited effectiveness in chronic cases. Consequently, there has been increasing interest in non-pharmacological interventions, particularly physical therapy, as an alternative or adjunct to pharmacological management.

Unlike pharmacological treatments that primarily provide temporary symptomatic relief, physical therapy directly addresses the musculoskeletal dysfunctions contributing to TTH. Manual therapy techniques, such as trigger point release therapy, suboccipital inhibitory pressure, and instrument-assisted soft tissue mobilization, aim to reduce muscle tension, restore cervical mobility, and alleviate myofascial pain. Acupuncture-based therapies, including dry needling, target neuromuscular sensitization and pain modulation. In addition, structured exercise programs, such as aerobic exercise and deep breathing exercises, enhance muscular flexibility and reduce physiological stress responses.

Despite promising findings, the efficacy of physical therapy for TTH remains uncertain due to variability in treatment protocols, differences in outcome measures, and a lack of long-term follow-up studies. Furthermore, comparative analyses assessing the relative effectiveness of different physical therapy techniques are limited. Addressing these gaps is essential to establishing evidence-based guidelines for integrating physical therapy into standard TTH management.

In this study, we aim to evaluate the effectiveness of physical therapy in reducing headache intensity, frequency, and duration in adults diagnosed with TTH.

Methods

Search strategy

This review followed the Preferred Reporting Items for Systematic Review and Meta-Analysis Statement (PRISMA 2020) Page et al. [6]. A systematic search was performed in several databases, including PubMed, Google Scholar, Web of Science, the Cochrane Library,

and DOAJ, using a combination of keywords (tension-type headache OR tension headache OR TTH) AND (physical therapy OR physiotherapy OR physical treatment OR physical exercises OR physical therapy interventions OR physical therapy modalities). This review was prospectively registered in the International Prospective Register of Systematic Reviews (PROSPERO; registration ID: CRD42024561923).

Study selection and eligibility criteria

Two independent reviewers performed a literature search in several databases from inception to December 2024. The search was limited to the English language, and only studies with primary data were included. The combination of keywords used was: (tension-type headache OR tension headache OR TTH) AND (physical therapy OR physiotherapy OR physical treatment OR physical exercises OR physical therapy interventions OR physical therapy modalities). The studies have been evaluated using a two-level assessment approach. Level 1 screening included titles and abstracts of retrieved records. The complete text of all potentially eligible citations was obtained and reviewed for final eligibility. All studies were screened based on the specified eligibility criteria outlined below. Studies that met the following criteria were included in this meta-analysis: (i) studies published without time frame limitations; (ii) studies in the English language; (iii) the review included studies with adult patients above 18 who were diagnosed with TTH; and (iv) the included studies reported outcomes of interest relevant to the clinical questions and included randomized controlled trials (RCTs) only.

Data extraction

Two independent reviewers extracted data from acceptable studies, including authors, study design, mean age, sex, sample size, number of sessions, and follow-up period, into a tested Excel spreadsheet. The two examiners cross-checked and verified the extraction sheet to eliminate errors and ensure the data's integrity.

Bias assessment

We performed a quality assessment for 51 studies. Following the ROB-2 Quality Assessment Tool guidelines Sterne et al. [7], 23 studies were considered to have good quality (low risk of bias). However, 28 studies were considered to have moderate quality (moderate risk of bias) due to some concerns in different domains, like the randomization process, deviations from the intended intervention, and missing data.

Results

Study selection and characteristics

The initial search yielded 1,923 potential articles. After screening titles and abstracts, 1,630 full-text articles were assessed for eligibility. Ultimately, 51 randomized controlled trials met the inclusion criteria (Figure 1). These studies included a total of 3,404 participants, aged between 20 and 55 years (Table 1). The sample consisted

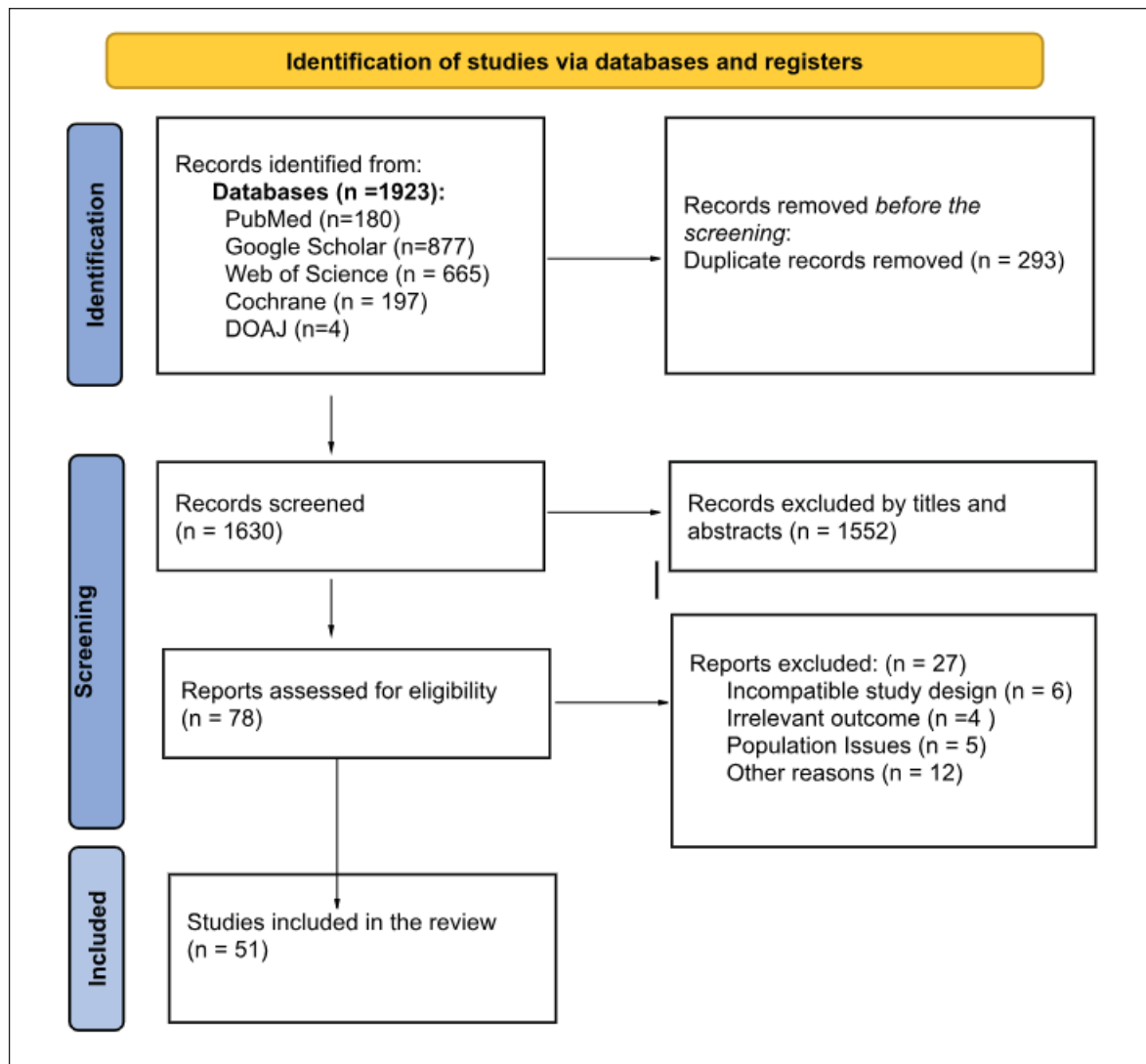


Figure 1. PRISMA flow diagram of study selection for the included studies in the systematic review.

predominantly of females (approximately 70% of participants) diagnosed with either chronic tension-type headache or episodic tension-type headache (Table 2).

Risk of bias assessment

The methodological quality of the included studies varied. Random sequence generation and allocation concealment were adequately reported in approximately 60% of the trials. Blinding of participants and therapists was a common challenge due to the nature of physical therapy interventions; however, blinding of outcome assessors was maintained in the majority of high-quality studies (Figure 2).

Efficacy of interventions

The included studies utilized a wide range of physical therapy interventions. Manual therapy (including suboccipital release and spinal manipulation) was the most common modality, appearing in 15 studies, followed by therapeutic exercises (craniocervical strengthening and aerobic training) in 12 studies. Other

interventions included acupuncture, dry needling, and electrotherapy. Multimodal physical therapy programs, combining manual techniques with exercise, were evaluated in 14 studies. Across the 51 studies, physical therapy consistently resulted in significant reductions in headache frequency, intensity, and duration compared to control or placebo groups. Manual therapy techniques showed immediate improvements in pain pressure thresholds, while therapeutic exercises demonstrated long-term benefits, particularly in reducing headache frequency (Table 3).

Adverse events

Safety data were explicitly reported in 38 of the 51 studies. Generally, physical therapy interventions were well-tolerated. Reported adverse events were mild and transient, including post-treatment muscle soreness, temporary fatigue, and minor bruising (primarily associated with dry needling). No serious or life-threatening adverse events were recorded in any of the included trials (Table 4).

Table 1. The studies' characteristics.

Author name, year, country	Study design	Sample size			Age			Sex, Male N (%)			Sex, Female N (%)			BMI (kg/m ²)		
		Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total
Danapong et al. [8]	RCT	30	30	60	N/M	N/M	49.75 ± 10.93	4	1	5	26	29	55	N/M	N/M	N/M
Glidir et al. [9]	Double-blind, parallel-group RCT	80	80	160	36.7 ± 7.6 years	36.0 ± 8.3 years	N/M	N/M	N/M	N/M	41 (51.3%)	44 (55%)	85	N/M	N/M	N/M
Endres et al. [10]	Multicenter, patient- and observer-blinded RCT	209	200	409	39.2 ± 11.4 years	38.9 ± 12.2 years	N/M	N/M	N/M	N/M	163 (78%)	158 (79%)	321	24 (21–27)	24 (22–26)	N/M
Mohamadi et al. [11]	RCT	13	13	26	39 ± 11 years	38 ± 9 years	N/M	2	3	5	11	10	21	N/M	N/M	N/M
Soderberg et al. [12]	RCT	27	28 and Group3: 25	80	40.5 ± 10.4 years	43.1 ± 8.8 years and Group3: 40.9 ± 8.3 years	N/M	6	7 and Group3: 6	19	21	21 and Group3: 19	61	N/M	N/M	N/M
Kwon and Yoon [13]	RCT	11	11 and Group3: 11	33	40.73 ± 7.82 years	49.27 ± 11.37 years and Group3: 41.82 ± 7.50 years	N/M	5	6 and Group3: 5	16	6	5 and Group3: 6	17	N/M	N/M	N/M
Aslam et al. [14]	RCT	16	15	31	35.81 ± 9.30 years	30.73 ± 7.99 years	N/M	N/M	N/M	N/M	N/M	N/M	N/M	24.11 ± 3.49	22.43 ± 3.36	N/M
Espi-López et al. [15]	Factorial, single-blinded, RCT	19	19	76 (19 per group across four groups)	Mean 43.74 years (SD 13.73)	Mean 41.58 years (SD 10.02)	39.9 years (SD 10.9)	2	8	14	17	11	28	N/M	N/M	N/M
Kamali et al. [16]	Parallel single-blind RCT	20	20	40	Mean 37.45 years (SD 12.57)	Mean 33.70 years (SD 9.94)	N/M	4	1	5	16	19	35	N/M	N/M	N/M
Saad et al. [17]	Single-blind RCT	15	15	30	Mean 36.67 years (SD 4.18)	Mean 33.73 years (SD 3.26)	N/M	N/M	N/M	N/M	N/M	N/M	N/M	Mean 24.31 kg/m ² (SD 1.46)	Mean 24.65 kg/m ² (SD 1.04)	N/M
Espi-López et al. [18]	RCT	19	19	76 (62 women, 14 men)	39.9 ± 10.9 years	N/M	N/M	N/M	N/M	14	N/M	N/M	62	N/M	N/M	N/M

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Author name, year, country	Study design	Sample size			Age			Sex, Male N (%)			Sex, Female N (%)			BMI (kg/m ²)		
		Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total
Ferragut-Garcias et al. [19]	Double-blind, placebo-based RCT	Group A (Placebo): 24 Group B (Soft tissue): 23 (Combined): 25	Group C (Neural mobilization): 25 Group D (Combined): 25	97	Group A: 40.5 ± 12.0 years Group B: 38.1 ± 10.9 years	Group C: 39.4 ± 11.0 years Group D: 40.8 ± 12.1 years	39.7 ± 11.5 years	Group A: 4 (16.7%) Group B: 6 (26.0%)	Group C: 5 (20.0%) Group D: 4 (16.0%)	19 (19.6%)	Group A: 20 (83.3%) Group B: 17 (74.0%)	Group C: 20 (80.0%) Group D: 21 (84.0%)	78 (80.4%)	Group A: 25.3 ± 3.0 kg/m ² Group B: 24.7 ± 3.4 kg/m ²	Group C: 25.1 ± 3.3 kg/m ² Group D: 24.9 ± 3.0 kg/m ²	25.0 ± 3.2 kg/m ²
Ajimsha [20]	Single-blinded RCT	Group A (DT-MFR): 22	Group B (IDT-MFR): 22 Group C (Control): 12	56	Group A: 43.7 ± 5.6 years	Group B: 44.7 ± 5.2 years Group C: 43.0 ± 5.4 years	43.8 ± 5.4 years	Group A: 7	Group B: 8 Group C: 5	20	Group A: 15	Group B: 14 Group C: 7	36	Group A: 24.7 ± 5.9 kg/m ²	Group B: 24.8 ± 6.2 kg/m ² Group C: 24.2 ± 5.0 kg/m ²	24.6 ± 5.7 kg/m ²
Castien et al. [21]	Pragmatic, multicentre, RCT clinical trial	41	41	82	20–63	20–59	N/M	N/M	N/M	9	N/M	N/M	32	N/M	N/M	N/M
Hossainifar et al. [22]	RCT	15	15	30	25.06 ± 7.64	29.33 ± 11.63	N/M	N/M	N/M	N/M	15	15	30	N/M	N/M	N/M
Gopichandran et al. [23]	Prospective RCT	84	85	169	44.21 ± 10.7	46.167 ± 11.12	N/M	39 (46.5)	34 (40)	N/M	45 (53.5)	51 (60)	N/M	N/M	N/M	N/M
Shafiq et al. [24]	RCT	15	15	30	mean age: 20.27 ± 1.53 years	mean age: 21.27 ± 1.79 years	N/M	5	6	11	10	9	19	N/M	N/M	N/M
Pérez-Llanes et al. [25]	RCT	13	12	25	43.3 years	46.2 years	N/M	3	1	4	10	11	21	Mean BMI: 24.7 kg/m ²	Mean BMI: 26.3 kg/m ²	N/M
Cabanillas-Barea et al. [26]	RCT	43	43	86	37.25 years (±15.41)	39.49 years (±16.26)	38.35 years (±15.78)	12	12	24	31	31	62	N/M	N/M	N/M
Monti-Ballano et al. [27]	Single-blinded RCT	16	16	32	mean age 31.75 ± 10.75 years	mean age 41.44 ± 14.68 years	mean age 39.09 ± 12.88 years	N/M	N/M	8	N/M	N/M	24	N/M	N/M	N/M
Álvarez-Melcón et al. [23]	RCT	76	76	152	mean age 20.23 (SD ± 2.50)	mean age 20.62 (SD ± 2.21)	20.42 years (SD ± 2.36)	26	42	68	50	34	84	N/M	N/M	N/M
Park [28]	RCT	18	17	35	39.6 ± 15.9 years	41.0 ± 9.4 years	N/M	N/M	N/M	N/M	N/M	N/M	N/M	23.8 ± 2.6 kg/m ²	25.2 ± 4.2 kg/m ²	N/M

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Author name, year, country	Study design	Sample size			Age			Sex, Male N (%)			Sex, Female N (%)			BMI (kg/m ²)		
		Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total
Chatchawan et al. [29]	Prospective, parallel-group RCT	36	36	72	mean age: 27.39 ± 6.73 years	mean age: 27.36 ± 9.55 years	total mean age: 27.37 ± 8.37 years	10	7	17	26	29	55	22.48 ± 5.33 kg/m ²	22.17 ± 7.55 kg/m ²	22.33 ± 6.36 kg/m ²
Choi et al. [30]	RCT	Temporomandibular group 14, cervical group 14	13	41	TMJT 35.00 ± 11.82 years, CMT 34.71 ± 7.50 years	42.62 ± 11.00 years	N/M	15	8	23	13	5	18	N/M	N/M	N/M
Martín-Vera et al. [31]	RCT	20	20	40	33.9 ± 12.2 years	40.1 ± 14.0 years	N/M	3 15%	5 25%	8 20%	17 85%	15 75%	32	24.4 ± 3.9 kg/m ²	24.2 ± 4.2 kg/m ²	N/M
Antonia (Gomez) et al. [32]	RCT	62	22	84	N/M	N/M	Mean age: 39.76 years (Range: 18-65)	N/M	N/M	16	N/M	N/M	68	N/M	N/M	N/M
Van Ettehoven and Lucas [33]	RCT	38	42	81	48.3	43.4	45.85	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M
Kanji et al. [34]	RCT	17	20	37	40.7	44.3	ns	5	3	8	12	17	29	N/M	N/M	N/M
Madsen et al. [35]	RCT	23	21	44	33	36	ns	5	6	11	18	15	33	24	24	24
Xue et al. [36]	Single-blinded, sham-controlled, crossover clinical trial	20	20	40	42.6 (1.8)	41.5 (1.9)	N/M	7	7	14	13	13	26	N/M	N/M	N/M
Moraska et al. [37]	RCT	20 for massage	21 wait list, 21 for placebo	62	32.1 ± 12.0	32.1 ± 12.0	N/M	N/M	N/M	N/M	88.2	89.5	N/M	N/M	N/M	N/M
Rinne et al. [38]	RCT	57	69	116	45.8 (8.6)	42.6 (9.7)	N/M	All female	N/M	N/M	57	69	N/M	N/M	N/M	N/M
Karst et al. [39]	RCT	34	35	69	47.9 (13.8)	48.2 (14.6)	N/M	17	14	31	17	21	38	N/M	N/M	N/M
Torelli et al. [40]	RCT	24	24	48	43.9 (24-63)	46.8 (29-59)	44.9 (24-63)	5	10	15	19	14	33	N/M	N/M	N/M
Bove and Nilsson 41]	RCT	37	38	75	37 (22-59)	38 (20-58)	38 (20-59)	15	11	26	23	26	49	N/M	N/M	N/M

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Author name, year, country	Study design	Sample size			Age			Sex, Male N (%)			Sex, Female N (%)			BMI (kg/m ²)		
		Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total
Hamed [42]	Parallel RCT	15 for each group SOES (group A); SPTP (group B);	15	45	Group A = 37.0 ± 7.1 Group B = 36.6 ± 7.2	35.6 ± 4.6	N/M	5 in group A, 6 in group B	5	16	10 IN GROUPE, 9 in group B	10	29	N/M	N/M	N/M
Karakurum et al. [43]	Randomized, double-blind study	15	15	30	28.4 ± 11.6	27.9 ± 10	N/M	all females	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M
Cabanillas-Barea et al. [44]	RCT	41	41	82	37.25 (15.41)	39.39 (16.26)	N/M	11	11	22	29	28	57	24.08 (2.17)	23.83 (1.70)	N/M
Corum et al. [45]	RCT	15 for each group (group A) and (group B);	15	45	Group A = 33.5 ± 8.2, Group B = 30.7 ± 8.0	32.5 ± 6.5	N/M	Group A = 4, Group B = 3	4	11	Group A = 9, Group B = 11	8	28	Group A = 22.1 ± 2.3, Group B = 21.9 ± 2.6	23.5 ± 3.2	N/M
Sentel and Bakar [46]	RCT	BAT (n = 20), aerobic exercise (n = 20)	20	60	42.6 ± 9.5 for BAT group, 36.20 ± 7.86 for AE Group	39.00 ± 9.53	N/M	N/M	N/M	N/M	N/M	N/M	N/M	27.31 ± 5.6 for BAT group, 27.10 ± 4.6 for AE Group	25.66 ± 3.71	N/M
Kwon and Yoon [47]	RCT	15	15	30	37.93 ± 10.32	38.27 ± 11.10	N/M	8	7	15	7	8	15	N/M	N/M	N/M
Espl-López et al. [48]	Single-blinded, RCT	52	52	104	37.69 ± 10.64	40.47 ± 11.33	N/M	10	13	23	41	38	79	N/M	N/M	N/M
Azhdari et al. [49]	Parallel design RCT	12	12	24	43.16	41.41	N/M	5	4	9	7	8	15	N/M	N/M	N/M
Georgoudis et al. [50]	A single-blind, prospective, multicentre, RCT	20	24	44	54.8 ± 14.7	43.0 ± 6.5	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M
Georgoudis et al. [51]	A single-blind multicenter-RCT	20	24	44	54.8 ± 14.7	43.0 ± 6.5	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M
Cho et al. [52]	RCT	SMI group = 15, SMIE = 15	15	45	SMI group = 36.60 ± 8.33, SMIE = 36.00 ± 8.47	39.07 ± 10.99	N/M	SMI group = 3, SMIE = 4	3	10	SMI group = 11, SMIE = 10	10	31	N/M	N/M	N/M
Ghanbari et al. [53]	RCT	15	15	30	37.66 ± 8.6	36.26 ± 7.46	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M
Ebneshahidi et al. [54]	single blind, placebo RCT	25	25	50	33 (25-52)	38.6 (26-54)	N/M	5	5	10	20	20	40	N/M	N/M	N/M

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Author name, year, country	Study design	Sample size			Age			Sex, Male N (%)			Sex, Female N (%)			BMI (kg/m ²)		
		Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total
Abaschian and Mansoursohani [55]	Single-blind, parallel-group RCT	12	12	24	37.4 ± 9.24	32.55 ± 7.65	N/M	N/M	N/M	N/M	N/M	N/M	N/M	25.27 ± 4.36	25.12 ± 4.55	N/M
Berggreen et al. [56]	RCT	20	19	39	38.8 (13.7)	42.3 (10.2)	N/M	ALL females	N/M	N/M	20	19	39	N/M	N/M	N/M
Espe-López et al. [57]	Double-blind RCT	62 divided to three sub groups	22	84	Treatment SI = 42.75 (18-65), OAA = 34 (18-50), SI and OAA = 40.70 (24-62)	41.95 (18-62)	39.76 (1-65)	Treatment SI = 5, OAA = 2, SI and OAA = 4	4	15	Treatment SI = 17, OAA = 20, SI and OAA = 18	18	68	N/M	N/M	N/M

Table 2. Baseline tension headache features of the participants.

Author, year, country	Type of headache		The pain intensity			Duration of tension-type headache			Frequency of tension-type headache			Mean No. Of analgesics		
	Chronic TTH #	Episodic TTH #	Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total
Danapong et al. [8]	55	N/M	6.3 ± -1.2	6.06 ± -0.94	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M
Gildir et al. [9]	160	N/M	4.5 ± 1.0 cm	4.6 ± 1.2 cm	N/M	3.9 ± 0.7 hours/day	3.9 ± 0.7 hours/day	N/M	18.5 ± 2.7 days/month	18.0 ± 2.4 days/month	N/M	N/M	N/M	N/M
Endres et al. [10]	93 (Group 1), 94 (Group 2)	116 (Group 1), 106 (Group 2)	68.3 ± 12.1	67.5 ± 12.5	N/M	11.2 ± 10.3 years	11.7 ± 10.7 years	N/M	14 (12-18) days/4 weeks	14 (12-19) days/4 weeks	N/M	N/M	N/M	N/M
Mohamadi et al. [11]	All participants	N/M	7.46 ± 1.80	6.53 ± 1.89	N/M	N/M	N/M	N/M	18.30 ± 6.29 days/month	16.69 ± 6.14 days/month	N/M	N/M	N/M	N/M
Soderberg et al. [12]	80	N/M	N/M	N/M	N/M	N/M	N/M	N/M	18.6 ± 6.7 days/month	20.5 ± 6.2 days/month and Group3: 18.2 ± 5.7 days/month	N/M	N/M	N/M	N/M

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Author, year, country	Type of headache		The pain intensity			Duration of tension-type headache			Frequency of tension-type headache			Mean No. Of analgesics		
	Chronic TTH#	Episodic TTH #	Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total
Kwon and Yoon [13]	33	N/M	35.36 ± 0.92	35.18 ± 1.25 and Group3: 35.36 ± 1.74	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M
Aslam et al. [14]	31	N/M	6.62 ± 1.58	6.73 ± 1.53	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M
Espí-López et al. [15]	17	21	6.5	N/M	6.5	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M
Kamali et al. [16]	N/M	N/M	Median 8.00	Median 9.50	N/M	N/M	N/M	N/M	Median 5 days/week	Median 7 days/week	N/M	N/M	N/M	N/M
Saad (Namb) et al. [17]	30	N/M	Mean 6.26 (SD 0.86)	Mean 6.45 (SD 0.96)	N/M	N/M	N/M	N/M	Mean 4.73 days/week (SD 1.03)	Mean 4.60 days/week (SD 0.91)	N/M	N/M	N/M	N/M
Espí-López et al. [18]	40.8% of 76	59.2% of 76	N/M	N/M	N/M	N/M	N/M	N/M	2.63 (pre-treatment frequency score for the control group)		N/M	N/M	N/M	N/M
Ferragut-Garcías et al. [19]	41.7% in Group A, 34.8% in Group B, 44.0% in Group C, 48.0% in Group D	58.3% in Group A, 65.2% in Group B, 56.0% in Group C, 52.0% in Group D	Group A: 5.6 ± 1.1 Group B: 4.4 ± 1.1	Group C: 5.7 ± 0.8 Group D: 5.1 ± 1.0	N/M	N/M	N/M	N/M	Group A: 7.2 ± 2.7 days/15 days Group B: 8.6 ± 2.3 days/15 days	Group C: 7.9 ± 2.7 days/15 days Group D: 8.0 ± 2.6 days/15 days	N/M	N/M	N/M	N/M
Ajimsha [20]	9 (16.1%)	47 (83.9%)	N/M	N/M	N/M	Group A: 7.1 ± 7.2 years Group B: 7.3 ± 8.1 years Group C: 7.7 ± 7.7 years		N/M	Group A: 12.0 ± 2.8 days/4 weeks Group C: 12.0 ± 2.5 days/4 weeks	Group B: 12.4 ± 2.8 days/4 weeks	N/M	N/M	N/M	N/M
Castien et al. [21]	83	0	7.5 (1.7) P value not reported	7.7 (1.6) P value not reported	N/M	12.8 (8.9) P value not reported	13.0 (9.1) P value not reported	N/M	23.7 (6.8) P value not reported	24.0 (7.0) P value not reported	N/M	N/M	N/M	2
Hosseinfar et al. [22]	30	0	4.06 ± 0.69 P value = 1.000	4.06 ± 1.27 P value = 1.000	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M
Gopichandran et al. [23]	169	0	6.7 ± 1.05	6.52 ± 1.2	N/M	<1 year = 2 (2.4) > 1-2 year = 38 (45.2) > 2-3 years = 30 (35.7) > 3 years = 14 (16.6)	<1 year = 5 (6) > 1-2 year = 32 (37.6) > 2-3 years = 30 (35.2) > 3 years = 18 (21.4)	N/M	19.44 ± 1.56 p = -0.03 (0.97)	19.43 ± 1.18 p = -0.03 (0.97)	N/M	Analgesics = 30 (35.3) Analgesics with amitriptyline and propranolol = 55 (64.7)	Analgesics 30 (35.7). Analgesics with amitriptyline and propranolol = 54 (64.3)	N/M

Continued

Author, year, country	Type of headache		The pain intensity		Duration of tension-type headache			Frequency of tension-type headache			Mean No. Of analgesics		Total
	Chronic TTH #	Episodic TTH #	Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Group 1 (Treatment)	Group 2 (Control)	
Shafiq et al. [24]	N/M	N/M	mean pain intensity: 2.53 ± 0.51	Mean pain intensity 2.53 ± 0.51	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M
Pérez-Llanes et al. [25]	25	0	4.6 (SD = 2.87)	6.9 (SD = 1.33)	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	paracetamol, NSAIDs, antidepressants, anxiolytics.
Cabanillas-Barea et al. [26]	86	0	Actual pain: 1.75 ± 1.74, usual pain: 3.42 ± 1.81, worse pain: 6.227 ± 1.64	Actual pain: 2.04 ± 1.76, usual pain: 2.99 ± 1.46, worse pain: 6.14 ± 1.71	N/M	N/M	N/M	N/M	13.28 ± 11.97 days/2 weeks	13.26 ± 12.39 days/2 weeks	NSAIDs 18, Acetaminophen and NSAIDs 8, Acetaminophen 16	NSAIDs 16, Acetaminophen and NSAIDs 7, Acetaminophen 16	N/M
Monti-Ballano et al. [27]	CTTH and ETTH		N/M	N/M	N/M	N/M	N/M	N/M	13.69 ± 9.00 days/months	13.19 ± 11.55 days/months	18.94 ± 28.21 doses/months	39.81 ± 69.98 doses/months	29.38 ± 53.73 doses/months
Álvarez-Melcón et al. [58]	52	100	Mean intensity 5.82 (1.26)	Mean intensity 5.57 (1.32)	Total mean intensity 5.695	Mean duration 6.35 hours/day	Mean duration 5.94 hours/day	Total mean duration 6.15 hours/day	Mean frequency 12.96 days in 4 weeks	Mean frequency 12.71 days in 4 weeks	Pre-treatment = 7.43 days in 4 weeks, post-treatment 5.13 days in 4 weeks, follow-up 4.19 days in 4 weeks	Pre-treatment 7.46 days in 4 weeks, post-treatment 6.13 days in 4 weeks, follow-up 5.31 days in 4 weeks	N/M
Park [28]	Only chronic	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M
Chatchawan et al. [29]	42	0	5.54 ± 2.16	4.66 ± 2.40	5.10 ± 2.23	8.28 ± 13.81 hours	4.65 ± 4.67 hours	N/M	16.26 ± 2.02 times/month	16.35 ± 6.68 times/month	25	25	50
Choi et al. [38]	41	0	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M
Martín-Vera et al. [31]	40	0	6.9 ± 1.3	7.1 ± 1.6	N/M	18.3 ± 7.5 hours/month	14.0 ± 8.3 hours/month	N/M	10.1 ± 9.5 days/month	10.5 ± 7.2 days/month	N/M	N/M	N/M
Gómez-Conesa et al. [32]	0.571	0.429	N/M	N/M	6.49 (SD: 1.69)	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M
van Ettekoven et al. [33]	ETTH and CTTH		5.72	5.86	N/M	8.1 hours per day	8.0 hours per day	N/M	5.49	4.86	1.13 days	0.98 days	ns
Kanji et al. [34]	All participants	N/A	4.3	3.5	ns	8.3 hours per day	6.6 hours per day	N/M	23.6	23.6	N/M	N/M	N/M
Madsen et al. [35]	N/M	N/M	4.5	3.6	NS	252 hours per month (mean)	198 hours per month (mean)	N/M	19 days	17 days	3.5 days	4.7 days	N/M

Continued

Author, year, country	Type of headache		The pain intensity			Duration of tension-type headache			Frequency of tension-type headache			Mean No. Of analgesics		
	Chronic TTH #	Episodic TTH #	Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total
Xue et al. [36]	11 in group A, 11 in group B	9 in each group	45.2 (2.4)	49.3 (2.4)	N/M	47.7 (7.4)	40.8 (4.8)	N/M	13.8 (1.7)	12.5 (1.6)	N/M	N/M	N/M	N/M
Moraska et al. [37]	64.7 % in massage group, 52.6% in placebo group	35.3% in massage group, 47.4% in the placebo group	31.4 ± 2.69	33.3 ± 2.52	N/M	3.15 ± 0.43	3.20 ± 0.55	N/M	3.72 ± 0.23	3.81 ± 0.21	N/M	Dose/week = 2.97 ± 0.82	Dose/week = 1.75 ± 0.76	N/M
Rinne et al. [38]	N/M	N/M	4.7 (95% confidence intervals, CI 4.4 to 5.0)	4.8 (4.5 to 5.1)	N/M	30.8 (95% CI 24.7 to 36.9) hours/ week in the exercise group	30.5 (23.9 to 37.1) in the control group	N/M	4.5 (95% CI 3.9 to 5.1) in the exercise group	and 4.4 (3.6 to 5.1) in the control group	N/M	N/M	N/M	N/M
Karst et al. [39]	25 in treatment, 22 in placebo	9 in treatment group, 12 in placebo group	6.4 (2.0)	6.3 (1.9)	N/M	N/M	N/M	N/M	21.1 (10.2)	20.5 (10.3)	N/M	Analgesics/month 9.0 (11.1)	Analgesics/month 15.6 (32.4)	N/M
Torelli et al.[40]	8 in treatment group, 16 in control group	16 in treatment group, 8 in control	1.6 (1-3)	1.7 (1-3)	N/M	9.8 (7-16)	12.0 (5-22)	N/M	14.5 (11-18)	18.1 (13-22)	N/M	19.7 (5-33)	23.6 (6-32)	N/M
Bove and Nilsson [41]	N/M	N/M	Per day (95% CI) 37/100 (33-41/100)	37/100 (33-41/100)	38/100 (35-41/100)	per day (95% CI) 2.8 (2.1-3.5)	3.4 (2.4-4.4)	3.1 (2.5-3.7)	N/M	N/M	N/M	Per day (95% CI) 0.66 (0.49-0.83)	0.82 (0.50-1.14)	0.74 (0.56-0.92)
Hamed [42]	N/M	N/M	group A = 70 ± 1, Group b = 71 ± 12	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M
Karakurum et al. [43]	20 (66.7%)	8 (26.7%) episodic and 2 (6.7%) had episodic TTH in combination with migraine without aura.	N/M	N/M	N/M	N/M	N/M	N/M	29.6/month	25.2/ month	N/M	N/M	N/M	N/M
Cabanillas-Barea et al. [44]	N/M	N/M	1.75 (1.74)	2.04 (1.77)	N/M	N/M	N/M	N/M	13.28 (11.90)	13.26 (12.29)	14 (35%) for Acetaminophen, 18 (45%) for NSAIDs	16 (41%) for Acetaminophen, 16 (41%) for NSAIDs	N/M	

Continued

Author, year, country	Type of headache		The pain intensity			Duration of tension-type headache			Frequency of tension-type headache			Mean No. Of analgesics		
	Chronic TTH #	Episodic TTH #	Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total	Group 1 (Treatment)	Group 2 (Control)	Total
Corum et al. [45]	3 in group A, 5 Group B, 4 Group C	9 in group A, 10 Group B, 8 Group C	Group A = 5.6 ± 1.1, Group B = 5.8 ± 1.1	5.8 ± 1.2	N/M	N/M	N/M	N/M	Group A (Day/2 week) = 5.4 ± 2.3, Group B (Day/2 week) = 5.6 ± 2.5	Day/2 week = 5.8 ± 2.7	N/M	N/M	N/M	N/M
Sertel and Bakar [46]	N/M	N/M	BAT = 6.15 ± 0.74, AE = 6.1 ± 1.02	5.90 ± 0.71	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M
Kwon et al. [47]	N/M	N/M	6.27 ± 1.38	5.33 ± 1.17	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M
Espi-López et al. [48]	24 in control group, 25 in treatment group	27 (control group, 26 in treatment group)	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M
Azhdari et al. [49]	N/M	N/M	7.37 (2.01)	6.70 (2.37)	N/M	344.18 (333.39)	279.05 (165.66)	N/M	4.91 (2.39)	5.25 (2.05)	N/M	N/M	N/M	N/M
Georgoudis et al. [50]	N/M	N/M	6.5 ± 2.5	7.8 ± 1.8	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M
Georgoudis et al. [51]	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	8.1 ± 2.8	8.2 ± 3.6	N/M	N/M	N/M	N/M
Cho et al. [52]	N/M	N/M	N/M	N/M	N/M	SMI group = 3.63 ± 1.67 (years), SMIEx = 3.29 ± 2.22 (years)	3.28 ± 1.43	N/M	N/M	N/M	N/M	N/M	N/M	N/M
Ghanbari et al. [53]	N/M	N/M	5.80 ± 1.75	6.03 ± 0.99	N/M	6.42 ± 5.70	5.37 ± 3.47	N/M	12.40 ± 2.22	11.13 ± 1.99	N/M	N/M	N/M	N/M
Ebneshahidi et al. [54]	N/M	N/M	10 (3.0)	10 (1.0)	N/M	10 (4.0)	8 (4.5)	N/M	N/M	N/M	N/M	N/M	N/M	N/M
Abaschian and Mansoursohani [55]	N/M	All cases (24)	6.08 ± 1.32	4.83 ± 1.37	N/M	N/M	N/M	N/M	11.44 ± 3.77	11.06 ± 3.36	N/M	N/M	N/M	N/M
Berggreen et al. [56]	N/M	N/M	28.0 (15.9)	26.6 (12.6)	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M
Espi-López et al. [57]	N/M	N/M	Treatment SI = 4.79 (2.26), OAA = 5.12 (1.95), SI and OAA = 4.80 (1.68)	5.24 (1.80)	N/M	N/M	N/M	N/M	Treatment SI = 3.25 (2.29), OAA = 2.90 (1.86), SI and OAA = 55.67 (7.74)	3.24 (1.57)	N/M	N/M	N/M	N/M

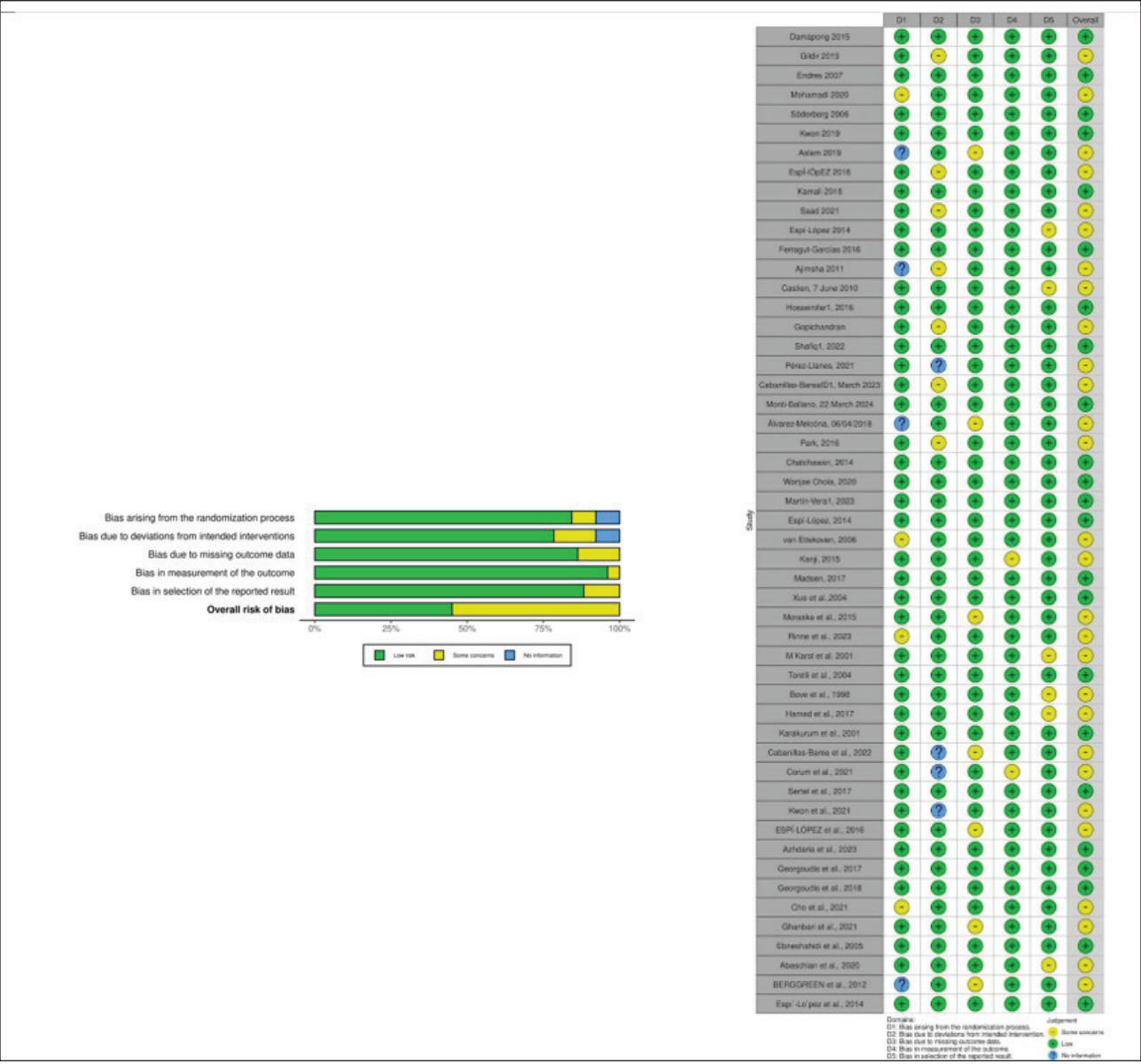


Figure 2. Risk of bias summary using ROB-2 quality assessment tool for the included randomized clinical trials.

Discussion

This systematic review synthesized evidence from 51 randomized controlled trials to evaluate the effectiveness of physical therapy for TTH. The comprehensive analysis of over 3,400 participants confirms that physical therapy is a robust, non-pharmacological treatment option.

The results align with the multifactorial pathophysiology of TTH. Our analysis supports the findings of Repiso-Guardeño et al. [59], whose systematic review highlighted the efficacy of manual therapy in reducing central sensitization. Similarly, Cumplido-Trasmonte et al. [60] emphasized that manual therapy techniques are particularly effective for adults with chronic TTH when applied to the cranio-cervical region.

Regarding multimodal care, our review suggests that combining passive therapies with active exercise yields superior outcomes. This is consistent with the recent meta-analysis by Onan et al. [61], which concluded

that physiotherapy approaches are most effective when tailored to address both muscular tension and functional disability. Furthermore, Qin et al. [62] noted that complementary strategies, including acupuncture, significantly reduce headache frequency compared to routine care.

Safety remains a key advantage of physical therapy. Krøll et al. [63] reported that adverse events in non-pharmacological interventions are rare and mild. Our review corroborates this, as the included RCTs (Appendix A) reported mostly transient muscle soreness, reinforcing physical therapy as a safe alternative for patients at risk of medication-overuse headaches.

Despite the heterogeneity in treatment protocols—ranging from single sessions to 12-week programs—the consistent positive outcomes across diverse populations strengthen the external validity of these findings. However, as noted by Jung et al. [64], the quality of evidence is often limited

Table 3. Characteristics of the physiotherapies conducted in TTH patients.

Author, year, country	Type of physiotherapy used		Home based program	Conducted by qualified physiotherapist	Duration of physiotherapy	Number of sessions	Follow up period	Withdraw (N)			Withdrawal reasons
	Group 1 (Treatment)	Group 2 (Control)						Group 1 (Treatment)	Group 2 (Control)	Total	
Damapong et al. [8]	Court type traditional Thai massage	Amitriptyline	N/M	Yes	45 minutes per session, twice a week for 4 weeks	8	2 weeks	None	None	None	N/A
Gildir et al. [9]	Trigger point dry needling (DN)	Sham dry needling (SDN)	No	Yes	3 sessions/week for 2 weeks	6 sessions (for each group)	1 month	0	1	1	1 patient in the control group used tricyclic antidepressants
Endres et al. [10]	Verum acupuncture	Sham acupuncture	No	Yes	10 sessions over 6 weeks (with an additional 5 sessions for partial response)	10 (or up to 15 for partial response)	6 months	5	6	11	N/M
Mohamadi et al. [11]	Positional release technique (PRT)	Ibuprofen 200mg	No	Yes	10 sessions over 5 weeks (2 sessions per week)	10 sessions	5 weeks	2	3	5	Ineligibility (detected by MRI) or discomfort with the procedures
Soderberg et al. [12]	Acupuncture	Physical training (strength and endurance) and Group3: Relaxation training	No	Yes	12 weeks	24 sessions	6 months	1	2 and Group3: 1	4	N/M
Kwon and Yoon [13]	Temporomandibular joint and cervical vertebra treatment	Temporomandibular joint treatment and Group3: Cervical vertebra treatment	NO	Yes	4 weeks, 3 times per week, 50 minutes per session	12 sessions	N/M	N/M	N/M	N/M	N/M
Aslam et al. [14]	Trigger point release therapy (20 minutes)	Heat therapy, TENS, and stretching exercises	No	Yes	3 sessions over 1 week	3	N/M	N/M	N/M	N/M	N/M
Espi-López et al. [15]	Suboccipital inhibitory pressure (SI), suboccipital manipulation (SM), combination	None	N/M	Yes	4 weeks	4 sessions	1 month	None	None	None	None
Kamali et al. [16]	Dry needling versus friction massage at myofascial trigger points	Manual circular pressure instead of needles	N/M	N/M	1 week	3 sessions	48 hours after the last session	N/M	N/M	4	1 withdrew due to needle phobia, 3 did not attend the follow-up
Saad (Nambi) et al. [17]	Instrument-assisted soft tissue mobilization (IASTM)	Conventional physical therapy	N/M	Yes	4 weeks	12 sessions	Post-treatment evaluation after 4 weeks	1	1	2	N/M

Continued

Author, year, country	Type of physiotherapy used		Home based program	Conducted by qualified physiotherapist	Duration of physiotherapy	Number of sessions	Follow up period	Withdraw (N)			Withdrawal reasons
	Group 1 (Treatment)	Group 2 (Control)						Group 1 (Treatment)	Group 2 (Control)	Total	
Espí-López et al. [18]	Suboccipital soft tissue inhibition (SI), Occiput-Atlas-Axis (OAA) manipulation, combination	None	No	Qualified physiotherapists with >10 years of experience	4 weeks	4 sessions	4 weeks	N/M	N/M	N/M	N/M
Ferragut-Garcías et al. [19]	Group A: Placebo superficial massage Group B: Soft tissue techniques	Group C: Neural mobilization techniques Group D: Combined soft tissue and neural mobilization techniques	No	Qualified physiotherapists	4 weeks	6 (15 minutes per session)	30 days	N/M	N/M	N/M	N/M
Ajimsha [20]	Group A (DT-MFR): Direct technique myofascial release (3 minutes on each side for multiple head, neck, and face regions)	Group B (IDT-MFR): Indirect technique myofascial release (10-minute stretches and pulls) Group C (Control): Slow soft stroking over the head	No	Certified myofascial release practitioners	12 weeks (2 sessions per week)	24	4 weeks post-intervention (Weeks 17-20)	Group A: 2	Group B: 2 Group C: 1	5	Not provided
Castien et al. [21]	Combination of mobilisations of the cervical and thoracic spine, exercises and postural correction	GP	yes	yes	30 minutes with a maximum of nine treatments	N/M	N/M	none	none	none	N/M
Hosseinfar et al. [22]	MFR technique and exercise therapy	No intervention	No	Yes	Each session last 45 minutes	4 times a week for 3 weeks	At least 3 months	None	None	None	N/M
Gopichandran et al. [23]	Perform PMR and deep breathing exercises	Standard routine care	Yes	Yes	20 minutes daily in the evening with an instruction booklet	Were asked to perform them for 12 weeks	2 weeks-6 weeks- 12 weeks	None	None	None	N/M
Shafiq et al. [24]	Soft tissue mobilization and trigger point release	Stretching	N/M	yes	3 weeks	9 sessions	N/M	N/M	N/M	22	N/M
Pérez-Llanes et al. [25]	Suboccipital muscle inhibition, interferential current therapy	Received standard treatment	N/M	Yes	4 weeks	8 sessions	4 weeks	0	3	3	Due to lack of adherence to the study
Cabanillas-Barea et al. [26]	Diacutaneous fibrolysis	None	N/M	Yes	1 week	3 sessions	1 month	N/M	N/M	7	4 did not complete the intervention protocol, 3 did not attend the evaluation sessions

Continued

Author, year, country	Type of physiotherapy used		Home based program	Conducted by qualified physiotherapist	Duration of physiotherapy	Number of sessions	Follow up period	Withdraw (N)			Withdrawal reasons
	Group 1 (Treatment)	Group 2 (Control)						Group 1 (Treatment)	Group 2 (Control)	Total	
Monti-Ballano et al. [27]	Dry needling	None	N/M	N/M	N/M	3 sessions	N/M	N/M	N/M	N/M	N/M
Álvarez-Melcón et al. [58]	Combination of relaxation techniques and physical therapy, Autogenic Training (AT), Cervical spine kinesiotherapy, posture correction.	Relaxation techniques only (Autogenic training)	Yes	Yes	3 weeks	28 sessions	3 months	4	4	8	N/M
Park et al. [28]	Manual therapy (relaxation approaches+self-exercise at peri-neck muscles	Conventional rehabilitation approaches	N/M	Yes	4 weeks	12 sessions	N/M	N/M	N/M	N/M	N/M
Chatchawan et al. [29]	Combined massage for 25 minutes and stretching for 5 minutes according to the pattern of royal Thai massage	Sham US application of a detuned device with circular kneading on the upper neck or upper back in the supine and side lying positions	N/M	Yes	3 weeks	9 sessions	12 weeks	6	5	11	Mild fever, mild soreness, and discomfort.
Choi et al. [30]	Temporomandibular joint therapy, Cervical manual therapy	Conservative percutaneous stimulation therapy, ultrasound therapy, and warm compress	N/M	Yes	3 weeks	3 sessions	N/M	N/M	N/M	4	N/M
Martín-Vera et al. [31]	The participants performed specific exercises for craniocervical, shoulder girdle, shoulder muscles.	Daily activities without monitoring	Yes	Yes	12 weeks	2 day in first 6 weeks, 3 days the last 6 weeks	N/M	N/M	N/M	N/M	N/M
Antonia (Gomez) et al. [32]	Manipulative treatment, manual therapy, and combined treatment	Did not receive treatment but stayed in the supine position with neutral ranges for 10 minutes	N/M	Yes	Four treatment sessions (1 session per week) were administered, with an interval of 7 days, Each session lasted for approximately 20 minutes.	4	8 weeks	2 participants	2 participants	4	Control group: 1 due to transient illness, 1 for no pain relief. treatment group: 1 due to mild cervical pain, 1 for personal reasons.

Continued

Author, year, country	Type of physiotherapy used		Home based program	Conducted by qualified physiotherapist	Duration of physiotherapy	Number of sessions	Follow up period	Withdraw (N)			Withdrawal reasons
	Group 1 (Treatment)	Group 2 (Control)						Group 1 (Treatment)	Group 2 (Control)	Total	
van Ettekov en et al. [33]	Standard physiotherapy and craniocervical training	Standard physiotherapy only	Yes, participants practiced craniocervical training at home twice daily	Yes	6 weeks	NS	Immediately after treatment and at 6 months	1	2	3	NS
Kanji et al. [34]	Sauna bathing 3 times per week for 20 minutes and education and self-directed soft tissue massage	Education and self-directed soft tissue massage	Yes (self-directed soft tissue massage)	N/M	8 weeks (sauna intervention period)	3 times a week (Group 1)	12 weeks	2	4	6	N/M
Madsen et al. [35]	Strength training (progressive with elastic resistance bands)	Ergonomics and posture correction	Yes (self-directed exercises in both groups)	Yes	10 weeks	3 times per week for the strength training group	19–22 weeks	7	9	16	N/M
Xue et al. [36]	Group A had real electroacupuncture (REA) in phase I, then SEA in phase II.	Group B reGroup B had SEA in phase I, then REA in phase II.	N/M	N/M	6 weeks	N/M	3 months	N/M	N/M	N/M	N/M
Moraska et al. [37]	Myofascial Trigger Point-focused Head and Neck Massage	Placebo (detuned ultrasound)	N/M	N/M	Over 6 weeks,	N/M	4 weeks	4	2	6	N/M
Rinne et al. [38]	Exercise (three included low-load exercises for the neck–shoulder region, and the remaining three specific strengthening exercises for the neck and upper body	Placebo-dosed treatment with 20 minutes of transcutaneous electrical nerve stimulation	N/M	N/M	6 months	N/M	6-month period.	9	9	18	Personal reasons (9), un known reason (5), low back pain (1), malaise 1)
Karst et al. [39]	Verum needle Seirin B-type needles no. 8 (0.3 × 0.3 mm) and no. 3 (0.2 × 0.15 mm) were used for verum acupuncture.	Placebo needle The tip of the needle is blunt in order to cause a pricking sensation without actually puncturing the skin	N/M	N/M	6 weeks	Two treatments per week for a total of 10 treatments.	6 month period	N/M	N/M	N/M	N/M
Torelli et al. [40]	Standardized physiotherapy	Observation period of similar length with clinic visits to the neurologist	N/M	N/M	8 weeks	Twice a week for 4 weeks	12 weeks	6	6	11	N/M

Continued

Author, year, country	Type of physiotherapy used		Home based program	Conducted by qualified physiotherapist	Duration of physiotherapy	Number of sessions	Follow up period	Withdraw (N)			Withdrawal reasons
	Group 1 (Treatment)	Group 2 (Control)						Group 1 (Treatment)	Group 2 (Control)	Total	
Bove et al. [41]	Joint manipulation of the cervical spine	Deep friction massage.	N/M	N/M	4 weeks	N/M	N/M	2	2	4	N/M
Hamed et al. [42]	Both group (A) and group (B) received standardized physiotherapy treatment program	Only analgesic medications	yes	An educational session	8 weeks	3/week	N/M	N/M	N/M	N/M	N/M
Karakurum et al. [43]	Intramuscular stimulation		N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M
Cabanillas-Barea et al. [44]	DF group received three interventions on alternate days.	Standard care intervention based on pharmacological therapy	N/M	N/M	3 treatment session	3	1 month	1	2	3	N/M
Corum et al. [45]	HVLA manipulation plus exercise (intervention) (group A) and: myofascial release group, which received suboccipital inhibition plus exercise	Receive exercise only	N/M	N/M	N/M	N/M	3 month	1	3	4	Attendance failure = 3, received prophylactic treatment = 1
Sertel et al. [46]	Treatment group (20 individuals), BAT program was applied, to the second group (20 individuals), the aerobic exercise program	N/M	N/M	N/M	N/M	N/M	N/M	2	0	2	1 had transport difficulty, 1 had family problems
Kwon et al. [47]	Hamstring relaxation program (HR)	Interferometric current treatment and infrared heat treatment simultaneously	N/M	N/M	8 weeks	3 session/ week	N/M	N/M	N/M	N/M	N/M
Espí-López et al. [48]	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M
Azhdari et al. [49]	Myo-fascial release of SCM,	Drugs (nortriptyline 50 qsh and valproate Na 200 mg BID).	N/M	N/M	1 week	3/week	1 week	0	0	0	N/M
Georgoudis et al. [50]	Acupuncture followed by stretching protocol followed by physiotherapy by microwave diathermy and myofascial release techniques.	Biomedical acupuncture followed by stretching protocol	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M

Continued

Author, year, country	Type of physiotherapy used		Home based program	Conducted by qualified physiotherapist	Duration of physiotherapy	Number of sessions	Follow up period	Withdraw (N)			Withdrawal reasons
	Group 1 (Treatment)	Group 2 (Control)						Group 1 (Treatment)	Group 2 (Control)	Total	
Georgoudis et al. [51]	Acupuncture followed by stretching protocol followed by physiotherapy by microwave diathermy and myofascial release techniques.	Biomedical acupuncture followed by stretching protocol	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M	N/M
Cho et al. [52]	Sub- occipital muscle inhibition for one group and another group SMI with FHP correction exercises (SMIEx)	None	N/M	N/M	4 weeks	2/week	N/M	2	2	4	N/M
Ghanbari et al. [53]	Positional Release Therapy	Medication therapy	N/M	N/M	2 weeks	5 session in 2 weeks	2 weeks	N/M	N/M	N/M	N/M
Ebneshahidi et al. [54]	low energy laser acupuncture to LU7, LI4,GB14, and GB20 bilaterally.	A similar way except that the output power of the equipment was set to zero.	N/M	N/M	1 month	10session, 2 per week	3 month	N/M	N/M	N/M	N/M
Abaschian et al. [55]	dry needling with passive stretching treatment	Received only passive stretching.	N/M	N/M	4 weeks	N/M	N/M	N/M	N/M	N/M	N/M
Berggreen et al. [56]	(one session of trigger point massage per week for 10 weeks)	None	N/M	N/M	4 weeks	N/M	4 weeks	1	3	4	One participant in the treatment group left the study after seven treatments because of unbearable pain. One participant in the control group was excluded because she started treatment elsewhere. Two participants from the control group did not fill out the diary and questionnaires at follow-up.
Espí-López et al. [57]	SI treatment	None	N/M	N/M	4 weeks	N/M	8 weeks	2	2	2 in treatment group by OAA due to adverse effect, 2 in control group due to not feeling of improvement	N/M

Table 4. Summary of clinical outcomes and adverse events of the included studies.

Study (References)	Pain intensity outcomes (group 1 vs. group 2)	Frequency and duration outcomes	Adverse events/complications
Damapong et al. [8]	Significant reduction: 2.6 ± 0.7 vs. 2.9 ± 1.0 ($p < 0.05$)	No significant difference in frequency.	Mild ache in shoulders (group 1); drowsiness (group 2).
Gildir et al. [9]	Significant reduction: 0.7 ± 0.8 vs. 4.6 ± 0.7 cm ($p < 0.001$).	Frequency reduced: 3.8 ± 1.8 vs. 7.9 ± 2.0 days/month ($p < 0.001$).	Pain and fear during procedure; One severe headache in sham group.
Endres et al. [10]	No significant difference: 68.3 ± 12.1 vs. 67.5 ± 12.5 .	Frequency: 6.0 vs. 8.4 days/4 weeks ($p = 0.002$).	One severe headache; 10 serious events reported (mostly unrelated).
Mohamadi et al. [11]	Significant reduction: 4.38 ± 1.66 vs. 7.00 ± 1.47 ($p = 0.002$).	Frequency reduced: 5.84 ± 3.76 days/month ($p = 0.001$).	None reported.
Söderberg et al. [12]	N/M	Frequency reduced: 11.1 ± 8.4 vs. 14.0 ± 9.3 days/month ($p < 0.001$).	None reported.
Kwon et al. [13]	Significant reduction: 22.64 ± 1.80 vs. 26.45 ± 2.42 ($p = 0.001$).	N/M	None reported.
Aslam et al. [14]	Significant reduction: 3.75 ± 1.39 vs. 5.93 ± 1.62 ($p < 0.001$).	N/M	None reported.
Espí-López et al. [15]	Significant reduction ($p = 0.02$).	N/M	None reported.
Kamali et al. [16]	Significant reduction: Mean 3.00 vs. 4.22 ($p < 0.05$).	Mean frequency: 1.95 vs. 2.85.	None reported.
Saad (Nambi) et al. [17]	Significant reduction: Mean 4.34 vs. 2.29 ($p < 0.05$).	Mean frequency: 2.83 vs. 1.73.	None reported.
Espí-López et al. [18]	Significant reduction ($p < 0.05$).	Significantly reduced frequency ($p < 0.05$).	N/M
Ferragut-Garcías et al. [19]	Significant reduction in combined group ($p < 0.001$).	Frequency reduced significantly in combined group ($p < 0.001$).	N/M
Ajimsha et al. [20]	N/M	Frequency reduced: 4.9 ± 1.7 vs. 10.4 ± 2.7 days/4 weeks ($p < 0.001$).	No serious adverse events; some mild headaches reported.
Castien et al. [21]	Significant difference: 1.8 points improvement.	Frequency difference: 6.4 days ($p < 0.001$).	No adverse events were reported.
Hosseinifar et al. [22]	Significant reduction: 2.33 ± 1.11 vs. 4.06 ± 1.16 ($p = 0.000$).	N/M	No side effects.
Gopichandran et al. [23]	Significant reduction at 12 weeks: 2.84 ± 1.49 ($p < 0.001$).	Frequency reduced: 15.03 ± 3.39 ($p < 0.001$).	No any side effect.
Shafiq et al. [24]	N/M	Frequency reduced from >5/week to 1/week.	N/M
Pérez-Llanes et al. [25]	No significant difference in pain intensity ($p = 0.27$).	N/M	No adverse events reported.
Cabanillas-Barea et al. [26]	Change: -0.88 vs. 0.22 . Significant between-group change ($p < 0.001$).	Frequency Change: -7.45 vs. 2.31 ($p < 0.001$).	N/M
Monti-Ballano et al. [27]	Pain intensity: 10.50 ± 16.79 vs. 27.93 ± 52 ($p = 0.160$).	Frequency reduced: 13.00 vs. 46.63 days/month ($p = 0.009$).	N/M
Álvarez-Melcón et al. [58]	Significant reduction: Mean 4.23 vs. 4.58 ($p < 0.001$).	Duration reduced ($p < 0.001$).	N/M
Park et al. [28]	N/M	Frequency reduced: 9.10 vs. 10.25 days/4 weeks ($p < 0.001$).	N/M
Chatchawan et al. [29]	Significant reduction: 2.32 vs. 2.93 ($p < 0.001$).	Duration reduced: 6.88 vs. 10.38 hours.	Mild fever, mild soreness, and discomfort.
Choi et al. [30]	Significant reduction: 5.1 ± 2.1 vs. 7.1 ± 1.6 ($p = 0.001$).	Frequency: 7.4 vs. 10.5 days/month ($p = 0.109$).	N/M
Martín-Vera et al. [31]	Significant reduction: 5.1 ± 2.1 vs. 7.1 ± 1.6 ($p = 0.001$).	Frequency: 7.4 vs. 10.5 days/month.	N/M
Antonia (Gomez) et al. [32]	Significant reduction with combined treatment ($p = 0.001$).	N/M	N/M
van Ettekovén et al. [33]	N/M	Frequency reduced: 1.95 days/week ($p < 0.0001$).	N/M
Kanji et al. [34]	Significant reduction (44%, $p < 0.0001$).	No significant change in frequency.	Lack of quantification of sauna attendance; no adverse effects.
Madsen et al. [35]	No significant difference ($p = 0.231$).	Frequency reduced: 19 to 17 days ($p = 0.041$).	N/M
Xue et al. [36]	Significant reduction: 13.3h vs. 6.3h (duration/pain).	Frequency reduced: 3.0 vs. 1.2 per month.	N/M
Moraska et al. [37]	No significant difference ($p = 0.3$).	Frequency: 3.38 vs. 3.21 ($p = 0.026$).	N/M

Continued

Study (References)	Pain intensity outcomes (group 1 vs. group 2)	Frequency and duration outcomes	Adverse events/complications
Rinne et al. [38]	No between-group difference ($p = 0.66$).	Duration decreased but no sig diff ($p = 0.24$).	Feasible and safe to perform.
Karst et al. [39]	N/M	No significant differences in frequency.	N/M
Torelli et al. [40]	N/M	Reduced days: 14.5 to 10.5 ($p < 0.001$).	N/M
Bove et al. [41]	No significant effect.	Duration unchanged.	N/M
Hamed et al. [42]	Significant reduction ($p = 0.0001$).	Decrease in headache frequency.	N/M
Karakurum et al. [43]	N/M	Frequency: 29.6 vs. 25.2/month.	N/M
Cabanillas-Barea et al. [44]	Significant reduction: 0.86 vs. 1.82 ($p < 0.003$).	Frequency reduced: 5.82 vs. 15.49 ($p < 0.001$).	N/M
Corum et al. [45]	Significant diff between manipulation and soft tissue ($p = 0.014$).	Frequency improvement ($p < 0.001$).	N/M
Sertel et al. [46]	Significant reduction: 2.50 vs. 5.65 ($p = 0.001$).	N/M	N/M
Kwon et al. [47]	Significant reduction: 2.73 vs. 3.13 ($p = 0.057$).	N/M	N/M
Espí-López et al. [48]	N/M	N/M	N/M
Azhdari et al. [49]	Significant reduction: 3.04 vs. 6.75 ($p = 0.001$).	Frequency reduced: 4.91 vs. 5.25.	N/M
Georgoudis et al. [50]	Significant effect of time on VAS ($p < .001$).	N/M	N/M
Georgoudis et al. [51]	N/M	Frequency: 3.25 vs. 4.50 ($p = 0.004$).	N/M
Cho et al. [52]	N/M	N/M	N/M
Ghanbari et al. [53]	No significant difference ($p = 0.486$).	Frequency: 9.33 vs. 10.03 ($p = 0.508$).	N/M
Ebneshahidi et al. [54]	Significant reduction: -5 vs. -1 ($p = 0.001$).	Duration reduced: -6 vs. -1 hours ($p = 0.001$).	N/M
Abaschian et al. [55]	Significant reduction: 5.52 vs. 5.12 ($p = 0.001$).	Frequency: 10.33 vs. 11.77 ($p = 0.018$).	N/M
Berggreen et al. [56]	Significant reduction ($p = 0.047$).	N/M	N/M
Espí-López et al. [57]	Significant reduction ($p = 0.03$).	N/M	N/M

by short follow-up periods, necessitating future research with longer observation timelines.

Conclusion

Based on high-quality evidence from 51 RCTs, physical therapy is an effective and safe intervention for TTH. It significantly reduces headache burden with minimal risk of adverse events. To optimize clinical practice, a multimodal approach combining manual therapy and active exercise is recommended. Future studies should focus on long-term follow-up and standardized treatment protocols.

Acknowledgments

None.

List of Abbreviations

GBD	global burden of disease
NSAIDs	non-steroidal anti-inflammatory drugs
PRISMA	Preferred Reporting Items for Systematic Reviews and Meta-Analyses
PROSPERO	International Prospective Register of Systematic Reviews
RCTs	randomized controlled trials
TTH	tension-type headache

Conflict of interest

The authors declare that there is no conflict of interest regarding the publication of this paper.

Funding

None.

Ethical approval

Not applicable (Systematic review of existing literature).

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Supplementary content (If any) is available online.

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