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Which muscles exhibit increased stiffness in people with chronic neck pain? A systematic review with meta-analysis

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Introduction: Chronic neck pain (CNP) substantially impacts quality of life, posing both personal challenges and economic strains. This systematic review sought to discern muscle-specific stiffness differences between CNP patients and individuals without CNP.

Methods: We searched the PubMed, Scopus, and PEDro databases for studies using ultrasound elastography or myotonometry to compare muscle stiffness between CNP patients and asymptomatic controls. Using a meta-analysis with a random-effects model, we derived the pooled effect as standardized mean difference (SMD).

Results: Out of the six studies selected, the adjusted Newcastle-Ottawa rating scale for cross-sectional studies denoted three as moderate-quality and three as high-quality. Our findings indicate that the upper trapezius (UT) stiffness was elevated in CNP patients compared to their counterparts without CNP (SMD = 0.39, 95% CI = 0.05 to 0.74; p = 0.03; small effect size). The data for other muscles remained inconclusive.

Discussion: Given the case-control design of all reviewed studies, a direct causative link between UT stiffness and CNP is yet to be confirmed. As such, recommending a reduction in trapezius muscle stiffness as a primary rehabilitation strategy for CNP patients is still inconclusive and further research is needed.

KEYWORDS

chronic neck pain, shear modulus, elastography, myotonometry, upper trapezius

1. Introduction

Neck pain (NP), ranking as the fourth major cause of disability, adversely impacts quality of life, leading to dissatisfaction at work, challenges in daily activities, and substantial personal and economic implications (1–3). With an annual prevalence rate of 37.2%, NP affects up to 67% of individuals at some point in their lives (4, 5). NP lasting over three months is termed chronic NP (CNP) (6). People suffering from CNP may experience restlessness, visual disturbances, neuromuscular dysfunction, and limited mobility of the cervical spine (7–10). Pain experiences can prompt individuals to rely less on superficial neck and back muscles, leading to weakening of deeper muscles (11). Such neuromuscular adaptations and muscular property shifts in NP patients are often associated with complaints of tension, tightness, or stiffness in neck muscles (12, 13). Previous studies reported altered muscle cross-sectional area, thickness, size, and activity of deep neck muscles, decreased neck muscle strength, and existing myofascial trigger points in NP patients (14–17). An increase in neck muscle stiffness can result in muscle dysfunction and excess strain on the cervical spine, intensifying pain sensations and potentially fueling a persistent pain cycle (18, 19).

While CNP patients often subjectively report increased neck stiffness (13, 20), objective measures of muscle stiffness can offer deeper insights into CNP's underlying mechanisms and guide therapeutic interventions. Ingram et al. (19) identified heightened joint stiffness at the cervical spine in those with nonspecific neck pain. Yet, this stiffness was not directly linked to pain severity or disability levels. A emerging area of research is the assessment of muscle-specific stiffness through ultrasound elastography (21, 22). Elastography is based on stress application and tracking of tissue displacement (23). It entails generating an acoustic or mechanical wave and subsequently tracking its propagation speed; higher speeds indicate stiffer tissues (24, 25). Ultrasound elastography has proven particularly reliable in assessing stiffness in neck muscles such as the trapezius (26, 27), sternocleidomastoid (28) and deep cervical extensors (29). Another method that allows the assessment of muscle-specific stiffness is myotonometry, which is based on tracking the mechanical oscillations after a mechanical impulse (30). However, muscle stiffness results obtained with myotonometers do not always agree with elastography-based stiffness (31, 32). This discrepancy might stem from shear-wave elastography's ability to scrutinize specific regions of interest, while myotonometry largely captures superficial tissues and is influenced by factors like skin thickness and body fat percentage. Nonetheless, owing to its affordability, ease of use, and high reliability, myotonometry is gaining attraction as a valuable clinical tool in assessing muscle stiffness (30, 33).

Recent research indicates heightened stiffness in several neck muscles among CNP patients. For example, Tas et al. (21) found increased stiffness in the upper trapezius, levator scapulae, and sternocleidomastoid (but not the splenius capitis) in CNP patients compared to those without symptoms. While other studies have similarly highlighted elevated stiffness in the trapezius muscle of CNP patients (34, 35), there is no documented increased stiffness in muscles such as the multifidus and semispinalis cervicis (36). Although muscle stiffness may not be related to pain symptoms and degree of disability (19, 21, 36), muscle-specific assessments of stiffness could be used to guide rehabilitation programs. Muscles displaying increased stiffness might be prioritized for stretching interventions, while those with reduced stiffness may benefit more from strengthening exercises. To date, a comprehensive systematic review on muscle-specific stiffness variations in CNP patients remains absent. Hence, our systematic review, complemented with a meta-analysis, aims to discern which muscles show elevated stiffness in those suffering from CNP.

2. Methods

A systematic review was performed to identify randomized clinical trials exploring the differences in muscle stiffness between patients with CNP and asymptomatic control groups. This review was executed in line with the guidelines established by the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) (37).

2.1. Eligibility criteria

We considered case-control cross-sectional studies that compared muscle stiffness between idiopathic CNP patients with pain lasting >3 months and a matched asymptomatic control group. Experimental studies involving both CNP patients and an asymptomatic control group were also considered, with only baseline data being retrieved. Studies were excluded if participants suffered from specific pathologies associated with CNP, such as migraines or myofascial syndrome. Regarding the muscle stiffness outcomes, we considered ultrasound elastography and myotonomtery. Segment or joint-level stiffness assessments (e.g., passive resistance to externally-induced neck movement) were not considered, as our research question was focused on stiffness changes in specific muscles. Aiming to offer a comprehensive review, we incorporated both passive (relaxed muscle) and active (contracted muscle) stiffness outcomes. We then performed subgroup and sensitivity analyses to discern if the differences between CNP patients and the control group were contingent upon the stiffness measurement type. Inclusion criteria included original articles, either full-length or brief communications, published in peer-reviewed journals. While our database search didn't restrict based on language or publication date, only English articles were selected for data extraction.

2.2. Information sources and selection process

The search was performed in January 2023 and revised in August 2023. Both authors independently conducted all steps of the review. Potential disagreements were resolved by discussion. We searched the PubMed, Scopus and PEDro databases. In MeSH terms database, "neck pain", "muscle tone", and "elastography" were identified. To ensure a comprehensive search, further synonyms were added. For PubMed and Scopus, we used the following search string: ("neck pain" OR "neck syndrome" OR "cervical pain") AND (muscle stiffness OR muscle tone OR elastography OR myotonometry). During the manuscript revision stage (August 2023), additional MeSH terms "muscle tonus" and "Elasticity Imaging Techniques" were added to the search string, with no additional papers identified. In the PEDro database, a simple search was used with individual combinations of words, such as "neck pain stiffness", "muscle stiffness neck" "elastography neck pain" and "myotonometry neck pain" were used. While Scoups and PEDro databases were also explored for studies that used terminology similar to the specified MeSH terms, it has to be recognized that they do not utilize PubMed's MeSH terms or subject headings. In addition, we (1) reviewed the reference lists of identified systematic reviews on the topic (2) reviewed the reference lists papers already obtained through the abovementioned databased (3) performed an additional nonsystematic search of the Google Scholar search engine with the same key words as used in PEDro database. The records were imported into Mendeley (version 1.19.8) to remove the duplicates, and then exported into Microsoft Excel software for

further steps. After the duplicates were removed, the authors examined the titles and abstracts, and excluded the studies that obviously did not fit the research question. In case there was a disagreement between the authors, the paper was carried over to the next stage. Then, a full-text examination was performed, and data extraction was performed for the articles that met the exclusion criteria. The reasons for exclusions were noted and listed in the PRISMA flowchart (Figure 1).

2.3. Data collection and data items

Both authors independently undertook the data extraction process. Any discrepancies encountered were settled through meticulous review and discussion. The data collated encompassed: (a) muscle stiffness figures for CNP and control groups, including means and standard deviations; (b) demographic details such as gender, age, height, weight, and body mass index; (c) specifics of the measurement processes, including the device used, type and units of the outcome measure, measurement location, and participant's posture.

All data were organized using Microsoft Excel 2016 (Microsoft, Redmond, WA, USA). When data was solely presented graphically, we utilized the GetGraphDataDigitalizer software (version 2.26.0.20) to extract the means and standard deviations. For any instances of incomplete data, we reached out to the respective article authors via email and ResearchGate. If there was no communication 14 days post the second reminder, we deemed the data as irretrievable.

2.4. Assessment of study quality

We employed the Newcastle–Ottawa quality assessment scale for Cross-Sectional studies (38) to evaluate the quality of the included studies. To align more closely with our research question, we made slight modifications to the scale. The "respondents" item was omitted, and "ascertainment of exposure" was substituted with "patient criteria." A study met this criterion if it clearly outlined the inclusion parameters (such as symptom duration and nature) and exclusion factors (like the presence of specific pathologies) for the CNP groups. The adapted scale featured six items, classifying studies as high quality (score of 6), moderate quality (scores of 4–5), low quality (scores of 2–3), and poor quality (scores of 0–1).

2.5. Effect measures and synthesis methods

The meta-analysis was conducted in Review Manager (Version 5.3, Copenhagen: The Nordic Cochrane Center, The Cochrane Collaboration, London, UK). There is no consensus on the minimal number of studies required for a meta-analysis (39), therefore, we performed the analyses when two or more studies could be included for each muscle. We used continuous outcomes meta-analysis with a random-effects model and inverse variance method to obtained the pooled difference between CNP and control groups. The effect sizes were expressed as standardized mean difference (SMD) with 95% confidence intervals (CI), as the units of measurement were not consistent across studies. Statistical heterogeneity among studies was assessed with the I^2 statistics, following the Cochrane guidelines (the I^2 statistics of 0%–30% might not be important, 30%–60% may represent moderate heterogeneity, 60%–90% may represent substantial heterogeneity, and 90%–100% indicates considerable heterogeneity). The threshold for statistical significance was set at $p \leq 0.05$ for the pooled effect size. In case of substantial heterogeneity between the studies, a sensitivity analysis was performed by examining the effect of exclusion of studies that differed significantly from the pooled effect.

3. Results

3.1. Study selection

The literature search is summarized in Figure 1, and the basic details for each study is presented in Table 1. We included 6 studies into the meta-analysis (Table 1).

3.2. Study characteristics

The muscles represented were upper trapezius (6 studies), levator scapulae (3 studies), sternocleidomastoideous (2 studies), splenis capitis (2 studies), multifidus (2 studies), semispinalis capitis (1 study), rhomboid major (1 study) and semispinalis cervicis (1 study). All studies included adult population (range of means for age: 24.5-52.5 years). Most studies (5/6) reported body mass index or a combination of body height and body mass data, with all groups means fitting into the normal weight category (range of means for body mass index: 20.6-24.9). Most studies assessed passive muscles stiffness, with participants lying prone our supine on an examination table, except for Wolff et al. (40), who investigated muscle stiffness during functional reach tasks. Two studies (36, 41) included only women and the rest included both men and women. Four studies reported neck disability index scores that ranged from 15.1 ± 1.3 points (40) to 32.5 ± 12.3 points (36). Four studies also reported mean pain using VAS or NRS scales, with scores ranging from 2.4 ± 1.6 (22) to 4.6 ± 1.9 (21). Inclusion criteria for CNP group differed substantially across studies; for instance, three studies included participants with CNP lasting >3 months (21, 40, 41), while >6 months (36) have also been used. To assess muscle stiffness, one study used myotonometry (41) and the other five used ultrasound elastography. The studies were somewhat heterogenous regarding participant positioning and measurement locations. The details concerning participant characteristics, methodology of each study and extracted outcome data are available in Supplementary File S1.



3.3. Assessment of study quality

The studies we included were categorized as either high quality (3 studies) or moderate quality (3 studies). A notable omission across these studies was the calculation of sample size; only half of them (3 out of 6) furnished a comprehensive description. Detailed information can be found in Table 2.

3.4. Differences between CNP patients and control groups

Upper trapezius stiffness was assessed in 6 studies (Figure 2), with a total of 157 participants in CNP groups and 163 in control groups. The pooled difference (SMD = 0.39, 95% CI = 0.05–0.74) was statistically significant (p = 0.03). The heterogeneity between the studies was moderate (I^2 = 46%). However, the sensitivity analysis showed that the pooled effect was relatively robust, ranging from 0.25 to 0.49 (p = 0.003–0.040) when studies were removed one-by-one.

Levator scapulae was assessed in three studies (21, 22, 42), with a total of 71 participants in CNP groups and 78 in control groups (Figure 2). The pooled effect did not indicate any differences between CNP and control groups (SMD = 0.01; 95% CI = -0.32-0.33; p = 0.970), and the effects were very homogenous across the studies ($I^2 = 0$ %).

Sternocleidomastoideus was assessed in two studies (40, 42), with a total of 53 participants in CNP groups and 53 in control groups (Figure 2). The pooled effect did not indicate any differences between CNP and control groups (SMD = 0.02; 95% CI = -0.83-0.86; p = 0.970). The heterogeneity between the studies was substantial ($I^2 = 77\%$).

Splenius capitis was assessed in two studies (21, 36), with a total of 71 participants in CNP groups and 78 in control groups (**Figure 3**). There were no differences between the CNP and control groups (SMD = -0.01; 95% CI = -0.39-0.36; p = 0.940) and the two studies very homogenous ($I^2 = 0\%$). Finally, multifidus was assessed in two studies (36, 42), with a total of 38 participants in CNP groups and 49 in control groups (**Figure 3**). Again, the pooled effect did not indicate a difference between the

Authors		Participants		Pain symptoms	Stiffness	Muscles included and position	
	Numerus	Age in years (mean ± SD)	Sex distribution (M/F)		outcome	during testing	
Dieterich et al. (36)	n = 38 CNP = 20 C = 18	$CNP = 52.5 \pm 12.0$ $C = 48.5 \pm 9.0$	Only women.	>6 months	Shear modulus (kPa)	U. trapezius, splenius capitis, semispinalis capitis/cervicis, multifidus (all P)	
Heredia-Rizo et al. (41)	n = 40 CNP = 20 C = 20	$CNP = 46.7 \pm 6.1$ C = 23-67 years	Only women.	>3 months	Myotonometry stiffness (N/m)	U. trapezius (P)	
Ishikawa et al. (22)	n = 30 CNP = 18 C = 12	$CNP = 25.7 \pm 2.9$ $C = 24.5 \pm 3.9$	P = 9/9 C = 1 8/4	Neck and shoulder complaints for >3 months in the past year, at least once in the past week and on the day of the measurement procedures.	Strain ratio	U. trapezius (SIT), levator scapulae (SIT), rhomboid major (SIT)	
Taș et al. (21)	n = 70 CNP = 35 C = 35	$CNP = 35.6 \pm 8.3$ $C = 35.2 \pm 9.2$	P = 8/27 C = 11/24	>3 months	Shear-wave velocity (m/s)	U. trapezius (P), levator scapulae (P), SCM (S), splenius capitis (P)	
Wolff et al. (40)	n = 36 CNP = 18 C = 18	$CNP = 34.4 \pm 2.5$ $C = 32.3 \pm 3.0$	P = 7/11 C = 7/11	>3 months	Shear-wave velocity (m/s)	U. trapezius (P), SCM (S)	
Xie et al. (42)	n = 49 CNP = 18 C = 31	$CNP = 39.3 \pm 11.5$ $C = 38.8 \pm 10.8$	P = 3/15 C = 5/26	Patients with mild, moderate and severe neck disability	Shear modulus (kPa)	Upper/middle/lower/posterior/anterior trapezius, spinalis capitis, semispinalis capitis/cervicis, multifidus, levator scapulae, serratus anterior (all SIT)	

TABLE 1 Characteristics of included studies.

CNP, chornic neck pain group; C, control group; S, supine; P, prone; SIT, seated.

TABLE 2 Summary of study quality assessment.

Study	Representativeness of the sample	Sample-size	Patient criteria defined	Comparability of groups	Assessment of the outcome	Statistical testing	Total
Dieterich et al. (36)	1	0	1	1	1	1	5
Heredia-Rizo et al. (41)	1	1	1	1	1	1	6
Ishikawa et al. (22)	1	0	1	1	1	1	5
Taș et al. (21)	1	1	1	1	1	1	6
Xie et al. (42)	1	0	0	1	1	1	4
Wolff et al. (40)	1	1	1	1	1	1	6

two groups (SMD = -0.22; 95% CI = -0.65-0.21; *p* = 0.310) and the two studies were very homogenous ($I^2 = 0\%$).

4. Discussion

The meta-analysis revealed a slight elevation in trapezius muscle stiffness among CNP patients in comparison to those without symptoms. Stiffness levels of the levator scapulae, splenius capitis, SCM, and multifidus appeared comparable between CNP patients and controls. It it crucial to emphasize that this systematic review, based on case-control studies, does not clarify whether the enhanced trapezius stiffness is a result of, or a reason for, CNP. However, further research is encouraged to elucidate if addressing trapezius stiffness might be an integral objective in CNP rehabilitation strategies.

Several studies, though they didn't meet the criteria for inclusion in our meta-analysis, are worth mentioning for context. One study (43) analyzed the stiffness of upper trapezius and pectoralis minor by shear wave elastography in 39 female patients with rounded shoulder posture and neck pain. This study identified a significant positive correlation between the stiffness of the left upper trapezius and the shoulder angle in the rounded shoulder position (r = 0.44; p = 0.003). However, there was no observed relationship between the stiffness of the right upper trapezius, bilateral pectoralis minor muscles, and the angles of the shoulder and neck. A limitation of this study was the undefined duration of symptoms. The second study (44) assessed neck muscle stiffness in 20 participants, 3 of which had chronic neck pain and other 17 were healthy. Findings indicated a markedly higher stiffness in the trapezius muscle for those with chronic neck pain (p = 0.008) and a modest correlation between trapezius stiffness and body mass index (r = 0.34; p = 0.034). Yet, this study lacked clarity on the symptom duration for the CNP group. Kocur et al. (35) found increased stiffness in both upper trapezius and SCM, but failed to report the duration of the symptoms in their patients. Likewise, Onda et al. (34) reported increased stiffness in upper trapezius, but did not report the duration of the symptoms. Moreover, the meta-analysis could not be conducted for two muscles that were the focus of just a single study. Specifically, no significant difference was found between groups in the study of semispinalis cervicis and

Upper trapezius Experimental Control Std. Mean Difference Std. Mean Difference SD SD IV, Random, 95% CI IV, Random, 95% CI Study or Subgroup Mean Total Mean Total Weight Dieterich 2020 2.1 6.9 2.15 18 16.2% -0.09 [-0.73, 0.54] 6.7 20 Heredia-Rizo 2020 258.7 41.1 19 237.8 42.8 20 16.2% 0.49 [-0.15, 1.13] Ishikawa 2017 7.66 5.59 18 3.74 1.75 12 13.0% 0.85 [0.08, 1.62] Tas 2018 3.2 0.5 35 2.8 0.4 35 20.9% 0.87 [0.38, 1.37] Wolff 2022 5.53 1.27 5.57 15.8% 18 1.14 18 -0.03 [-0.69, 0.62] Xie 2023 17.11 5.66 18 16 4.75 31 17.9% 0.21 [-0.37, 0.80] Total (95% CI) 134 100.0% 0.39 [0.05, 0.74] 128 Heterogeneity: Tau² = 0.09; Chi² = 9.32, df = 5 (P = 0.10); l² = 46% -2 Test for overall effect: Z = 2.22 (P = 0.03) Higher in Control Higher in CNP Levator Scapulae Control Std. Mean Difference Std. Mean Difference Experimental Study or Subgroup IV, Random, 95% CI IV, Random, 95% CI Mean SD Total Mean SD Total Weight Ishikawa 2017 7.82 3.44 18 8.79 7.11 12 19.9% -0.18 [-0.91, 0.55] Tas 2018 2.5 2.2 35 2.2 0.3 35 48.5% 0.19 [-0.28, 0.66] Xie 2023 10.62 5.6 18 11.62 6.75 31 31.6% -0.15 [-0.74, 0.43] Total (95% CI) 71 78 100.0% 0.01 [-0.32, 0.33] Heterogeneity: Tau² = 0.00; Chi² = 1.13, df = 2 (P = 0.57); l² = 0% -2 -1 Test for overall effect: Z = 0.04 (P = 0.97) Higher in Control Higher in CNP Sternocleidomastoideous Std. Mean Difference Std. Mean Difference Experimental Control Study or Subgroup Mean SD Total Mean SD Total Weight IV, Random, 95% CI IV, Random, 95% CI Tas 2018 4 0.8 35 3.7 0.6 35 53.7% 0.42 [-0.05, 0.89] Wolff 2022 3.71 0.8 18 4.17 1.18 18 46.3% -0.45 [-1.11, 0.22] Total (95% CI) 53 53 100.0% 0.02 [-0.83, 0.86] Heterogeneity: Tau² = 0.29; Chi² = 4.34, df = 1 (P = 0.04); l² = 77% .2 Test for overall effect: Z = 0.04 (P = 0.97) Higher in Control Higher in CNP FIGURE 2

Meta-analyses for upper trapezius, levator scapulae and sternocleidomastoid.

semispinalis capitis (36), with no difference between the groups. One study (22) also assessed rhomboid major, and also reported no differences between the groups.

The underlying mechanisms of increased trapezius stiffness in CNP patients are difficult to determine. It is plausible that pain and inflammation activate type III and IV sensory endings. This could stimulate gamma-motoneurons, potentially altering muscle spindle excitability, leading to increased muscle tone (45). Notably, most included studies [with one exception (40)] assessed passive muscle stiffness at rest (without voluntary contraction), examined passive muscle stiffness during rest, without any voluntary contraction. This raises questions about whether the observed stiffness in CNP patients indicates enhanced muscle activity at rest

(26). There is prior evidence pointing to low-amplitude activity in the trapezius as a risk factor for neck and shoulder pain onset (46, 47). Alternatively, this augmented resting activation might act as a protective strategy to limit painful neck movements. This line of thought mirrors the hypotheses proposed for the observed increased stiffness in trunk extensors amongst low back pain patients (48). However, this does not fully account for why only the trapezius (and possibly SCM) displays increased stiffness. Additionally, stiffness measurements can vary based on muscle length (49), and thus on joint position. Poor neck posture may increase tension in the SCM muscle (50), which could be one of the risk factors for CNP, but also a confounding methodological factor when neck posture is not standardized in the assessment of F

Splenius capitis

<u>Mean</u> 8.1 2.6	SD 4.9	Total	Mean			¢	sta. Mean Difference	Stu. Mean Difference
8.1 2.6	4.9			SD	Total	Weight	IV, Random, 95% CI	IV, Random, 95% Cl
2.6	- ·	20	8.3	5.1	18	35.1%	-0.04 [-0.68, 0.60]	
	0.4	35	2.6	0.4	35	64.9%	0.00 [-0.47, 0.47]	
		55			53	100.0%	-0.01 [-0.39, 0.36]	•
.00; Chi ²	² = 0.0)1, df =	1 (P =	0.92);	l² = 0%	D	_	
= 0.07 (P = 0	.94)						Higher in Control Higher in CNP
Exper	imen	tal	c	ontro			0/ I M D'//	
		u	U	onuo			Std. Mean Difference	Std. Mean Difference
Mean	SD	Total	Mean	SD	Total	Weight	IV, Random, 95% Cl	Std. Mean Difference IV, Random, 95% Cl
Mean 15.5	SD 11.1	Total 20	Mean 17.1	SD 11.7	Total	Weight 45.6%	Std. Mean Difference IV, Random, 95% CI -0.14 [-0.78, 0.50]	Std. Mean Difference IV, Random, 95% Cl
<u>Mean</u> 15.5 9.55	SD 11.1 4.53	<u>Total</u> 20 18	<u>Mean</u> 17.1 11.32	SD 11.7 6.65	Total 18 31	Weight 45.6% 54.4%	Std. Mean Difference IV, Random, 95% CI -0.14 [-0.78, 0.50] -0.29 [-0.88, 0.29]	Std. Mean Difference IV, Random, 95% Cl
<u>Mean</u> 15.5 9.55	SD 11.1 4.53	<u>Total</u> 20 18 38	<u>Mean</u> 17.1 11.32	<u>SD</u> 11.7 6.65	Total 18 31 49	Weight 45.6% 54.4% 100.0%	Std. Mean Difference IV, Random, 95% CI -0.14 [-0.78, 0.50] -0.29 [-0.88, 0.29] -0.22 [-0.65, 0.21]	Std. Mean Difference IV, Random, 95% CI
Mean 15.5 9.55	<u>SD</u> 11.1 4.53 ² = 0.1	<u>Total</u> 20 18 38 2, df =	<u>Mean</u> 17.1 11.32	<u>SD</u> 11.7 6.65	Total 18 31 49 ² = 0%	Weight 45.6% 54.4% 100.0%	Std. Mean Difference IV, Random, 95% CI -0.14 [-0.78, 0.50] -0.29 [-0.88, 0.29] -0.22 [-0.65, 0.21]	Std. Mean Difference IV, Random, 95% CI
	.00; Chi ^a = 0.07 (Exper	.00; Chi ² = 0.0 = 0.07 (P = 0.	.00; Chi ² = 0.01, df = = 0.07 (P = 0.94)	.00; Chi ² = 0.01, df = 1 (P = (= 0.07 (P = 0.94)	.00; Chi ² = 0.01, df = 1 (P = 0.92); = 0.07 (P = 0.94)	.00; Chi ² = 0.01, df = 1 (P = 0.92); l ² = 0% = 0.07 (P = 0.94)	.00; Chi ² = 0.01, df = 1 (P = 0.92); l ² = 0% = 0.07 (P = 0.94)	.00; Chi ² = 0.01, df = 1 (P = 0.92); l ² = 0% -

stiffness. It would be beneficial for upcoming research on CNP patients to integrate EMG with shear-wave elastography. This approach could discern whether the increased stiffness is attributed to passive mechanical characteristics or elevated muscle activity. Such a distinction could profoundly impact treatment decisions. For instance, if addressing muscle mechanical properties is the goal, specific interventions like prolonged static stretching might be required (51).

While the exact role of stiffness in the trapezius in the onset of CNP remains uncertain, prioritizing the reduction of this stiffness could be a central objective of rehabilitation programs. In line with this, neck muscle stretching, especially of the trapezius, has been associated with immediate relief from neck pain (52). This is corroborated by a study which noted a simultaneous decrease in pain scores and upper trapezius muscle stiffness, as measured by elastography, in patients afflicted with myofascial pain syndrome. Contrary to our initial assumptions, we observed no heightened stiffness in the levator scapulae among CNP patients. Only one research (21) indicated a significant difference between CNP patients and control subjects, though the effect size was relatively small (specific study SMD = 0.42). This is particularly intriguing since the levator scapulae is frequently reported as tender among adults suffering from nonspecific neck pain (53). A potential limitation of the included studies is the lack of control for myofascial trigger points, which are also associated with increased elastographybased stiffness scores (54). Such active trigger points appear more frequently in the upper trapezius (40%) compared to the levator scapulae (15%) in those with mechanical neck pain (55).

It is plausible that the observed trapezius stiffness in CNP could be attributed to the presence of myofascial trigger points. Consequently, stretching might alleviate pain by targeting these trigger points rather than purely addressing stiffness. This theory gains traction when considering research that has observed pain mitigation following stretching or other "release" techniques aimed at neck trigger points (56, 57). Irrespective of the exact mechanism at play, trapezius stretching may be a pivotal component in CNP treatment.

As we expected, no difference in the stiffness of other muscles (splenius capitis and multifidus) was found between the CNP and control groups. However, these results are based on a very limited number of studies (2 studies for each muscle); therefore, further studies are needed to reach a more solid conclusion. Multifidus stiffness tended to be even lower in CNP patients, suggesting that this muscle may benefit from strengthening in this group. Along with the transversus abdominis, the multifidus is often considered critical for healthy spinal function (58), and structural changes (e.g., atrophy and fat infiltration) in the multifidus muscle at the lumbar level have already been demonstrated in patients with low back pain (59). However, multifidus stiffness is increased in low back pain (60), which is contrary to the trends we have observed for the multifidus in CNP. This discrepancy is difficult to explain because different factors such as the decrease in muscle function and protective co-contraction could have opposite effects on the stiffness results. Nevertheless, in CNP, it might be justified to perform strengthening or stabilization exercises in addition to stretching exercises. Indeed, it has been shown that neck strengthening (61) and stabilization (62) may be

beneficial in CNP patients, which could be related to improved multifidus function.

This systematic review does come with certain limitations that merit attention. Firstly, our conclusions are drawn strictly from case-control studies, meaning we cannot infer direct cause-andeffect relationships. For clarity on whether muscle stiffness is a precursor or an outcome of CNP, future research should adopt a prospective approach. Even though the case-control studies we included were generally of commendable quality, they employed varied assessment techniques (spanning elastography to myotonometry) and had participants with differing durations of symptoms. Notably, three of these studies were solely focused on women, while others incorporated both genders without conducting gender-specific analyses. Assessment of muscle stiffness may be influenced by factors such as adipose tissue content (63), which could cofound the relationship between stiffness scores and pain. Such discrepancies are especially significant for myotonometry, which concentrates on the muscle's superficial portions. However, the findings from myotonometry-centric studies generally mirrored those from elastography-centric ones, with no notable statistical discrepancies. Further reasons for heterogeneity include different elastography methods (strain elastography and shear-wave elastography) and different stiffness units (strain rate, shear modulus, shear-wave velocity). Yet, the limited number of studies at our disposal did not highlight any consistent effect of the assessment tool or outcome units on the results. Upcoming research should contemplate incorporating moderator variables, including age, sex, and body composition. Lastly, our literature exploration was constrained to three databases, possibly leading to the omission of some pertinent studies. Nevertheless, additional non-systematic searches in Google Scholar and reviews of reference lists uncovered 53 potential articles in our search's preliminary stage. However, none of these additional articles were integrated into the final review. This suggests that the search strategy was sufficient to include all relevant studies.

5. Conclusion

This article aimed to investigate which muscles exhibit increased stiffness in patients with CNP. Only a small increase in stiffness for the trapezius muscle was confirmed, whereas stiffness of the other muscles was similar in CNP patients and asymptomatic individuals. Nonetheless, a definitive link between

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muscle stiffness and CNP remains unestablished. As such, recommending a reduction in trapezius muscle stiffness as a primary rehabilitation strategy is still inconclusive and further research is needed.

Data availability statement

The original contributions presented in the study are included in the article/Supplementary Material, further inquiries can be directed to the corresponding author.

Author contributions

Conceptualization, MO and ZK; methodology, ZK; software, ZK; literature search and data extraction, MO; writing—original draft preparation, MO; writing—review and editing, ZK; All authors contributed to the article and approved the submitted version.

Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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Supplementary material

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fspor.2023. 1172514/full#supplementary-material.

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