Conservative Management of Mechanical Neck Disorders: A Systematic Review

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ABSTRACT. Objective. To determine if conservative treatments (manual therapies, physical medicine methods, medication, and patient education) relieved pain or improved function/disability, patient satisfaction, and global perceived effect in adults with acute, subacute, and chronic mechanical neck disorders (MND) by updating 11 systematic reviews of randomized controlled trials (RCT).

> Methods. Two independent authors selected studies, abstracted data, and assessed methodological quality from computerized databases. We calculated relative risks and standardized mean differences (SMD) when possible. In the absence of heterogeneity, we calculated pooled effect sizes.

> **Results.** We studied 88 unique RCT. The mean methodological quality scores were acceptable in 59% of the trials. We noted strong evidence of benefit for maintained pain reduction [pooled SMD -0.85 (95% CI –1.20, –0.50)], improvement in function, and positive global perceived effect favoring exercise plus mobilization/manipulation versus control for subacute/chronic MND. We found moderate evidence of longterm benefit for improved function favoring direct neck strengthening and stretching for chronic MND, and for high global perceived effect favoring vertigo exercises. We noted moderate evidence of no benefit for botulinium-A injection [pooled SMD -0.39 (95% CI -01.25, 0.47)]. We found many treatments demonstrating short-term effects.

> Conclusion. Exercise combined with mobilization/manipulation, exercise alone, and intramuscular lidocaine for chronic MND; intravenous glucocorticoid for acute whiplash associated disorders; and low-level laser therapy demonstrated either intermediate or longterm benefits. Optimal dosage of effective techniques and prognostic indicators for responders to care should be explored in future research. (First Release Jan 15 2007; J Rheumatol 2007;34:1083-102)

Key Indexing Terms:

NECK RADICULAR

WHIPLASH **TREATMENTS**

DEGENERATIVE SYSTEMATIC REVIEW

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Neck pain is still a major contributor to disability worldwide¹⁻⁴, with about 70% of the population experiencing an episode of neck pain at some point in their lives 1,5 and 15% experiencing chronic neck pain⁶. Chronic pain accounts for \$150 to \$215 billion US each year in economic loss (i.e., lost workdays, therapy, disability)^{7,8}, yet very little is known about the effectiveness of many of the available treatments. In this report, we update our previous systematic reviews from the Cervical Overview Group on conservative management for mechanical neck disorders⁹⁻¹⁹.

MATERIALS AND METHODS

The medical and alternative-medicine literature was searched from 1997 to September 2004 with no language restrictions using a sensitive search strategy²⁰. It included computerized bibliographic databases: Cochrane Register of Controlled Trials (Central), Medline, Embase, Manual Alternative and Natural Therapy, Cumulative Index to Nursing and Allied Health Literature, Index to Chiropractic Literature, an acupuncture database in China (root to September 2005). Medical Subject Headings key words included terms related to anatomic, disorder/syndrome, treatment, and methodology. Figure 1 depicts the review retrieval flow from selection to metaanalyses. Two independent reviewers conducted study selection using pilot-tested forms (qw kappa 0.82, SD 0.05)²¹.

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Gross, et al: Management of neck disorders

Selection criteria

Type of study. Published or unpublished (quasi-) randomized controlled trials.

Type of participant. Adults with acute (< 30 days), subacute (30–90 days), or chronic (> 90 days) neck disorders categorized as: (1) mechanical neck disorders (MND), including whiplash associated disorders (WAD I/II) 22,23 , myofascial neck pain, and degenerative changes or OA 24 ; (2) neck disorder with headache (NDH) $^{25-27}$; and (3) neck disorder with radicular findings (NDR), including WAD III 22,23 .

Type of intervention. Medication, medical injections¹⁸, acupuncture¹⁹, electrotherapy¹⁷, exercise¹⁶, low-level laser therapy¹¹, orthosis, thermal agents¹², traction¹³, massage¹⁵, mobilization, manipulation¹⁰, and patient education¹⁴.

The control group consisted of a placebo, wait-list/no treatment control; active treatment control (e.g., exercise and ultrasound vs ultrasound); or inactive treatment control (e.g., sham transcutaneous electrical nerve stimulation). Other comparisons were excluded.

Type of outcome. Pain, disability/function including work related measures, patient satisfaction, and global perceived effect (GPE)²⁸. Followup periods were defined as post-treatment (≤ 1 day), short-term (> 1 day to < 3 months), intermediate term (≥ 3 months to < 1 year), and longterm (≥ 1 year).

Two independent reviewers conducted data abstraction using pilot-tested forms. We calculated standard mean difference (SMD), relative risk (RR), number needed to treat, absolute benefit, and treatment advantage (Table 1, Figures 2 and 3). In the absence of heterogeneity (p ≥ 0.05), data were pooled statistically (random effects model) when we judged the studies to be clinically and statistically similar by Q-test (Figure 4). We categorized our findings using levels of evidence (Table 2)^{29,30}.

Methodological quality. We had at least 2 authors independently assess each selected study for methodological quality, based on the validated Jadad criteria³¹ (maximum score 5, high/acceptable score ≥ 3) and the van Tulder criteria³⁰ (maximum score 11, high/acceptable score ≥ 6; Table 2). The mean scores were 2.9 (SD 1.2) for Jadad, et al³¹ or 6.0 (SD 2.3) for the van Tulder, et al³⁰ criteria lists. Using a cutoff value of 50% (6/11) on the van Tulder criteria list, 59% of the included studies had "acceptable" methodological quality. Table 3 shows methodological quality scores of all studies and Figure 5 the main methodological limitations of the studies by treatment category. Sensitivity analysis for methodological quality using the Jadad scale (high score ≥ 3) upheld our primary analysis. Metaregression was not possible.

RESULTS

We detailed trial findings by "level of evidence" and "treatment category" in the later sections. Table 1 details the magnitude of the effect in terms of effect size (SMD or RR), number needed to treat, and treatment advantage; Table 4 gives a summary of the level of evidence by treatment category.

Evidence of benefit

Strong evidence

We found that multimodal approaches including stretching/strengthening exercise and mobilization/manipulation for sub-acute/chronic MND, NDR, and NDH reduced pain (Figure 4³²⁻³⁶), improved function, and resulted in favorable GPE in the long term.

Moderate evidence

Exercise. We noted 7 trials that supported various methods of direct neck strengthening and stretching exercises for chronic NDH³⁵ and chronic MND^{32,37-39} (Figure 4^{40,41}) in the intermediate or long term for multiple outcomes. However, strengthening and stretching of only the shoulder region plus general conditioning^{38,42} did not alter pain in the short or long

term, but did assist in improving function in the short term for chronic MND. One study found an effect favoring active range of motion exercises for acute pain reduction of WAD in the short term^{43,44}. Other studies favored cervical proprioceptive training and eye-fixation exercises to achieve pain reduction, improved function and GPE in the short term, and GPE in the long term for cases of chronic MND^{45,46} (Figure 4). The effect for pain was not maintained in the long term.

Medicine. We found 2 controlled trials favoring specific medicines in the intermediate or long term, as follows: intravenous glucocorticoid for pain reduction and reduced sick leave in cases of acute WAD⁴⁷, and epidural injections for pain reduction and improved function in cases of chronic neck disorder with radiculopathy⁴⁸.

Low-level laser therapy. Using sensitivity analysis by disorder subtype, we found evidence to support the use of low-level laser therapy (830 or 904 nm) for pain reduction and functional improvement in the intermediate term for acute/subacute and chronic MND/degenerative changes⁴⁹⁻⁵². Although the frequency and duration of treatment were similar, other aspects of dosage (radiant power, energy density, emission frequency, duration of disorder) were diverse and precluded a metaanalysis.

Electrotherapy. We found a short course of low-frequency pulsed electromagnetic field was helpful to palliate pain for acute WAD I and II, acute MND, or chronic MND with associated degenerative changes. We noted an immediate posttreatment effect; this was not maintained into the short term⁵³⁻⁵⁷.

Intermittent traction. For pain, we determined that there was moderate evidence of benefit favoring intermittent traction compared to control or placebo for chronic MND, NDR, degenerative changes^{58,59}. These were short-term results.

Acupuncture. Acupuncture was found to be effective for pain relief compared to inactive treatments either immediately posttreatment or in short-term followup for chronic MND⁶⁰⁻⁶² (Figure 4) and NDR⁶³. However, we noted that the evidence suggests no benefit for pain relief in the intermediate and long term and no functional improvements in the short, intermediate, or long term^{61,62}. Additionally, one high-quality study assessed the traditional Chinese medicine procedure of dryneedling to trigger points⁶⁴ and another low-quality trial on local "standard points" did not relieve pain in the short term.

Limited evidence

We found limited evidence that suggested there may be benefit in the use of repetitive magnetic stimulation⁶⁶, traditional Chinese massage⁶⁷, orthopedic pillow⁶⁸, and intramuscular injection of local anesthetic (lidocaine)⁶⁹.

Evidence of no benefit

We found evidence that varied between moderate and limited, for both intermediate and longterm use, suggesting that home exercise, hot packs, electromechanical stimulation, ultra-

sound, and combination of manipulation/mobilization/modalities do not relieve chronic pain or improve function in MND. Additionally, we found that short-term evidence suggests the following treatments do not aid pain reduction: medicines notably botulinum-A⁷⁰⁻⁷⁵ (Figure 4), morphine added to an epidural injection, manipulation alone, various massage techniques, laser for myofascial pain, infrared light, static traction, spray and stretch^{76,77} (Figure 4), electrotherapies (diadynamic current, galvanic current, iontophoresis, magnetic necklace), ultra-reiz, oral splint, neck school, and advice [to rest for acute WAD pain relief was inferior to active treatments in the short term^{43,44,78} (Figure 4); advice to activate; or on pain and stress coping skills].

Conflicting evidence

We have recorded numerous trials with conflicting/unclear evidence in Table 5.

Adverse events

We found that minor, transient, and reversible side effects consisting of increased symptoms were occasionally reported. A valid estimate of clinically significant, uncommon, and rare adverse events cannot be made from these trials. Adverse effects of longterm steroid therapy⁸¹ and manipulation⁸² have been well described.

DISCUSSION

For treatment of subacute and chronic MND or NDH, our review found evidence favoring a multimodal strategy (exercise and mobilization/manipulation); exercise alone; intramuscular lidocaine injection; and low level laser therapy (for OA) for pain, function, and GPE in the short and long term. Acupuncture, low-frequency pulse electromagnetic field, repetitive magnetic stimulation, cervical orthopedic pillow, and traditional Chinese massage are favored for either immediate or short-term pain management. For acute WAD, we found that studies of intravenous glucocorticoid show reduction of work disability at 1 year, while stretching exercises and low-frequency pulse electromagnetic field reduce pain. For chronic NDR, we determined that epidural methylprednisolone and lidocaine improved function and pain in the short and long term, while intermittent traction improved pain in the short term. Other commonly used interventions were either not studied, were unclear, or were not compatible with any evidence of benefit.

Interpretation of the magnitude of these treatment effects can benefit communication with our patients, third-party payers, and policy-makers in terms of treatment advantage, expected absolute benefit, and number needed to treat. For example, as shown in Table 1, a multimodal management approach (exercise, mobilization, and manipulation) is compatible with a 28% to 70% treatment advantage over a control, and with a longterm absolute benefit in pain reduction of 25 mm on a numeric rating scale (0-100 mm) from baseline for 1

in 2 to 5 patients with subacute or chronic MND/NDH. Similarly, intramuscular lidocaine injection for chronic myofascial neck pain is associated with a 45% treatment advantage, 40 mm absolute benefit, and a number needed to treat of 3. Table 1 provides corresponding data for treatment types shown to be beneficial.

Despite a large increase in the number of trials since our 1996 review, the advances in our understanding of the effectiveness of treatments are modest. No substantive change in methodological quality has occurred since the 1980s. The main flaws were in concealment of allocation; blinding of patients, caregivers, and outcome assessors; avoidance of cointervention; and compliance. There continues to be ample room for improving the methodological quality of trials, as proposed in the Consolidated Standards of Reporting Trials (CONSORT) statement⁸³.

To date, few trials on neck disorders have looked at costs⁸⁴. However, given the lack of large treatment differences between interventions, economic evaluations are becoming increasingly important and should be performed in randomized clinical trials⁸⁵.

What are the most important unanswered questions with regard to treating mechanical neck disorders? Information on commonly used pain medications (nonsteroidal antiinflammatory drugs, acetaminophen, opioids) is needed. Glucocorticoid studies suggest reduction of work disability at 1 year; if this can be confirmed, it has important public health implications for acute whiplash injury. We need to understand the most effective treatment techniques, combinations, or approaches, and the optimal dosages. This is especially true for different forms of exercise therapy and manual therapy. Are there prognostic indicators for those who will or will not respond to care? Increased insight into compliance with treatments like exercise will help address application barriers. These are the challenging questions requiring focused attention.

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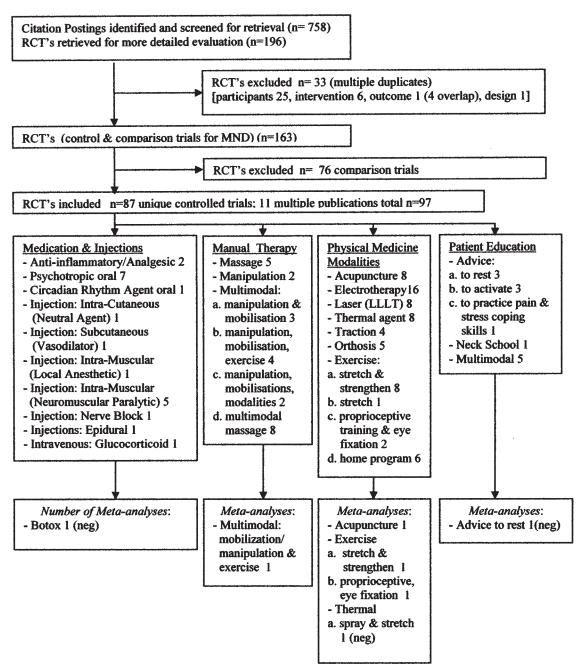


Figure 1. Study selection and metaanalysis for the 2004 Cervical Overview Group update. RCT: randomized controlled trial, MND: mechanical neck disorders, LLLT: low-level laser therapy, neg: negative metaanalysis.

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Review: Conservative Management of Mechanical Neck Disorders (M-A)

Comparison: 01 Assessment of All Trials vs Control

Outcome: 08 Pain Intensity @ intermediate or long term follow-up

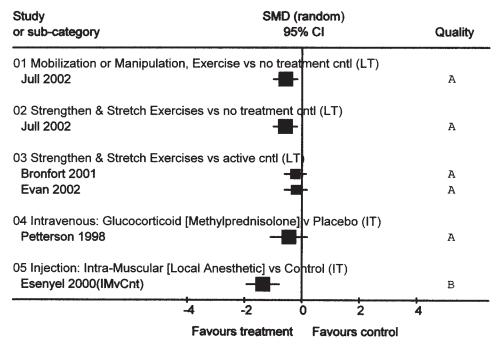


Figure 2. Intermediate (IT) and longterm (LT) results for continuous data, reported in standard mean difference (SMD), show evidence of benefit favoring pain reduction. In Pettersson's 1998 $trial^{47}$, although there was no significant effect on pain reduction, there was a clinically important effect on return to work. Direct comparison across all data is hampered by the various forms of controls (cntl) and would require a head-to-head trial comparing the various treatments. "A": high/acceptable methodological quality (≥ 3), "B": low quality on the Jadad scale³¹.

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Review: Conservative Management of Mechanical Neck Disorders (M-A)

Comparison: 01 Assessment of All Trials vs Control

Outcome: 09 Pain Intensity @ post or short term follow-up

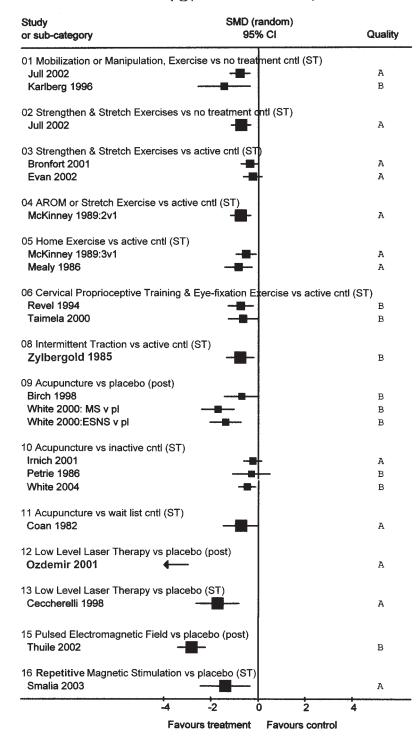


Figure 3. Short (ST) and posttreatment effects across treatment categories are depicted for continuous data on pain relief. "A": high/acceptable methodological quality (≥ 3), "B": low quality on the Jadad scale³¹. ST: short-term, AROM: active range of motion.

Review: Conservative Management of Mechanical Neck Disorders (M-A)
Comparison: 01 Assessment of All Trials vs Control

Outcome: 01 Pain Intensity

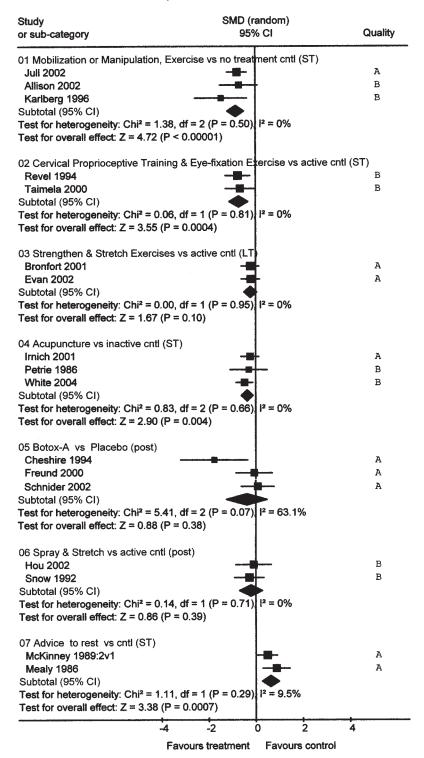


Figure 4. Metaanalyses for conservative treatments. "A": high/acceptable methodological quality (≥ 3), "B": low quality on the Jadad scale³¹.

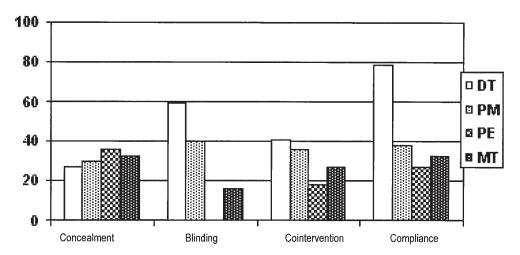


Figure 5. Proportion of studies (%) meeting quality criteria for concealment, blinding, cointervention, and compliance by treatment category. Proportion of studies meeting the van Tulder 2003 blinding criteria³⁰ across all treatments was: care provider 30%, patient 56%, outcome assessor 67%. DT: drug therapy, PM: physical medicine methods, PE: patient education, MT: manual therapy.

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Table 1. Evidence of benefit translated into clinically meaningful terms. For example, a multimodal management approach (exercise, mobilization, and manipulation) is compatible with a 28% to 70% treatment advantage over a control and a sustained absolute benefit in pain reduction of 25 mm (0–100 mm numeric rating scale) from baseline for 1 in 2 to 5 patients with subacute or chronic MND/NDH. cntl deteriorated: **baseline values different between treatment and control; LT/IT/ST: longterm/intermediate/short-term results; SMD: standard mean difference; RR: relative risk; NA: not applicable; NPQ: Nordwick Park Questionnaire 0–36 scale converted to 0–100 scale; NDI: Neck Disability Index 0–50 scale converted to 0–100 scale; NPD: Neck Pain Disability VAS 0–100; MPQ: McGill Pain Questionnaire; DC: degenerative changes; OA: osteoarthritic.

₹x	Disorder Author	vs Control Outcome	!	Effect Size (95%CI)	NNT	Treatment Advantage	Absolute Benefit
	Mobilization or Manipula		rcise				
	Chronic MND &	ation & DAC	10150				
	subacute/chronic NDH	vs active c	ntl				
	Allison 2002 ³²	Pain:	ST	pSMD -0.85 (-1.20,-0.50)	3	69%	25mm
aj	Karlberg 1996 ³⁶	i aiii.	D1	psivib =0.03 (=1.20,=0.30)	2	41%	23mm
pog	Jull 2002 ³⁵		LT		5	28%	23mm
tin.	Brodin 1985 ^{33,34}		LI	RR 0.67(0.43, 1.04)	4	NA	NA
Multimodal	DIQUII 1703			KK 0.07(0.43, 1.04)	4	INA	IVA
2	Jull 2002 ³⁵	Function:	ΙТ	pSMD -0.57 (-0.94, 0.21)	6	31%	14 NPQ
	Allison 2002 ³²	i unction.	LI	psivib -0.37 (-0.54, 0.21)	11	13%	8 NPQ
	Amson 2002				11	1370	0141 Q
	Jull 2002 ³⁵	GPE:	LT	SMD -2.73 (-3.30,-2.16)	NA	NA	69%
	Strengthen & Stretch Exe						
	Chronic NDH	vs no Rx c					
	Juli 2002 ³²	Pain:	LT	SMD -0.59 (-1.00, -0.18)	6	28%	28mm
		Function:	LT	SMD -0.59 (-1.00, -0.18)	6	32%	16 NPQ
		GPE:	LT	SMD -2.51 (-3.05, -1.97)	NA	NA	60%
	Chronic MND	vs no Rx c	<u>ntl</u>				
	Gam 1998 ³⁷	Pain:	ST	SMD -0.75 (-1.42,-0.07)	2	122%**	11mm
		vs active c	entl.				
	Bronfort 2001 ⁴⁰	Pain:	ST	pSMD -0.32 (-0.59,-0.04)	9	13%	12mm
	Evans 2002 ⁴¹				13	10%	32mm
	Bronfort 2001 ⁴⁰		LT	pSMD -0.23 (-0.50, 0.04)	15	10%	20mm
	Evans 2002 ⁴¹				14	9%	26mm
	Bronfort 2001 ⁴⁰	Function:	ST	pSMD -0.31 (-0.59,-0.04)	11	3%	14 NDI
	Evans 2002 ⁴¹				14	3%	13 NDI
	Bronfort 2001 ⁴⁰		LT	pSMD -0.34 (-0.62,-0.07)	NA	11%	11 NDI
ise	Evans 2002 ⁴¹			_		11%	11 NDI
Exercise	Bronfort 2001 ⁴⁰	GPE:	ST	pSMD -0.24 (-0.52, 0.04)	NA	NA	14%
ΕX	Evans 2002 ⁴¹			-		NA	15%
	Bronfort 2001 ⁴⁰		LT	pSMD -0.26 (-0.54, 0.02)	NA	NA	22%
	Evans 2002 ⁴¹			- , , , ,		NA	22%
	Bronfort 2001 ⁴⁰	Satisfaction	ST	pSMD -0.18 (-0.46, 0.10)	NA	NA	11%
	Evans 2002 ⁴¹					NA	13%
	Bronfort 2001 ⁴⁰		LT	pSMD -0.26 (-0.54, 0.02)	NA	NA	13%
	Evans 2002 ⁴¹			F	NA	NA	15%
	AROM or Stretch Exerc	ise alone					
	Acute WAD	vs active o	entl				
	McKinney 1989 ^{43,44} 2v1	Pain:	ST	SMD -0.77 (-1.20, -0.35)	14	17%	23mn
	Home Exercise						
	Acute WAD	vs active	cntl				
	McKinney 1989 ^{43,44} 3v1	Pain:	ST	SMD -0.58 (-0.96, -0.11)	12	19%	35mn
	Mealy 1986	Pain:	ST	SMD -0.86 (-1.44, -0.28)	5	32%	41mn

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Rx	Disorder Author	vs Control	<u>l</u>	Effect Size (95%CI)	NNT	Treatment Advantage	Absolute Benefit
	Chronic MND Taimela 2000 ⁴⁶ Revel 1994 ⁴⁵	vs active c	entl ST	pSMD -0.72 (-1.12,-0.32)	5 4	32% 34%	27mm 22mm
	Revel 1994 ⁴⁵ Taimela 2000 ⁴⁶	Function: GPE:	ST post IT	RR 0.55 (0.33, 0.89) SMD -2.32 (-3.10, -1.53) SMD -1.60 (-2.30,-0.91)	3 NA NA	NA NA NA	NA 77% 72%
	Home Exercise		11	SMD -1.00 (-2.30,-0.91)	INA	INA	1270
	Acute WAD McKinney 1989 ^{43,44} 3v1 Mealy 1986 ⁷⁸	vs active of Pain: Pain:	ST ST	SMD -0.58 (-0.96, -0.11) SMD -0.86 (-1.44, -0.28)	12 5	19% 32%	35mm 41mm
ge	Traditional Chinese Ther						
Massage	Chronic MND Cen 2003 ⁶⁷	vs wait cni Function:		SMD -1.75 (-2.82, -0.68)	2	72%	19NPQ
	Intermittent Traction						
Traction	Acute to chronic MND, NDR, DC	vs active c					
Tra	Zybergold 1985 ⁵⁸ Goldie 1970 ⁵⁹ Goldie 1970 ⁵⁹	Pain: Pain: GPE:	ST ST ST	SMD -0.78 (-1.36,-0.21) RR 0.50 (0.27, 0.90) RR 0.50 (0.27, 0.90)	5 3 3	36% 35% NA	2MPQ NA NA
	Subacute/chronic MND Petrie 1986 ⁶⁰ Irnich 2001 ⁶¹ White 2004 ⁶²	v inactive Pain:		pSMD -0.37 (-0.61,-0.12)	17 13 12	-11%* 14% 16%	15mm 30mm 29mm
Acupuncture	Petrie 1983 ⁶⁵ Irnich 2002 ⁶⁴		post post	RR 0.14 (0.02, 0.88) SMD -0.49 (-0.98, -0.01)	2 2	86% 38%	NA 16mm
Acı	Birch 1998 ⁷⁹ White 2000 ⁸⁰ (MS) White 2000 ⁸⁰ (ESNS) Chronic NDR	Pain: Pain:	post post post	SMD -0.72 (-1.45, 0.01) SMD -1.73 (-2.41,-1.04) SMD -1.40 (-2.05,-0.75)	5 3 3	30% 29% 25%	29mm 29mm 25mm
	Coan 1982 ⁶³ Low Level Laser Therap	Pain:	ST	SMD -0.74 (-1.49, 0.00)	3	41%	24mm
	Acute/subacute/chronic MND with DC (OA) Ceccherelli 1998 ⁷⁰	vs placebo	_	SMD 174 (264 092)	2	104%**	20,,,,,,
¥	Özdemer 2001 ⁵⁰ Soriano 1996 ⁵²	Pain: IT/p Pain: IT/p	ost ost	SMD -1.74 (-2.64, -0.83) SMD -3.86 (-4.73, -2.98) RR 0.39 (0.24, 0.64)	2 2 3	63% 50%	38mm 54mm NA
Laser	Taverna 1990 ⁵¹ Özdemer 2001 ⁵⁰	Pain: IT/p Function:		RR 0.30 (0.12, 0.76) SMD -4.51 (-5.48, -3.53)	3 2	47% 62%	NA 68mm

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outhor Alsed Electromagnetic Formic MND/DC Rigato 2002 ⁵⁵ Trock 1994 ⁵⁷	TAMMEA vs placeb Pain: Extreme vs placeb Pain: ADL: po	IF) E o o post LF	RR 0.30 (0.18, 0.51) Significant	2 NA	Advantage 70% NA	NA
Aronic MND/DC Rigato 2002 ⁵⁵	TAMMEA vs placeb Pain: Extreme vs placeb Pain: ADL: po	post LF post	Significant	_		NA
Rigato 2002 ⁵⁵	vs placeb Pain: Extreme vs placeb Pain: ADL: po	post <u>LF</u> post	Significant	_		NA
	Pain: Extreme vs placeb Pain: ADL: po	post <u>LF</u> <u>o</u> post	Significant	_		NA
	Extreme vs placeb Pain:	<u>LF</u> oo post	Significant	_		NA
Trock 1994 ⁵⁷	vs placeb Pain: ADL: po	post		NA	NΔ	
Trock 1994 ⁵⁷	Pain: ADL: po	post		NA	NΔ	
Trock 1994 ⁵⁷	ADL: po			NA	NΔ	
		ST	3.7 . · · · · · · · ·		1477	NA
			Not significant	NA	NA	NA
		st,ST	Not significant	NA	NA	NA
	GPE: pos	st, ST	Not significant	NA	NA	NA
	HF vs pla	<u>icebo</u>				
Foley-Nolan 1990 ⁵³	Pain:	post	Significant	NA	NA	NA
Rigato 2002 ⁵⁵			RR 0.31 (0.19, 0.52)	2	69%	NA
cute WAD I & II		<u>acebo</u>				
		post	SMD -2.82 (-3.41, -2.24)	3	57%	44mm
		post	Not significant	NA	NA	NA
epetative Magnetic Stin	nulation (r	MT)				
hronic MND -	vs placeb	0				
yofascial)	Pain:	ST	SMD -1.39 (-2.44, -0.33)	3	56%	37mm
Smaria 2003 ⁶⁶	Function	ST	SMD -1.39 (-2.44, -0.33)	2	57%	29NPD
travenous Glucocortico	id					
cute WAD	vs placeb	0				
Petterson 1998 ⁴⁷	Pain:	ST	SMD -0.90 (-1.57, -0.24)	NA	NA	NA
tra-muscular injection:	local anes	sthetic	(lidocaine)			
ronic MND						
yofascial)	vs placeb	0				
Esenvel 2000 ⁶⁹	Pain:	ST	SMD -1.36 (-1.93, -0.80)	3	45%	40mm
Figure	Rigato 2002 ⁵⁵ ute WAD I & II Chuile 2002 ⁵⁶ Foley-Nolan 1992 ⁵⁴ petative Magnetic Stim ronic MND - yofascial) Gmaria 2003 ⁶⁶ ravenous Glucocortico ute WAD Petterson 1998 ⁴⁷ ra-muscular injection: ronic MND	Rigato 2002 ⁵⁵ Pain: **wite WAD I & II HF vs pla Thuile 2002 ⁵⁶ Pain: **poley-Nolan 1992 ⁵⁴ Pain: **poletative Magnetic Stimulation (r **ronic MND - vs placeb* **yofascial) Pain: **Smaria 2003 ⁶⁶ Function **ravenous Glucocorticoid **we WAD vs placeb* **poletterson 1998 ⁴⁷ Pain: **ra-muscular injection: local aneseronic MND **yofascial) vs placeb* **yofascial* **yofas	Rigato 2002 ⁵⁵ Pain: post Mute WAD I & II Chuile 2002 ⁵⁶ Pain: post Poley-Nolan 1992 ⁵⁴ Pain: post Petative Magnetic Stimulation (rMT) Pronic MND - vs placebo Pofascial) Pain: ST Fravenous Glucocorticoid Mute WAD Petterson 1998 ⁴⁷ Pain: ST ra-muscular injection: local anesthetic Pronic MND Pronic MND Pronic Model Pronic ST Pronic ST Pronic Model Pronic ST Pronic Model Pronic ST Pro	Rigato 2002 ⁵⁵ Pain: post RR 0.31 (0.19, 0.52) HF vs placebo Chuile 2002 ⁵⁶ Pain: post SMD -2.82 (-3.41, -2.24) Foley-Nolan 1992 ⁵⁴ Pain: post Not significant Petative Magnetic Stimulation (rMT) ronic MND - vs placebo Smaria 2003 ⁶⁶ Pain: ST SMD -1.39 (-2.44, -0.33) Function: ST SMD -1.39 (-2.44, -0.33) ravenous Glucocorticoid ute WAD Petterson 1998 ⁴⁷ Pain: ST SMD -0.90 (-1.57, -0.24) ra-muscular injection: local anesthetic (lidocaine) ronic MND vofascial) vs placebo	Rigato 2002 ⁵⁵ Pain: post RR 0.31 (0.19, 0.52) 2 Let WAD I & II HF vs placebo Chuile 2002 ⁵⁶ Pain: post SMD -2.82 (-3.41, -2.24) 3 Foley-Nolan 1992 ⁵⁴ Pain: post Not significant NA Petative Magnetic Stimulation (rMT) ronic MND - vs placebo Vofascial) Pain: ST SMD -1.39 (-2.44, -0.33) 3 Smaria 2003 ⁶⁶ Function: ST SMD -1.39 (-2.44, -0.33) 2 ravenous Glucocorticoid Let WAD vs placebo Petterson 1998 ⁴⁷ Pain: ST SMD -0.90 (-1.57, -0.24) NA ra-muscular injection: local anesthetic (lidocaine) ronic MND Vofascial) vs placebo	Rigato 2002 ⁵⁵ Pain: post RR 0.31 (0.19, 0.52) 2 69% WHE WAD I & II HF vs placebo Chuile 2002 ⁵⁶ Pain: post SMD -2.82 (-3.41, -2.24) 3 57% Foley-Nolan 1992 ⁵⁴ Pain: post Not significant NA NA Petative Magnetic Stimulation (rMT) **ronic MND - vs placebo** Vofascial) Pain: ST SMD -1.39 (-2.44, -0.33) 3 56% Smaria 2003 ⁶⁶ Function: ST SMD -1.39 (-2.44, -0.33) 2 57% **ravenous Glucocorticoid** **we WAD vs placebo** Petterson 1998 ⁴⁷ Pain: ST SMD -0.90 (-1.57, -0.24) NA NA **ra-muscular injection: local anesthetic (lidocaine)* **ronic MND** **vs placebo** **vs placebo** Pain: ST SMD -0.90 (-1.57, -0.24) NA NA **ra-muscular injection: local anesthetic (lidocaine)* **ronic MND** **vs placebo**

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Table 2. Jadad, et al^{31} and van Tulder, et al^{30} methodological quality criteria lists and classification of "Level of Evidence" 29,30 .

Methodological Quality Criteria Lists

Jadad et al (Total score 5; high quality ≥ 3)

1a. Was the study described as randomized? (Score 1 if yes)

1b and 1c. Was the method of randomization described and appropriate to conceal allocation? (Score 1 if appropriate and -1 if not appropriate)

2a. Was the study described as double-blinded? (Score 1 if yes)

2b and 2c. Was the method of double blinding described and appropriate to maintain double-blinding? (Score 1 if appropriate and -1 if not appropriate)

3. Was there a description of how withdrawals and dropouts were handled?

(Score1 if yes)

Van Tulder et al (Total score 11; high quality ≥ 6)

A. Was the method of randomization adequate?

(Score1 if yes)

- B. Was the treatment allocation concealed?
- C. Were groups similar at baseline regarding the most important prognostic indicators?
- D. Was the patient blinded to the intervention?
- E. Was the care provider blinded to the intervention?
- F. Was the outcome assessor blinded to the intervention?
- G. Were cointerventions avoided or similar?
- H. Was the compliance acceptable in all groups?
- I. Was the withdrawal/dropout rate described and acceptable?
- J. Was the timing of the outcome assessment in all groups similar?
- K. Did the analysis include an intention-to-treat analysis?

Level of Evidence

Strong Consistent findings in multiple high-quality RCT

Moderate Findings in a single, high-quality RCT or consistent findings in

multiple low-quality trials

Limited A single low-quality RCT

Conflicting/unclear Inconsistent results in multiple RCT

No evidence No studies were identified

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Table 3. Methodological quality of selected trials. Agreement between both methodological criteria list scores was acceptable (Spearman rank correlation: rho = 0.76). Specific major gaps continue to be dominant for concealment of treatment allocation, blinding (outcome assessor, patient, and treater), avoiding cointervention, and compliance to intervention (see Figure 5). Mobs/manip: mobilization and/or manipulation.

Total 8

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Table 3. Continued.

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Table 3. Continued

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Table 4. Review article findings by intervention characteristics categorized as showing evidence of benefit/no benefit. Strong level of evidence denotes consistent findings in multiple high-quality randomized controlled trials; Moderate evidence denotes findings in a single, high-quality randomized controlled trial or consistent findings in multiple low-quality trials; Limited evidence indicates a single low-quality randomized trial. The comparisons noted after the author in column 2 are those noted by the author. ST/IT/LT: short-term, intermediate, longterm; neg: negative results; MND: mechanical neck disorder; NDH: neck disorder with headache; NDR: neck disorder with radicular findings; DC: degenerative changes; WAD: whiplash associated disorder; M-A: results based on a metaanalysis; s: session; w: week; Rx: treatment; mobs: mobilizations; manip: manipulation.

Rx	Treatment Details and Disorder	Author		vel of Evidence	* * * *
			Strong	Moderate	Limited
EVI	DENCE of BENEFIT	Allison 2002 ³² : NT v CG	LT pain (M-A)		
Multimodal	Multimodal: stretching & strengthening exercise, mobilization, and manipulation for subacute/chronic MND/NDH/NDR	Allison 2002 : NT V CG Brodin 1985 ^{33,34} : 3 v 1 Karlberg 1996 ³⁶ Jull 2002 ³⁵ : MTExT v Cntl	LT gain (M-A) LT GPE LT function (M-A neg)		
se	Strengthening & stretching of neck region • for chronic MND	Bronfort 2001 ⁴⁰ : SMT/Ex v SMT Evans 2002 ⁴¹ : SMT/Ex v SMT		M-A all outcomes ST pain/function ST GPE (neg) ST Patient satisfaction (neg) LT pain (neg) LT function LT GPE (neg) LT Patient satisfaction (neg)	
Exercise		Gam 1998 ³⁷		ST pain	
щ		Lundblad 1999 ³⁸		LT pain (neg) LT function (neg)	
		Ylinen 2003 ³⁹ : EvC; SvC		LT pain	
	• for chronic NDH	Jull 2002 ³⁵		LT pain LT function LT GPE	
	Strengthen & stretching of shoulder region plus total body conditioning for chronic MND				
	Group exercise class (gymnastics) at work	Takala 1994 ⁴²		ST pain (neg) ST function ST/LT pain (neg)	
	• Feldenkrais intervention	Lundblad 1999 ³⁸		ST function LT function (neg)	
	Active range of motion or stretch exercise for acute WAD	McKinney 1989 ^{43,44} : 2 v 1		ST pain	
	Cervical proprioceptive training and eye fixation exercises • for chronic MND	Revel 1994 45		ST pain/ function ST pain/ function /GPE;	
		Taimela 2000 ⁴⁶		LT pain (neg) LT GPE	
	Home exercise for acute WAD	McKinney 1989 ^{43,44} : 3 v 1 Mealy 1986 ⁷⁸ : Active v Standard		ST pain ST pain	
	Intravenous glucocorticoid for acute WAD	Petterson 1998 ⁴⁷		ST pain IT sick leave	
Medicine	Epidural injections with methylprednisolone and lidocaine for chronic neck disorder with radiation	Stav 1993 ⁴⁸		ST/LT pain ST/LT function	· · · · · · · · · · · · · · · · · · ·
4	Intramuscular injection local anesthetic for chronic MND (myofascial pain)	Esenyel 2000 ⁶⁹		and the state of t	IT pain
Massage	Traditional Chinese therapeutic massage for chronic MND	Cen 2003 ⁶⁷			Post function

Rx	Treatment Details and Disorder	Author	Level of Evidence	
		Strong	Moderate	Limite
	Pulsed electromagnetic field			
	for acute MND/WAD & chronic OA			
			ST pain	
	 extremely low frequency 	Trock 1994 ⁵⁷	ST function (neg)	
			GPE (neg)	
	• low frequency	FF 11 000056		
<u>a</u>	3 low nequency	Thuile 2002 ⁵⁶	Post pain	
er:		Rigato 2002 ⁵⁵	Post pain	
Electrotherapy	• high frequency @ 2-3 W Rx	Foley-Nolan 1990 ⁵³	Post pain	
Ě	81	Foley-Nolan 1992 ⁵⁴	Post pain	
<u>ğ</u>	@ 4-6 W Rx	Foley-Nolan 1990 ⁵³	Post pain (neg)	
Ц	(a) 4-0 W RX	Foley-Nolan 1992 ⁵⁴	Post pain	
	0.44.44	Foley-Nolan 1992 ⁵⁴	Post pain (neg)	
	@ 12 W Rx		2 220 Farm (1148)	
	• TAMMEF	Rigato 2002 ⁵⁵	Post pain	
	Repetitive magnetic stimulation	Smania 2003 ⁶⁶		ST pain
	for chronic myofascial pain			
	Low level laser therapy			
	(GaAl-830 or 904 nm)			
Laser	for chronic MND/DC (OA)	Ceccherelli 1998 ⁴⁹	Post/IT pain	
SS .	for emonie with Direct (Ort)	Özdemir 2001 ⁵⁰	Post/IT pain	
-			Post function	
		Taverna 1990 ⁵¹	Post/IT pain	
		Soriano 1996 ⁵² Goldie 1970 ⁵⁹ : trac v Cntl	Post/IT pain	
¤.	Intermittent traction	Zylbergold 1985 ⁵⁸ : 2 v 4	ST pain, ST GPE	
22	for chronic MND, NDR, DC	Zylbergold 1985 : 2 V 4		
raction				
Ξ				
·^	Orthopedic pillow	Jochems ⁶⁸		ST pain
SIS	for chronic MND			ST GPE
Orthosis				
Ö				
	Acupuncture		ST pain (M-A)	
	Traditional Chinese medicine for chronic	Coan 1982 ⁶³	ST pain,	
	MND, NDR	Irnich 2002 ⁶⁴	Post pain	
	• Japan-style for subacute/chronic MND &	Birch 1998 ⁷⁹	ST pain	
			F	
	WAD			
	• Electroacupuncture for chronic MND & DC	White 2000 ⁸⁰	Post pain	
	Western for chronic MND	VVII 1: 000 462	D	
n re	Western for enrollie MIVE	White 2004 ⁶²	Post pain, ST/IT/	
덛			LT pain (neg) ST/IT/LT	
∄			function (neg)	
Acupuncture	 Local standard points for chronic NDR 	Petrie 1983 ⁶⁵	Post pain	
A	- Eocai standard points for enforce NDR	Teure 1965	r ost pani	
		- 4 4 4 60		
	 Local standard points for chronic MND 	Petrie 1986 ⁶⁰	ST pain (neg)	
	• Dry needling over ear TP for chronic MND	Irnich 2001 ⁶¹	ST pain,	
	m 12: 101: 22: 23: 23: 23: 23: 23: 23: 23: 23: 23		IT pain (neg)	
	• Traditional Chinese medicine (dry needling			
	TP) for chronic MND	Irnich 2002 ⁶⁴	Post pain (neg)	w <u> </u>
(V)	DENCE of NO BENEFIT			
ЭG	Botulinum-A injection	Cheshire 1994 ⁷⁰	ST pain (neg M-A)	
CII	for chronic MND with or without	Schnider 2002 ⁷¹		
-=	radiculopathy or headache	Freund 2000 ^{72,73}		
2				
Medicine	radiculopatily of headache	Wheeler 1998 ⁷⁴ Wheeler 2001 ⁷⁵		

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- importance of the neck in postural control. J Vestib Res 1996;6:439-53.
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Rx	Treatment Details and Disorder	Author		Level of Evidence	
			Strong	Moderate	Limited
	Intracutaneous injection of sterile water for NDH	Sand 1992 ⁸⁶		ST pain (neg)	
	Subcutaneous injection of a vasodilator for chronic MND	Brockow 200187		Post pain (neg)	
•	Melatonin for chronic WAD, MND	van Wieringen 2001 ⁸⁸		Post function (neg)	
,	Morphine added to epidural injection of triamcinolone plus lidocaine for chronic NDR	Castagnera 1994 ¹³⁶			Post/LT pain (neg
	Home exercise	Allison 2002 ³² : AT v Cntl		ST pain (neg)	
Exercise	for chronic MND and NDR	Koes 1992 ⁹⁰⁻⁹⁶ : GP v pl		ST function (neg) ST pain (neg) ST function (neg)	
i Xi		Brodin 1985 ^{33,34} : 2 v 1		ST pain (neg)	
***		Horneij 2001 ⁴⁷ : IT v Cntl		LT pain (neg) LT function (neg)	
	Manipulation alone • 1 session (s) for acute, subacute, chronic MND	Sloop 1982 ⁹⁸		ST pain (neg)	
_	• 3-4s/3w for chronic NDR/NDH	Bitterli 1977 ¹⁰¹ : B v C		ST pain (neg)	
rapy	Manipulation, mobilization and modalities	Brodin 1985 ^{33,34} : 3 v 2		ST pain (neg)	
Lhe	for chronic NDR/DC	Kogstad 1978 ¹²⁶ : MT v pl		LT GPE (neg)	
Manual Therapy	Massage alone • Ischemic compression for chronic myofascial neck pain and MND	Hou 2002 ⁷⁶ : B2 v B1 Hanton 2000 ¹⁰²		Post/ST pain (neg)	
Ž	occipital release for MND Western massage for subacute MND	Hanton 1997 ⁹⁹ Irnich 2001 ⁶⁴ : M v S		Post pain (neg)	
	occipital release for MND	Hanton 1997 ⁹⁹ : 1 v 3		Post pain (neg)	
	Ultrasound				
	for chronic MND (myofascial) • at 8s/4w, 3 w/cm ² , 3 min per TP	Gam 1998 ³⁷ :US v pl		Post pain (neg) Post function (neg) Post/IT GPE (neg)	
	• at 10s/2w, 1.5w/cm ² , 6 min per TP	Esenyel 2000 ⁶⁹ : US v Cr	tl	Post/IT pain	
ts	for acute WAD	Flynn 1986 ¹⁰⁵ : US v pl		Post pain (neg)	
Thermal Agents	Hot pack	Hurwitz 2002 ⁸² :		LT pain (neg)	
1 A	for chronic MND	(heat, mobs v mobs); (heat, manip v manip);		LT function (neg) LT patient	
ma		(heat, mobs, EMS v		satisfaction (neg)	
her		mobs, EMS),			
I		(heat, manip, EMS v manip, EMS)			
	Infrared light • for subacute/chronic MND/DC	Lewith 1981 ¹⁰⁴		ST pain (neg)	
	for myofascial neck pain	Birch 1998 ⁷⁹		ST pain (neg)	
	Spray & stretch	Snow 1992 ⁷⁷		Post pain	
	for chronic MND (myofascial)	Hou 2003 ⁷⁶		(neg M-A)	
apy	Modulated galvanic current (diadynamic current) for chronic NDR/NDH	Philipson 1983 ¹⁶³			Post pain (neg) Post GPE (neg)
Electrotherapy	Iontophoresis for acute WAD	Fialka 1989 ¹⁰⁶			Post pain (neg)
Elect	Magnetic necklace (static magnet) for chronic MND	Hong 1982 ¹⁰⁷			Post pain (neg)

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Rx	Treatment Details and Disorder	Author		Level of Evidence	
			Strong	Moderate	Limited
	Electrical muscle stimulation for chronic MND, NDR, NDH	Hseuh 1997 ¹⁰⁸ Hurwitz 2002 ⁸² : (EMS, manip v manip); (EMS, mobs v mobs); (EMS, heat, manip v heat, manip); (EMS, heat, mobs)		LT pain (neg) LT function (neg) LT patient satisfaction (neg)	
	Low-level laser therapy (GaAs-830 nm) for subacute/chronic MND (myofascial)	Thorsen 1991 ¹⁰⁹		ST pain (neg)	
Lasei	for chronic MND (myofascial)	Thorsen 1992 ¹¹⁰ Seidel 2002 ¹¹¹		ST pain (neg) ST pain (neg)	
	Low-level laser therapy (HeNe-632.8 nm) for chronic MND (myalgia)	Waylonis 1988 ¹¹²			Post pain (neg)
	Oral splint for chronic MND, NDH	Karppinien 1999 ¹¹³		ST/LT pain (neg)	
Orthosis	Soft collar for acute WAD or NDH	Gennis 1996 ¹¹⁴ Borchgrevink 1998 ¹¹⁵ McKinney 1989 ^{43,44} : 1 v 3 Mealy 1986 ⁷⁸		ST pain (neg) IT pain (neg) IT sick leave (neg) IT GPE (neg) ST pain (neg M-A) ST pain (neg M-A)	
	Advice to activate for subacute/chronic MND	Koes 1992 ⁹⁰⁻⁹⁶ : GP v pl Kamwendo 1991 ¹¹⁶ : A v C Glossop 1982 ¹³⁷ : I v II		ST/IT pain	
Education	Advice to rest for acute WAD	McKinney 1989 ^{43,44} Mealy 1986 ⁷⁸ Borchgrevink 1998 ¹¹⁵		ST pain (neg M-A) ST pain (neg M-A) IT pain (neg) IT sick leave (neg) IT GPE (neg)	
	Advice on pain and stress coping skills for chronic MND	Horneij 2001 ⁴⁷ : SM v Cntl			LT pain (neg)
	Neck school for acute/ subacute/ chronic MND	Kamwendo 1991 ¹¹⁶ : A v C, B v C			ST pain (neg)
Traction	Static traction for acute to chronic MND, NDR, DC	Brewerton 1966 ¹¹⁷ Klaber-Moffet 1990 ¹¹⁶ Zylbergold 1985 ^{58:} 1v3		ST pain (neg) ST function (neg)	

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Table 5. Treatments with conflicting evidence.

Treatments with Conflicting Evidence	Author	Outcome
Massage (multimodal)	** 177-5-8-001	
 plus electrotherapy for chronic MND, DC 	Hou 2002 ⁷⁶ ; Brodin 1985 ^{33,34} Gam 1998 ³⁷ ; Karlberg 1996 ³⁶ ; Brodin 1985 ^{33,34} ; Fialka 1989 ¹⁰⁶ ; Schnabel 2002 ¹¹⁹ ; Koes 1992 ⁹⁰⁻⁹⁶ ; Hanton 2000 ¹⁰²	Post, ST, IT, LT for pain
• plus exercise (+/- other treatment)	Gam 1998 ³⁷ ; Karlberg 1996 ³⁶ ; Brodin	(variable), function
for acute WAD, subacute or chronic	1985 ^{33,34} ; Fialka 1989 ¹⁰⁶ ; Schnabel 2002 ¹¹⁹ ;	(variable)
MND/NDH/NDR, chronic DC	Koes 1992 ⁹⁰⁻⁹⁶ ; Hanton 2000 ¹⁰²	•
Mobilization and manipulation	Koes 1992 ⁹⁰⁻⁹⁶ : MT v pl; Bitterli 1977 ¹⁰¹ : A v	ST, LT for pain (M-A,
for subacute/chronic MND; chronic NDH	C; Jull 2002 ³⁵ : MT v Cntl	neg), function (neg), GPE
Transcutaneous electrical nerve stimulation	Flynn 1986 ¹⁰³ : UR v Pl	Post pain (neg)
including Ultra-Reiz (143 Hz) for WAD, acute	Chee 1986 ¹²⁰	Post pain (neg)
MND, chronic NDR	Hsueh 1997 ¹⁰⁸	Post pain
, , , , , , , , , , , , , , , , , , ,	Nordemar 1981 ¹²¹	ST pain (neg)
Interferential (50 Hz)		
• for acute WAD/NDH,	Fialka 1989 ¹⁰⁶	Post pain
chronic myofascial neck	Hou 2002 ⁷⁶	Post, ST, LT pain (neg)
Education within multimodal context for acute	Karlberg 1996 ³⁶ ; Persson 2001 ¹²²⁻¹²⁵ : PT v	Post, ST, IT, or LT pain
WAD, subacute/chronic MND, NDH, NDR	Col; Horneij 2001 ⁴⁷ : IT v cntl; Lundblad	(variable), function
···, -·, -·, -·, -·, -·,	1999 ³⁸ : F v cntl, PT v cntl; Kogstad 1978 ¹²⁶ :	(variable)
	CT v Pl	(1414616)
Oral psychotropic agents		
Cyclobenzaprine for subacute MND	Nasswetter 1998 ¹²⁷	Post pain
Cyclocompanie for sucure in 12	Basmajian 1978 ¹²⁸	Post muscle spasm (neg)
		r est masere spasm (neg)
 Diazepam for subacute MDN 	Basmajian 1978 ¹²⁸	Post muscle spasm (neg)
for acute MND	Basmajian 1983 ¹²⁹	Post pain (neg)
for chronic DC	Thomas 1991 ¹³⁰	Post pain (neg)
200 000000 2 0		2 000 pm (110g)
Tetrazapam for acute MND	Salzman 1993 ¹³¹	Post pain, GPE
		F , -
 Eperison hydrochloride for chronic MND 	Bose 1999 ¹³²	ST pain
-r y		
 Phenobarbital for acute MND 	Basmajian 1983 ¹²⁹	Post pain (neg)
	Payne 1964 ¹³³	Post pain
Meprobamate for chronic NDR		
Oral antiinflammatory agents and oral analgesics		
 Ibuprofen for chronic NDH 	Dostal 1978 ¹³⁴	ST pain
 Oral antiinflammatory, analgesic, education 	Koes 1992 ⁹⁰⁻⁹⁶	ST pain (neg), GPE (neg),
		function (neg)
Nerve block injections	135	
 Prilocaine 2% anesthetic block of greater 	Terzi 2002 ¹³⁵	Post pain
occipital nerve for NDH		