

NKR-nakke pico 7. Massage vs.ingen massage

Review information

Authors

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Citation example: [Empty name]. NKR-nakke pico 7. Massage vs.ingen massage. Cochrane Database of Systematic Reviews [Year], Issue [Issue].

Characteristics of studies

Characteristics of included studies

Sherman 2009

Methods	
Participants	
Interventions	
Outcomes	
Identification	
Notes	

Risk of bias table

Bias	Authors' judgement	Support for judgement
Incomplete outcome data	Low risk	From Patel, 2012
Allocation concealment	Low risk	From Patel, 2012
Selective outcome reporting	Unclear risk	From Patel, 2012

Sequence Generation	Low risk	From Patel, 2012
Blinding of participants and personnel	High risk	From Patel, 2012
Other sources of bias	Unclear risk	
Blinding of outcome assessors	High risk	From Patel, 2012

Sherman 2014

<p>Methods</p>	<p>Study design: Randomized controlled trial Study grouping: Parallel group Open Label: Cluster RCT:</p>
<p>Participants</p>	<p>Baseline Characteristics</p> <p>Intervention</p> <ul style="list-style-type: none"> ● <i>Males (%)</i>: 30.8 ● <i>Mean age (SD)</i>: 49.0 (9.9) ● <i>Pain duration weeks</i>: > 5 år: 43.6% <p>Control</p> <ul style="list-style-type: none"> ● <i>Males (%)</i>: 29.7 ● <i>Mean age (SD)</i>: 44.4 (12.2) ● <i>Pain duration weeks</i>: >5 år: 29.7% <p>Included criteria: Adults aged 20 to 64 years with chronic nonspecific neckpain lasting at least 3 months who were able and willing to attend treatment Excluded criteria: We excluded individuals whose neck pain had a pathologically identifiable cause (eg, vertebral fracture, metastatic cancer), was complex (eg, cervical radiculopathy, recent automobile accident), or was too mild, defined as scoring less than 4 on a pain intensity scaleranging from 0 to 10 and less than 5 on the Neck Disability/Index (NDI) ranging from 0 to 50. We alsoexcluded those with potential contraindications formassage (eg, hypersensitivity to touch), any massagewithin the last 3 months, massage for neck pain withinthe last year, or an inability to give informed consentor speak English. Finally, we excluded persons withmedicolegal issues related to neck or back pain. Pretreatment: Ingen markante</p>

<p>Interventions</p>	<p>Intervention Characteristics Intervention</p> <ul style="list-style-type: none"> ● <i>Description:</i> On the basis of an earlier study, 18 we defined distinct treatment protocols for both 30- and 60-minute treatments, which included range of motion assessment, hands-on check-in, massage applied directly to the neck, addressing compensatory patterns, and integration (reestablishment within a patient of being in a unified body after having received intensive isolated work). Therapists were given time limits for each part of the massage and permitted to use a broad range of massage techniques. No self-care recommendations were permitted. Eight licensed massage therapists with at least 5 years of experience were trained in the study protocol and provided massage treatments in the research clinic at Group Health. Treatment fidelity was monitored by a research assistant who was also a massage therapist and who observed a treatment for all therapists and 34% of those randomized to massage (4% of all treatments). ● <i>Duration (weeks):</i> 4 ● <i>Number of treatments:</i> 12 <p>Control</p> <ul style="list-style-type: none"> ● <i>Description:</i> venteliste ● <i>Duration (weeks):</i> 4 ● <i>Number of treatments:</i> 0
<p>Outcomes</p>	<p><i>Smerte (pain) end of treatment</i></p> <ul style="list-style-type: none"> ● Outcome type: Continuous Outcome <p><i>Smerte (pain) 4-12 weeks follow up</i></p> <ul style="list-style-type: none"> ● Outcome type: Continuous Outcome <p><i>Funktionsevne (level of function) 4-12 weeks follow up</i></p> <ul style="list-style-type: none"> ● Outcome type: Continuous Outcome <p><i>Forbrug af medicin (use of medicine) 4-12 weeks follow up</i></p> <ul style="list-style-type: none"> ● Outcome type: Dichotomous Outcome <p><i>Tilbage til arbejde (return to work) 4-12 weeks follow up</i></p> <ul style="list-style-type: none"> ● Outcome type: Dichotomous Outcome <p><i>Sygefravær (sick leave) 4-12 weeks follow up</i></p> <ul style="list-style-type: none"> ● Outcome type: Dichotomous Outcome

	<p><i>Livskvalitet (Quality of life) 4-12 weeks follow up</i></p> <ul style="list-style-type: none"> ● Outcome type: ContinuousOutcome <p><i>Frafald (dropout) Behandlingsafslutning</i></p> <ul style="list-style-type: none"> ● Outcome type: DichotomousOutcome
<p>Identification</p>	<p>Sponsorship source: Our study was funded by grant R01 AT004411 from the National Center for Complementary and Alternative Medicine, National Institutes of Health.</p> <p>Country: USA</p> <p>Setting: Primary care and general population</p> <p>Comments: Recruited participants using mailed invitations to Group Health members with neck pain–related visits to primary care clinicians, advertisements in the health plan’s magazine, posters, a study website, neighborhood blogs, and direct-mail postcards</p> <p>Authors name: Karen J. Sherman</p> <p>Institution: Department of Epidemiology, University of Washington, Seattle, Washington</p> <p>Email: sherman.k@ghc.org</p> <p>Address: Group Health Research Institute 1730 Minor Ave, Ste 1600 Seattle, WA 9810, USA</p>
<p>Notes</p>	

Risk of bias table

Bias	Authors' judgement	Support for judgement
Incomplete outcome data	Low risk	Judgement Comment: Low drop out rate. Intention to treat analysis
Allocation concealment	Low risk	Quote: "NDI scores (5-14 and ≥ 15). They were embedded in the computer-assisted tele- phone interviewing program and inaccessible to study staff before randomization. Treatments For the 4-week primary"
Selective outcome reporting	Low risk	Judgement Comment: Alle nævnte outcomes rapporteret
Sequence Generation	Low risk	Quote: "of the 6 treatment groups. Treatment assignments were generated by a statistician (A.J.C.) using the freely available R software (version 2.11.0, R-Project for Sta- tistical Computing), with random block sizes of 6 and 12 within 2 strata, based on NDI scores (5-14 and ≥ 15). They were embedded in the"

Blinding of participants and personnel	High risk	Judgement Comment: Ikke muligt da kontrol er venteliste
Other sources of bias	Low risk	Judgement Comment: Apparently no other source of bias
Blinding of outcome assessors	Unclear risk	Judgement Comment: Assessor er blindet men patient er ikke og der er tale om patientrapporterede outcomes. UNCLEAR da det er uklart i hvilken grad denne form for blinding reducerer bias

Footnotes

Characteristics of excluded studies

Celenay 2016

Reason for exclusion	Wrong patient population
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Topolska 2012

Reason for exclusion	Wrong patient population
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Footnotes

Characteristics of studies awaiting classification

Footnotes

Characteristics of ongoing studies

Footnotes

References to studies

Included studies

Sherman 2009

Published and unpublished data

[Empty]

Sherman 2014

Sherman, Karen J.; Cook, Andrea J.; Wellman, Robert D.; Hawkes, Rene J.; Kahn, Janet R.; Deyo, Richard A.; Cherkin, Daniel C.. Five-week outcomes from a dosing trial of therapeutic massage for chronic neck pain.. *Annals of Family Medicine* 2014;12(2):112-120. [DOI: <http://dx.doi.org/10.1370/afm.1602>]

Excluded studies

Celenay 2016

Celenay, Seyda Toprak; Kaya, Derya Ozer; Akbayrak, Turkan. Cervical and scapulothoracic stabilization exercises with and without connective tissue massage for chronic mechanical neck pain: A prospective, randomised controlled trial.. *Manual therapy* 2016;21(Journal Article):144-150. [DOI: <http://dx.doi.org/10.1016/j.math.2015.07.003>]

Topolska 2012

Topolska, Marta; Chrzan, Sebastian; Sapula, Rafal; Kowerski, Mieczyslaw; Sobon, Marta; Marczewski, Krzysztof. Evaluation of the effectiveness of therapeutic massage in patients with neck pain.. *Ortopedia Traumatologia Rehabilitacja* 2012;14(2):115-124. [DOI: <http://dx.doi.org/10.5604/15093492.992301>]

Data and analyses

1 Intervention vs Control

Outcome or Subgroup	Studies	Participants	Statistical Method	Effect Estimate
1.1 Smerte (pain) end of treatment	2	132	Mean Difference (IV, Random, 95% CI)	-1.84 [-2.82, -0.87]
1.1.1 Time (change)	2	132	Mean Difference (IV, Random, 95% CI)	-1.84 [-2.82, -0.87]

1.2 Smerte (pain) 4-12 weeks follow up	1	58	Mean Difference (IV, Fixed, 95% CI)	-0.10 [-1.49, 1.29]
1.3 Funktionsevne (level of function) 4-12 weeks follow up	1	58	Mean Difference (IV, Fixed, 95% CI)	-1.90 [-4.83, 1.03]
1.3.1 14 weeks follow-up	1	58	Mean Difference (IV, Fixed, 95% CI)	-1.90 [-4.83, 1.03]
1.4 Livskvalitet (Quality of life) 4-12 weeks follow up	0	0	Mean Difference (IV, Fixed, 95% CI)	Not estimable
1.5 Forbrug af medicin (use of medicin) 4-12 weeks follow up	0		Risk Ratio (IV, Fixed, 95% CI)	No totals
1.6 Tilbage til arbejde (return to work) 4-12 weeks follow up	0		Risk Ratio (IV, Fixed, 95% CI)	No totals
1.7 Sygefravær (sick leave) 4-12 weeks follow up	0		Risk Ratio (IV, Fixed, 95% CI)	No totals
1.8 Frafald (dropout) Behandlingsafslutning	2	140	Risk Ratio (IV, Random, 95% CI)	0.33 [0.07, 1.62]
1.8.1 Time	2	140	Risk Ratio (IV, Random, 95% CI)	0.33 [0.07, 1.62]