NKR 40: PICO 7 Bør patienter med nyopståede lænderygsmerter tilbydes akupunktur teknikker i tillæg til vanlig behandling?

Review information

Authors

[Empty name]¹

Citation example: [Empty name]. NKR 40: PICO 7 Bør patienter med nyopståede lænderygsmerter tilbydes akupunktur teknikker i tillæg til vanlig behandling?. Cochrane Database of Systematic Reviews [Year], Issue [Issue].

Abstract

Background

Objectives

Search methods

Selection criteria

Data collection and analysis

¹[Empty affiliation]

Main results

Authors' conclusions

Characteristics of studies

Characteristics of included studies

Kennedy 2008

Methods	Study design: Randomized controlled trial Study grouping: Parallel group Open Label: Cluster RCT:
Participants	Baseline Characteristics Acupuncture Sham acupuncture Included criteria: Adults able to give informed consent, males and femalesaged 18—70 years. Current episode of non-specific LBP, withor without referred pain, of up to 12 weeks duration. LBPwas defined as pain localised below the lowest ribs, andabove the inferior gluteal folds, with or without radiation to the lower extremities.12 Excluded criteria: Pain lasting more than 12 weeks. Presentation of red flagsas defined by the Clinical Standards Advisory Guidelines13;contra-indications to acupuncture or previous acupuncturetreatment; conflicting or ongoing treatment. Pretreatment: No significant differences between the two groups were identified;however the placebo group had higher scores on VASand RMDQ at baseline
Interventions	Intervention Characteristics Acupuncture • Acupuncture style was based on a western medical approach, 3-12 session over 4-6 weeks: x • Park Sham Device (non-penetrating device), with follow up session, no ?:

	 Sham acupuncture ◆ Acupuncture style was based on a western medical approach, 3-12 session over 4-6 weeks: ◆ Park Sham Device (non-penetrating device), with follow up session, no ?: x
Outcomes	 pain Outcome type: ContinuousOutcome Reporting: Fully reported Scale: VAS Range: 0-100 Unit of measure: none Direction: Lower is better Data value: Endpoint Disability Outcome type: ContinuousOutcome
	 Reporting: Fully reported Scale: Roland Morris Disability Questionnaire Range: 0-23 Unit of measure: none Direction: Lower is better Data value: Endpoint
	Days off work Outcome type: ContinuousOutcome Reporting: Fully reported Scale: days Range: 0-48 Unit of measure: none Direction: Lower is better Data value: Change from baseline
Identification	Sponsorship source: Country: Setting:

	Comments:
	Authors name: S. Kennedya, G.D. Baxterb, D.P. Kerra, I. Bradburya, J. Parkc, S.M. McDonougha,*
	Institution: Health and Rehabilitation Sciences Research Institute, School of Health Sciences, University of Ulster,
	Northern Ireland, United Kingdom
	Email: s.mcdonough@ulster.ac.uk (S.M. McDonough).
	Address: Tel.: +44 2890366459; fax: +44 2890368068.
Notes	

Risk of bias table

Bias	Authors' judgement	Support for judgement			
Blinding of outcome assessors	Low risk	Quote: "The primary researcher, who was unaware of patient allocation until the completion of the data analysis, carried out data collection at all time points."			
Blinding of participants and personnel	Low risk	Judgement Comment: Blinding of patients, but not clinicians - however, unlikely to influence results			
Incomplete outcome data	Low risk	Quote: "Outcome data were obtained from 94% of participants at the end of treatment; at 3 months follow up this had reduced to 83%. Reasons given for not complet- ing treatment were: no benefit gained, no time available, and work commitments."			
Selective outcome reporting	Low risk	Judgement Comment: No protocol			
Allocation concealment	Unclear risk	Quote: "To ensure concealment of allocation, an administrative assis- tant not otherwise involved in the study held the randomization list."			
Sequence Generation	Low risk	Quote: "Subjects were randomly allocated into groups using a computer generated randomization table (http://www.randomization.com)."			
Other sources of bias	Low risk				

Liu 2010

Methods	Study design: Randomized controlled trial Study grouping: Parallel group Open Label: Cluster RCT:					
Participants	Baseline Characteristics Acupuncture Medicine Acupuncture + medicine Included criteria: Acute lower back pain, restricted movements, oraccompanied with radiating leg pain. The diseaseduration was less than 2 weeks, and there was nohistory of back pain in past 4 weeks before theonset. Excluded criteria: Lumbar trauma, spine-derived pain (tumor,inflammation, infection, fracture, or syndrome ofcauda equina); muscular weakness, sensoryparalysis, weakness or hyperfunction of tendonreflex; history of peptic ulcer; recent medicalhistory of nonsteroidal anti-inflammatory drug(NSAIDs) or anticoagulant, allergic history ofNSAIDs; severe abnormal function of heart, liver orkidney. Pretreatment: There were no significant difference ingender, age and duration among the three groups					
Interventions	Intervention Characteristics Acupuncture • The needles were retained for 30 min, and manipulated every 10 min. The treatment was done once daily for 5 d.: x • Diclofenac Sodium was administered orally, 50 mg per time, twice one day for 5 d.: Medicine • The needles were retained for 30 min, and manipulated every 10 min. The treatment was done once daily for 5 d.: • Diclofenac Sodium was administered orally, 50 mg per time, twice one day for 5 d.: x Acupuncture + medicine • The needles were retained for 30 min, and manipulated every 10 min. The treatment was done once daily for 5 d.: x • Diclofenac Sodium was administered orally, 50 mg per time, twice one day for 5 d.: x					
Outcomes	pain Outcome type: ContinuousOutcome Reporting: Fully reported					

	 Scale: NRS Range: 0-10 Unit of measure: none Direction: Lower is better Data value: Endpoint Disability Outcome type: ContinuousOutcome Reporting: Fully reported Scale: Roland Morris Disability Questionnaire Range: 0-24 Unit of measure: none Direction: Lower is better Data value: Change from baseline
Identification	Sponsorship source: not reported Country: Kina Setting: hospital - outpatients Comments: Authors name: LIU Jing (刘静), LI Ning (李宁) Institution: Jiangsu Provincial Hospital of Integrated Traditional Chinese and Western Medicine, Nanjing 210028, P. R. China Email: Address: Jiangsu Provincial Hospital of Integrated Traditional Chinese and Western Medicine, Nanjing 210028, P. R. China
Notes	

Risk of bias table

Bias	Authors' judgement	Support for judgement		
Blinding of outcome assessors	Unclear risk	No info		
Blinding of participants and personnel	Unclear risk	No info		
Incomplete outcome data	Low risk			
Selective outcome reporting	Low risk			
Allocation concealment	Unclear risk	No info		
Sequence Generation	Unclear risk	No info		
Other sources of bias	High risk	Judgement Comment: Statistical analysis not reported		

Footnotes

References to studies

Included studies

Kennedy 2008

Kennedy S.; Baxter GD.; Kerr DP.; Bradbury I.; Park J.; McDonough SM.. Acupuncture for acute non-specific low back pain: a pilot randomised non-penetrating sham controlled trial.. Complementary therapies in medicine 2008;16(3):139-46. [DOI: 10.1016/j.ctim.2007.03.001]

Liu 2010

Liu, J.; Li, N.. Clinical observation of a combination of acupuncture and drug administration for non-specific acute lumbar sprain. Journal of Acupuncture and Tuina Science 2010;8(1):47-49. [DOI:]

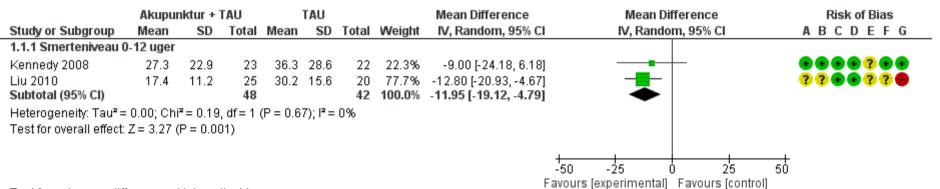
Data and analyses

1 Acupuncture + TAU vs TAU

Outcome or Subgroup	Studies	Participants	Statistical Method	Effect Estimate
1.1 Smerteniveau 0-12 uger	2		Mean Difference (IV, Random, 95% CI)	Subtotals only
1.1.1 Smerteniveau 0-12 uger	2	90	Mean Difference (IV, Random, 95% CI)	-11.95 [-19.12, -4.79]
1.2 Funktionsniveau 0-12 uger - final score	2		Mean Difference (IV, Random, 95% CI)	Subtotals only
1.2.1 Funktionsniveau 0-12 uger (EoT)	2	89	Mean Difference (IV, Random, 95% CI)	-1.16 [-4.09, 1.77]
1.4 Smerteniveau 6-18 måneder	0	0	Mean Difference (IV, Fixed, 95% CI)	Not estimable
1.5 Funktionsniveau 6-18 måneder	0	0	Mean Difference (IV, Fixed, 95% CI)	Not estimable
1.6 Sygefravær - antal sygedage 6-18 måneder	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
1.6.1 Sygefravær - antal sygedage 6-18 måneder	1	45	Mean Difference (IV, Fixed, 95% CI)	3.00 [-10.12, 16.12]
1.7 Livskvalitet 6-18 måneder	0	0	Mean Difference (IV, Fixed, 95% CI)	Not estimable
1.8 Recidiv - antal smerteepisoder 6-18 måneder	0	0	Odds Ratio (M-H, Fixed, 95% CI)	Not estimable
1.9 Sygefravær - tid tilbage-til-arbejde	0	0	Mean Difference (IV, Fixed, 95% CI)	Not estimable
1.10 Sygefravær - proportion i arbejde	0	0	Odds Ratio (M-H, Fixed, 95% CI)	Not estimable
1.11 Frafald pga. bivirkninger	0	0	Odds Ratio (M-H, Fixed, 95% CI)	Not estimable

Figures

Figure 1 (Analysis 1.1)



Test for subgroup differences: Not applicable

Risk of bias legend

- (A) Blinding of outcome assessors
- (B) Blinding of participants and personnel
- (C) Incomplete outcome data
- (D) Selective outcome reporting
- (E) Allocation concealment
- (F) Sequence Generation
- (G) Other sources of bias

Forest plot of comparison: 1 Acupuncture + TAU vs TAU, outcome: 1.1 Smerteniveau 0-12 uger.

Figure 2 (Analysis 1.2)

	True Acu	ıpuncture	(TA)	convention	al treatmen	rt (CT)		Mean Difference	Mean Difference	Risk of Bias
Study or Subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV, Random, 95% CI	IV, Random, 95% CI	ABCDEFG
1.2.1 Funktionsniveau	0-12 ugei	(EoT)								
Kennedy 2008	6	4.7	23	7	6.1	22	84.1%	-1.00 [-4.19, 2.19]		$lackbox{0.05}{$\bullet$} lackbox{0.05}{$\bullet$} lackbox{0.05}{$\bullet$} lackbox{0.05}{$\bullet$} lackbox{0.05}{$\bullet$}$
Liu 2010	4.44	12.85	24	6.45	11.95	20	15.9%	-2.01 [-9.35, 5.33]	-	? ? \varTheta 🕶 ? ? 👄
Subtotal (95% CI)			47			42	100.0%	-1.16 [-4.09, 1.77]	-	
Heterogeneity: Tau ² = (0.00; Chi ^z :	= 0.06, df:	= 1 (P = 0	$.80); I^2 = 0\%$						
Test for overall effect: 2	Z = 0.78 (P	= 0.44)								
								-	-10 -5 0 5 10	
								Δε	upuncture + TAU TAU	

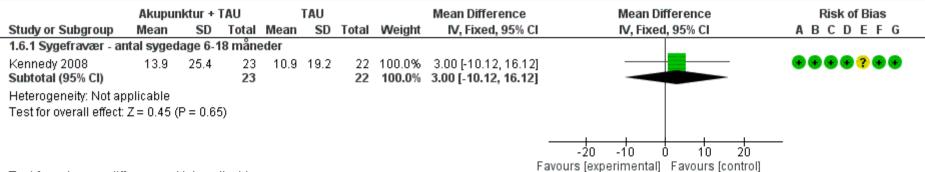
Test for subgroup differences: Not applicable

Risk of bias legend

- (A) Blinding of outcome assessors
- (B) Blinding of participants and personnel
- (C) Incomplete outcome data
- (D) Selective outcome reporting
- (E) Allocation concealment
- (F) Sequence Generation
- (G) Other sources of bias

Forest plot of comparison: 1 Acupuncture + TAU vs TAU, outcome: 1.2 Funktionsniveau 0-12 uger - final score.

Figure 3 (Analysis 1.6)



Test for subgroup differences: Not applicable

Risk of bias legend

- (A) Blinding of outcome assessors
- (B) Blinding of participants and personnel
- (C) Incomplete outcome data
- (D) Selective outcome reporting
- (E) Allocation concealment
- (F) Sequence Generation
- (G) Other sources of bias

Forest plot of comparison: 1 Acupuncture + TAU vs TAU, outcome: 1.6 Sygefravær - antal sygedage 6-18 måneder.