# PICO 1

# **Review information**

## Authors

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Citation example: S. Standard. Cochrane Database of Systematic Reviews [Year], Issue [Issue].

### **Characteristics of studies**

### **Characteristics of included studies**

## Comer 2013

Methods	Study design: Randomized controlled trial Study grouping: Parallel group
Participants	Baseline Characteristics     Intervention     Control     mean difference between groups     Overall     Included criteria: Age 50 years or overBilateral neurogenic claudication symptoms (ie exercise inducedleg pain on walking, relieved in sitting or flexion)Patient-reported limitation in walking tolerance due toNC symptoms     Excluded criteria: Cognitive impairment or other medical conditions preventing understanding or participation in the studyClearly defined radicular symptoms (ie single nerve root symptoms)Signs or symptoms of acute cauda equina syndrome or severe or worsening neurological statusrequiring medical or surgical assessment. (This includes significant or worsening nerve root orcauda equina function, significant or sinister weight loss, pyrexia, unremitting pain, significantinflammatory joint disease)     Pretreatment: Group differences at baseline:* Mean age 4.5 years younger in control group (70.3 yrs) compared to active group (75.3 yrs)* Smaller proportion of women in control group (47.4%) compared to active group (57.9%)* Median duration (time since onset of back/leg pain) longer in control group (10.0 years) compared to active group (3.5 years)*     Median number of shuttles completed greater in the control group (21.0) compared to the active group (15.0)* Mean Back pain VAS higher in the control group (63.2) compared to the active group (55.2)* Median number of physio sessions lower in control group (1) compared to active group (3)
Interventions	Intervention Characteristics     Intervention     • exercise description: Advice and education provided in both verbal and written format + condition-specific home exercise programme focusing on 1) flattening of lumbar lordosis, 2) lumbar flexion, 3) abdominal muscle activation, 4) trunk muscle strengthening, 5) aerobic fitness     • Exercise dose (frequency, duration): 2 x daily for 6 weeks     • supervised sessions: Training at home. First physiotherapy appointment, subsequent physiotherapy appointment.     Control     • exercise dose (frequency, duration): 1 initial physiotherapy appointment at which advice and education was given     • supervised sessions: no supersion     mean difference between groups     • exercise dose (frequency, duration): .     • Exercise dose (frequency, duration): .     • supervised sessions: no supersion
Outcomes	SSS physical function (ZCQ)     Outcome type: ContinuousOutcome     Reporting: Fully reported     Scale: SSS     Range: 0-100%     Direction: Lower is better     Data value: Change from baseline     OD/     Outcome type: ContinuousOutcome     Reporting: Fully reported     Scale: ODI     Range: 0-100     Direction: Higher is better     Data value: Change from baseline     Back Pain VAS     Outcome type: ContinuousOutcome     Reporting: Fully reported     Scale: ODI     Back Pain VAS     Outcome type: ContinuousOutcome     Reporting: Fully reported     Scale: VAS     Range: 0-100     Direction: Higher is better     Data value: ContinuousOutcome

	Deta value: Change from baseline Leg Pain VAS Outcome type: ContinuousOutcome Reporting: Fully reported Scale: VAS Range: 0-100 Direction: Higher is better SSS severity (ZCO) Outcome type: ContinuousOutcome Reporting: Fully reported Scale: SSS Range: 0-100 Direction: Higher is better Data value: Change from baseline N shuttles completed (excluding 2 outliers) outcome type: ContinuousOutcome Reporting: Fully reported Range: 0-7 Outfit of measure: number Outcome type: ContinuousOutcome Reporting: Fully reported Range: 0.7 Outione type: ContinuousOutcome Reporting: Fully reported Range: 0.7 Outcome type: ContinuousOutcome Reporting: Fully reported Range: 0.4 Outcome type: ContinuousOutcome Reporting: Ful
Identification	Sponsorship source: Arthritis Research UK Country: United Kingdom Setting: Leeds Musculoskeletal and Rehabilitation Service (a primary care-based musculoskeletal service Comments: . Authors name: Christine Comer Institution: 1) Leeds Musculoskeletal and Rehabilitation Service, Leeds Community Health Care, Leeds, United Kingdom; 2) Leeds Institute of Rheumatic and Musculoskeletal Disease, Faculty of Health, University of Leeds, Leeds, United Kingdom Email: p.conaghan@leeds.ac.uk (corresponding author). No email reported for lead author. Address: Not available
Notes	NKR 51 Stenose on 06/01/2017 21:09     Outcomes     Der er målt en forskel fra baseline og der er indtastet en forskel mellem de to grupper. Jo flere N i antal shuttles, jo bedre (der står ikke beskrevet hvor lang en shuttle er)Der er kun Cl for forskelle mellem de to behandlinger     Helle Algren Brogger on 30/01/2017 23:11     Outcomes     According to figure 1, page 4, n=38 in each group at baseline, but at week 8, n=35 (intervention) and n=36 (control), and at week 12 n=29 (intervention) and n=32 (control).Nonetheless, according to table 4, n=38 in both groups at all points of measurement. It is assumed that n=38 in both groups at all points of measurement is based on multiple imputation (Quote page 5: "Multiple imputation allowed all 38 patients in each group to be included in the analysis). However, it is not specified in table 4, whether mean change in control and active group is data in- or exclusive multiple imputation.

## Standard

### Risk of bias table

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	
Allocation concealment (selection bias)	Low risk	
Blinding of participants and personnel (performance bias)	High risk	
Blinding of participants and personnel (performance bias)	High risk	
Blinding of outcome assessment (detection bias)	Low risk	
Incomplete outcome data (attrition bias)	Low risk	
Selective reporting (reporting bias)	Low risk	
Other bias	Low risk	

Footnotes

## **References to studies**

#### **Included studies**

#### Comer 2013

Comer, C.; Redmond, A. C.; Bird, H. A.; Hensor, E. M. A.; Conaghan, P. G.: A home exercise programme is no more beneficial than advice and education for people with neurogenic claudication: results from a randomised controlled trial. PLoS ONE 2013;8(9):e72878. [DOI:]

#### **Excluded studies**

#### Akuthota 2012

Akuthota V.; Hammerich A.S.; Mintken P.E.; Cleland J.A.; Whitman J.M.; Singh J.R.; Knight E.; Santo, K. J.. Effectiveness of physical therapy as an adjunct to epidural steroid injections in the treatment of lumbar spinal stenosis: A pilot randomized controlled trial.. Spine Journal.Conference: 27th Annual Meeting of the North American Spine Society, NASS 2012.Dallas, TX United States.Conference Start: 20121024.Conference End: 20121027.Conference Publication: (var.pagings) 2012;12(9 SUPPL. 1):146S. [DOI: http://dx.doi.org/10.1016/j.spinee.2012.08.379]

#### Goren 2010

Goren, A.; Yildiz, N.; Topuz, O.; Findikoglu, G.; Ardic, F.. Efficacy of exercise and ultrasound in patients with lumbar spinal stenosis: a prospective randomized controlled trial [with consumer summary]. Clinical rehabilitation 2010;24(7):623-631. [DOI: ]

#### Haig 2012

Haig A.J.; Goodnight S.M.; Sandella D.; Tomkins C.; Yamakawa, K. S.. Exploration of the walking participation-to-capacity ratio in persons with neurogenic claudication, back pain, or asymptomatic volunteers.. PM and R.Conference: 2012 American Academy of Physical Medicine and Rehabilitation, AAPM&R Annual Assembly.Atlanta, GA United States.Conference Start: 20121115.Conference End: 20121118.Conference Publication: (var.pagings) 2012;4(10 SUPPL. 1):S243. [DOI: ]

#### Homayouni 2015

Homayouni, K.; Naseri, M.; Zaravar, F.; Zaravar, L.; Karimian, H.. Comparison of the effect of aquatic physical therapy and conventional physical therapy in patients with lumbar spinal stenosis (a randomized controlled trial). Journal of Musculoskeletal Research 2015;18(1):1550002. [DOI: ]

#### Kiralp 2009

Kiralp M.Z.; Cakar E.; Dincer U.; Durmus, O.: Effectiveness of the physical therapy agents on lumbar spondylosis treatment. Arthritis and Rheumatism 2009;60(Journal Article):1185. [DOI: http://dx.doi.org/10.1002/art.26259]

#### *Koc 2009*

Koc, Zarife; Ozcakir, Suheda; Sivrioglu, Koncuy; Gurbet, Alp; Kucukoglu, Selcuk. Effectiveness of physical therapy and epidural steroid injections in lumbar spinal stenosis.. Spine 2009;34(10):985-989. [DOI: http://dx.doi.org/10.1097/BRS.0b013e31819c0a6b]

#### Lindback 2016

Lindback, Yvonne; Tropp, Hans; Enthoven, Paul; Abbott, Allan; Oberg, Birgitta. PREPARE: Pre-surgery physiotherapy for patients with degenerative lumbar spine disorder: a randomized controlled trial protocol.. BMC Musculoskeletal Disorders 2016;17(Journal Article):270. [DOI: http://dx.doi.org/10.1186/s12891-016-1126-4]

#### Maher 2007

Maher, Christopher G., Re: Whitman JM, Flynn TW, Childs JD, et al. A comparison between two physical therapy treatment programs for patients with lumbar spinal stenosis: a randomized clinical trial. Spine 2006;31:2541-9.. Spine 2007;32(7):833-4. [DOI: ]

#### Sahin 2009

Sahin, Fusun; Yilmaz, Figen; Kotevoglu, Nurdan; Kuran, Banu. The efficacy of physical therapy and physical therapy plus calcitonin in the treatment of lumbar spinal stenosis.. Yonsei medical journal 2009;50(5):683-688. [DOI: http://dx.doi.org/10.3349/ymj.2009.50.5.683]

#### Schneider 2014

Schneider, Michael; Ammendolia, Carlo; Murphy, Donald; Glick, Ronald; Piva, Sara; Hile, Elizabeth; Tudorascu, Dana; Morton, Sally C.. Comparison of non-surgical treatment methods for patients with lumbar spinal stenosis: protocol for a randomized controlled trial.. Chiropractic & manual therapies 2014;22(Journal Article):19. [DOI: http://dx.doi.org/10.1186/2045-709X-22-19]

### Standard

### **Urper 2011**

Urper S.; Gunaydin R.; Karatepe A.G.; Kaya, T.. Effects of physical therapy and exercise program on clinical findings, functional status and disability in patients with lumbar spinal stenosis, Lomber spinal stenozlu olgularda fizik tedavi ve egzersiz programinin klinik bulgular, fonksiyonel durum ve ozurluluk uzerine etkisi.. Turkiye Fiziksel Tip ve Rehabilitasyon Dergisi 2011;57(Journal Article):248. [DOI: ]

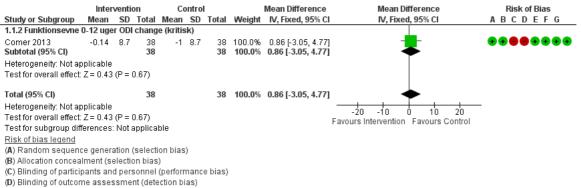
## **Data and analyses**

#### 1 Superviseret træning vs vanlig behandling

Outcome or Subgroup	Studies	Participants	Statistical Method	Effect Estimate	
1.3 Gangdistance 0-12 uger (kritisk)	0	0	Mean Difference (IV, Fixed, 95% CI)	Not estimable	
1.3.1 Gangdistance 0-12 uger (kritisk)	0	0	Mean Difference (IV, Fixed, 95% CI)	Not estimable	

### **Figures**

### Figure 1 (Analysis 1.1)



(E) Incomplete outcome data (attrition bias)

(F) Selective reporting (reporting bias)

(G) Other bias

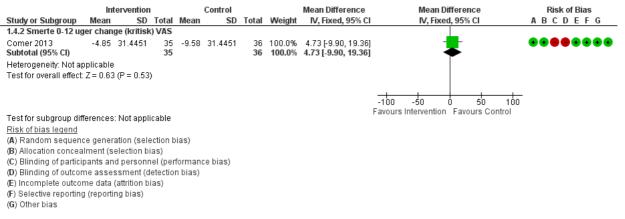
Forest plot of comparison: 1 Intervention vs Control, outcome: 1.1 Funktionsevne 0-12 uger (kritisk).

### Figure 2 (Analysis 1.3)

	Intervention 1 Intervention 2			Mean Difference Mean Difference			Risk of Bias			
Study or Subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV, Fixed, 95% Cl	IV, Fixed, 95% Cl	ABCDEFG
1.3.1 Gangdistance 0	-12 uger	(kritis	sk)							
Subtotal (95% CI)			0			0		Not estimable		
Heterogeneity: Not ap	plicable									
Test for overall effect:	Not appli	cable								
Total (95% CI)			0			0		Not estimable		
Heterogeneity: Not ap	plicable								-100 -50 0 50 100	
Test for overall effect. Not applicable							-100 -50 0 50 100 Favours Intervention 1 Favours Intervention 2			
Test for subgroup diffe	erences:	Not ap	pplicat	ole					Tavours intervention 1 Tavours intervention 2	
Risk of bias legend										
(A) Random sequenc	e genera	tion (s	selectio	on bias)						
(B) Allocation conceal	ment (se	lectior	n bias)	)						
(C) Blinding of particip	ants and	perso	onnel (	(perform	ance k	oias)				
(D) Blinding of outcom	ne asses	sment	t (dete	ction bia	s)					
(E) Incomplete outcom	ne data (a	attritior	n bias)	)						
(F) Selective reporting	(reportin	g bias	6)							
(G) Other bias										

Forest plot of comparison: 1 Intervention vs Control, outcome: 1.3 Gangdistance 0-12 uger (kritisk).

#### Figure 3 (Analysis 1.4)



Forest plot of comparison: 1 Intervention vs Control, outcome: 1.4 Smerte 0-12 uger (kritisk) NRSC.