NKR 53 demens og adfærdsforstyrrelser PICO 5 årsagsanalyse

Review information

Authors

Sundhedsstyrelsen¹

¹[Empty affiliation]

Citation example: S. NKR 53 demens og adfærdsforstyrrelser PICO 5 årsagsanalyse. Cochrane Database of Systematic Reviews [Year], Issue [Issue].

Characteristics of studies

Characteristics of included studies

Burgio 2003

Methods	
Participants	
Interventions	
Outcomes	
Identification	
Notes	Outcomes målt ved 6mdr er afrapporteret i dette studie, selvom interventionen strakte sig til 12 mdr. Data obtained from: Moniz, Cook E.; Swift K.; James I.; Malouf R.; De, Vugt M.; Verhey F. Functional analysis-based interventions for challenging behaviour in dementiaCochrane database of systematic reviews (Online) 2012;2(Journal Article):CD006929United Kingdom 2012

Risk of bias table

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	reference: Moniz Cook et al. 2012
Allocation concealment (selection bias)	Low risk	reference: Moniz Cook et al. 2012
Blinding of participants and personnel (performance bias)	High risk	reference: Moniz Cook et al. 2012
Blinding of outcome assessment (detection bias)	Low risk	reference: Moniz Cook et al. 2012
Incomplete outcome data (attrition bias)	Low risk	reference: Moniz Cook et al. 2012
Selective reporting (reporting bias)	Low risk	reference: Moniz Cook et al. 2012
Other bias	Low risk	reference: Moniz Cook et al. 2012

Chenoweth 2009

Methods	
Participants	
Interventions	
Outcomes	
Identification	
Notes	Data obtained from: Moniz, Cook E.; Swift K.; James I.; Malouf R.; De, Vugt M.; Verhey F. Functional analysis-based interventions for challenging behaviour in dementiaCochrane database of systematic reviews (Online) 2012;2(Journal Article):CD006929United Kingdom 2012 Outcomes Antipsychotic usage: not clear if the numbers are proportions or percent. We have assumed they are proportions

Risk of bias table

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	reference: Moniz Cook et al. 2012
Allocation concealment (selection bias)	Low risk	reference: Moniz Cook et al. 2012
Blinding of participants and personnel (performance bias)	Low risk	reference: Moniz Cook et al. 2012
Blinding of outcome assessment (detection bias)	Low risk	reference: Moniz Cook et al. 2012
Incomplete outcome data (attrition bias)	Low risk	reference: Moniz Cook et al. 2012
Selective reporting (reporting bias)	Low risk	reference: Moniz Cook et al. 2012
Other bias	Unclear risk	reference: Moniz Cook et al. 2012

Cohen Mansfield 2012

Methods	Study design: Randomized controlled trial Study grouping: Parallel group		
Participants	Baseline Characteristics Intervention 1 • Age y (SD): 85.9 (8.62) • MMSE (mean, SD): 7.62 (6.33) Control • Age y (SD): 85.3 (9.62) • MMSE (mean, SD): 9.38 (6.76) Included criteria: Resident 1) had been at the nursing home ⇒ 3 weeks 2) had been identified by nursing staff as agitatied at least several times a day 3) was aged ⇒ 60 years and 4) had a diagnosis of dementia Excluded criteria: Resident 1) had life expectancy of 3months 2) had a diagnosis of bipolar disorder, schizophrenia or mental retardation 3) was expected to leave nursing home within 4 months 4) had MMSE ⇒ 25 or 5) had participated in a previous TREA trial		

Interventions	Intervention • Description: Treatment Routes for exploring Agitation (TREA) • Length of treatment: 2 weeks • Longest follow-up after end of treatment: none Control • Description: Treatment as usual • Length of treatment: 2 weeks • Longest follow-up after end of treatment: none
Outcomes	Agitation (ABMI), SD ● Outcome type: ContinuousOutcome
Identification	
Notes	

Risk of bias table

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Judgement Comment: Participants were randomised either by units (for larger nursing homes with many eligible participants) or by nursing homes (when there were fewer eligible participants). Randomisation was performed using random numbers via a ration of 1.5:1
Allocation concealment (selection bias)	High risk	Judgement Comment: The research assistents were blind to group allocation, until the treatment began
Blinding of participants and personnel (performance bias)	Low risk	Judgement Comment: participants were blinded, but not the personnel
Blinding of outcome assessment (detection bias)	High risk	Judgement Comment: Blinding not possible

Incomplete outcome data (attrition bias)		Judgement Comment: 62 participants withdrew from the intervention group and 36 from the placebo group. However, as ITT analyses was performed, the risk of bias is considered low. No apparent sources of bias
Selective reporting (reporting bias)	Low risk	Judgement Comment: Matches study protocol
Other bias	Low risk	Judgement Comment: No apparent sources of bias

Gonyea 2006

Methods	
Participants	
Interventions	
Outcomes	
Identification	
Notes	Data obtained from: Moniz, Cook E.; Swift K.; James I.; Malouf R.; De, Vugt M.; Verhey F. Functional analysis-based interventions for challenging behaviour in dementiaCochrane database of systematic reviews (Online) 2012;2(Journal Article):CD006929United Kingdom 2012

Risk of bias table

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	reference: Moniz Cook et al. 2012
Allocation concealment (selection bias)	Low risk	reference: Moniz Cook et al. 2012
Blinding of participants and personnel (performance bias)	Unclear risk	reference: Moniz Cook et al. 2012
Blinding of outcome assessment (detection bias)	Unclear risk	reference: Moniz Cook et al. 2012
Incomplete outcome data (attrition bias)	Low risk	reference: Moniz Cook et al. 2012
Selective reporting (reporting bias)	Unclear risk	reference: Moniz Cook et al. 2012
Other bias	Unclear risk	reference: Moniz Cook et al. 2012

Gormley 2001

Methods	
Participants	
Interventions	
Outcomes	
Identification	
Notes	Data obtained from: Moniz, Cook E.; Swift K.; James I.; Malouf R.; De, Vugt M.; Verhey F. Functional analysis-based interventions for challenging behaviour in dementiaCochrane database of systematic reviews (Online) 2012;2(Journal Article):CD006929United Kingdom 2012

Risk of bias table

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	reference: Moniz Cook et al. 2012
Allocation concealment (selection bias)	Low risk	reference: Moniz Cook et al. 2012
Blinding of participants and personnel (performance bias)	Unclear risk	reference: Moniz Cook et al. 2012
Blinding of outcome assessment (detection bias)	Low risk	reference: Moniz Cook et al. 2012
Incomplete outcome data (attrition bias)	Low risk	reference: Moniz Cook et al. 2012
Selective reporting (reporting bias)	Low risk	reference: Moniz Cook et al. 2012
Other bias	Unclear risk	reference: Moniz Cook et al. 2012

Huang 2003

Methods	
Participants	
Interventions	
Outcomes	
Identification	
Notes	Data obtained from: Moniz, Cook E.; Swift K.; James I.; Malouf R.; De, Vugt M.; Verhey F. Functional analysis-based interventions for challenging behaviour in dementiaCochrane database of systematic reviews (Online) 2012;2(Journal Article):CD006929United Kingdom 2012

Risk of bias table

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	reference: Moniz Cook et al. 2012
Allocation concealment (selection bias)	Unclear risk	reference: Moniz Cook et al. 2012
Blinding of participants and personnel (performance bias)	Low risk	reference: Moniz Cook et al. 2012
Blinding of outcome assessment (detection bias)	Unclear risk	reference: Moniz Cook et al. 2012
Incomplete outcome data (attrition bias)	Low risk	reference: Moniz Cook et al. 2012
Selective reporting (reporting bias)	Low risk	reference: Moniz Cook et al. 2012
Other bias	Unclear risk	reference: Moniz Cook et al. 2012

Kovach 2006

Methods	Study design: Randomized controlled trial
	Study grouping: Parallel group

Participants	Baseline Characteristics Intervention • Age mean (SD): 7.35 • MMSE: 8.26 Control • Age mean (SD): 6.13 • MMSE: 6.29 Included criteria: MMSE score indicating moderate to severe cognitive impairment. Advanced funcitonal impairment. No chronic psychiatric diagnosi. At least 4 weeks postadmission to skilled nursing care at this nursing home
Interventions	Intervention Characteristics Intervention • Description: Serial trial intervention • Length of treatment: 2 weeks • Longest follow-up after end of treatment: 4 weeks Control • Description: Standard care • Length of treatment: 2 weeks • Longest follow-up after end of treatment: 4 weeks
Outcomes	BPSD (BEHAVE-AD), SD ● Outcome type: ContinuousOutcome
Identification	
Notes	

Risk of bias table

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Judgement Comment: Randomly assigned using coin toss
Allocation concealment (selection bias)	High risk	Judgement Comment: Randomly assigned using coin toss
Blinding of participants and personnel (performance bias)	Low risk	Judgement Comment: Research subjects described as blinded
Blinding of outcome assessment (detection bias)	Low risk	Judgement Comment: Data collectors described as blinded
Incomplete outcome data (attrition bias)	Low risk	Judgement Comment:127 ramdomized and 114 completed
Selective reporting (reporting bias)	Low risk	Judgement Comment: None detected
Other bias	Low risk	Judgement Comment: No other apparent sources of bias

McCabe 2015

Methods	Study design: Randomized controlled trial Study grouping: Parallel group
Participants	Baseline Characteristics Intervention 1 ● Age y (SD): 82.85 (8.45)
	Control • Age y (SD): 81.25 (11.03)
	Included criteria: The inclusion crite-ria required a positive diagnosis of dementia, and thepresence of at least one challenging behavior, defined as 'any behavior associated with dementia which causes dis-tress or danger to the person with dementia and/or others' (Bird et al., 2009).
Interventions	Intervention • Description: Training. A two-hour training session in which staff were helped towork through and identify probably causal factors for thebehavior of residents, and develop potential ways of ame-liorating these causes (training/support and trainingconditions). • Length of treatment: 3 months • Longest follow-up after end of treatment: 6 months

	Control • Description: Care as usual • Length of treatment: 3 months • Longest follow-up after end of treatment: 6 months
Outcomes	Agitation (CMAI), SD ● Outcome type: ContinuousOutcome
Identification	
Notes	

Risk of bias table

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Quote: "the study agreed to participate. Facilities were randomized to one of the four study/intervention conditions using a clus- ter randomized controlled design (i.e., the facility rather than the participants/residents or staff were the unit of ran- domization). Randomization occurred by facilities being allocated to one of the conditions as they were recruited into the study. The numbers 1, 2, 3, and 4 were placed in a box (in each of the two locations) in both year 1 and year 2. The number that was drawn out for the facility deter- mined which of the four conditions the facility was allo- cated to. /b> Aged-care residents were recruited through
Allocation concealment (selection bias)	Low risk	Quote: "were recruited into the study. The numbers 1, 2, 3, and 4 were placed in a box (in each of the two locations) in both year 1 and year 2. The number that was drawn out for the facility deter- mined which of the four conditions the facility was allo- cated to. Aged-care residents were recruited through"
Blinding of participants and personnel (performance bias)	Unclear risk	Judgement Comment: Nothing mentioned
Blinding of outcome assessment (detection bias)	Unclear risk	Judgement Comment: Nothing mentioned
Incomplete outcome data (attrition bias)	High risk	Judgement Comment: Number of withdrawers are not described

Selective reporting (reporting bias)	Low risk	Judgement Comment: No apparent sources of bias
Other bias	Low risk	Judgement Comment: No other apparent sources of bias

Pieper 2016

Methods	Study design: Cluster randomized controlled trial Study grouping: Parallel group		
Participants	Baseline Characteristics Intervention ● Age y (SD): 84.3 (7.4)		
	Control ■ Age y (SD): 83.3(6.9)		
	Included criteria: et thecriteria, that at least one psychogeriatric unit was willing toparticipate and no major organizational changes or buildingactivities were planned or performed during the study per-iod. In each nursing home, residents with moderate to sev-ere cognitive impairment (Reisberg Global DeteriorationScale (GDS) Stage 5, 6, or 7),15no psychiatric diagnosisother than dementia, and clinically significant symptoms ofchallenging behavior (Neuropsychiatric Inventory—NursingHome version (NPI-NH) score>4 or Cohen-Mansfield Agi-tation Inventory (CMAI) score>44)16,17were eligible for participation, providing that written proxy consent wasreceived		
Interventions	Intervention Characteristics Intervention • Description: STA OP! • Length of treatment: 3 months • Longest follow-up after end of treatment: 6 months Control • Description: Treatment as usual		
	 Length of treatment: 3 months Longest follow-up after end of treatment: 6 months 		

Outcomes	BPSD (NPI) CI ● Outcome type: ContinuousOutcome
	Antipsychotic usage, OR ● Outcome type: DichotomousOutcome
	Depression (Cornell), CI ● Outcome type: ContinuousOutcome
	Agitation (CMAI), CI ● Outcome type: ContinuousOutcome
Identification	
Notes	

Risk of bias table

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Quote: "An independent researcher (who was unaware of the identity of the units) performed the allocation using a computer-generated sequence program"
Allocation concealment (selection bias)	Low risk	Quote: "An independent researcher (who was unaware of the identity of the units) performed the allocation using a computer-generated sequence program (Random"
Blinding of participants and personnel (performance bias)	High risk	Quote: "The trial was single blinded (the researcher knew the condi- tion, but the research assistants performing the measure- ments were blinded)." Judgement Comment: Participants were not blinded
Blinding of outcome assessment (detection bias)	Low risk	Quote: "The trial was single blinded (the researcher knew the condi- tion, but the research assistants performing the measure- ments were blinded). 13 Residents"
Incomplete outcome data (attrition bias)	Low risk	Judgement Comment: No apparent sources of bias

Selective reporting (reporting bias)		Quote: "This trial is registered at the Netherlands National Trial Register (NTR1967)." Judgement Comment: According to the protocol, Quality of Life should have been measured using Qualidem. There is no reportings on Qualidem in thi study
Other bias	Low risk	Judgement Comment: No apparent sources of bias

Proctor 1999

Methods	
Participants	
Interventions	
Outcomes	
Identification	
Notes	Data obtained from: Moniz, Cook E.; Swift K.; James I.; Malouf R.; De, Vugt M.; Verhey F. Functional analysis-based interventions for challenging behaviour in dementiaCochrane database of systematic reviews (Online) 2012;2(Journal Article):CD006929United Kingdom 2012

Risk of bias table

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	reference: Moniz Cook et al. 2012
Allocation concealment (selection bias)	Low risk	reference: Moniz Cook et al. 2012
Blinding of participants and personnel (performance bias)	Low risk	reference: Moniz Cook et al. 2012
Blinding of outcome assessment (detection bias)	Unclear risk	reference: Moniz Cook et al. 2012
Incomplete outcome data (attrition bias)	Low risk	reference: Moniz Cook et al. 2012
Selective reporting (reporting bias)	Low risk	reference: Moniz Cook et al. 2012
Other bias	Unclear risk	reference: Moniz Cook et al. 2012

Teri 2005a

Methods	
Participants	
Interventions	
Outcomes	
Identification	
Notes	Data obtained from: Moniz, Cook E.; Swift K.; James I.; Malouf R.; De, Vugt M.; Verhey F. Functional analysis-based interventions for challenging behaviour in dementiaCochrane database of systematic reviews (Online) 2012;2(Journal Article):CD006929United Kingdom 2012 Outcomes Only total no. of patient was reported (n=31). For the analysis we assumed that there was 16 in the intervention og 15 in the control group

Risk of bias table

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	reference: Moniz Cook et al. 2012
Allocation concealment (selection bias)	Unclear risk	reference: Moniz Cook et al. 2012
Blinding of participants and personnel (performance bias)	Unclear risk	reference: Moniz Cook et al. 2012
Blinding of outcome assessment (detection bias)	Low risk	reference: Moniz Cook et al. 2012
Incomplete outcome data (attrition bias)	Low risk	reference: Moniz Cook et al. 2012
Selective reporting (reporting bias)	Low risk	reference: Moniz Cook et al. 2012
Other bias	Unclear risk	reference: Moniz Cook et al. 2012

Teri 2005b

Methods	
Participants	
Interventions	
Outcomes	
Identification	
Notes	Data obtained from: Moniz, Cook E.; Swift K.; James I.; Malouf R.; De, Vugt M.; Verhey F. Functional analysis-based interventions for challenging behaviour in dementiaCochrane database of systematic reviews (Online) 2012;2(Journal Article):CD006929United Kingdom 2012

Risk of bias table

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	reference: Moniz Cook et al. 2012
Allocation concealment (selection bias)	Unclear risk	reference: Moniz Cook et al. 2012
Blinding of participants and personnel (performance bias)	Unclear risk	reference: Moniz Cook et al. 2012
Blinding of outcome assessment (detection bias)	Low risk	reference: Moniz Cook et al. 2012
Incomplete outcome data (attrition bias)	Low risk	reference: Moniz Cook et al. 2012
Selective reporting (reporting bias)	Unclear risk	reference: Moniz Cook et al. 2012
Other bias	Low risk	reference: Moniz Cook et al. 2012

Footnotes

Characteristics of excluded studies

Ballard 2016

Reason for exclusion	Wrong intervention

Ballard 2017

Reason for exclusion	Wrong intervention	
----------------------	--------------------	--

Ballard 2017a

December evaluation	A hadrond make
Reason for exclusion	Abstract only

Dichter 2015

Reason for exclusion	Wrong patient population
neason for exclusion	Wrong patient population

Farran 2004

Reason for exclusion	Wrong intervention
----------------------	--------------------

Fernandez Calvo 2015

Reason for exclusion	Wrong patient population
----------------------	--------------------------

Fossey 2006

Reason for exclusion	Wrong patient population
----------------------	--------------------------

Gitlin 2003

Reason for exclusion	Wrong intervention
----------------------	--------------------

Gitlin 2010

Re	eason for exclusion	Wrong outcomes
		,

Goga 2017

December of the construction	Maranan linkan namblan	
Reason for exclusion	Wrong intervention	
1.00001. 10. 02.01001011	Triong intol volution	

Halek 2013

Reason for exclusion	Wrong study design	
I I Casoli Ioi Exclusion	I WIOLIG Study acsign	
		l l

LosadaBaltar 2004

Reason for exclusion	Not in English
----------------------	----------------

Mador 2004

Reason for exclusion Wrong patient population	Reason for exclusion
---	----------------------

Moniz Cook 2008

Reason for exclusion	Wrong patient population

Reisberg 2015

Reason for exclusion	Wrong study design

Sampson 2011

Reason for exclusion	Wrong intervention
----------------------	--------------------

Straubmeier 2017

Reason for exclusion	Wrong intervention	
----------------------	--------------------	--

Teri 2000

		,
December for evaluation	Musican intervioletica	,
Reason for exclusion	Wrong intervention	,
	1 · · · · · · · · · · · · · · · · · · ·	,

Teri 2003

Reason for exclusion	Wrong intervention
----------------------	--------------------

Thyrian 2017

Reason for exclusion	Wrong patient population
----------------------	--------------------------

van 2013

Reason for exclusion Wrong patient population	
---	--

VanHaitsma 2015

Reason for exclusion	Wrong intervention

Weiner 2002

Decree for controlled	W
Reason for exclusion	Wrong outcomes

Zarit 1987

Reason for exclusion	Wrong patient population
----------------------	--------------------------

Zwijsen 2013

	At a second	
Reason for exclusion	Abstract only	
ricusori for exolusion	7 loot act only	

Zwijsen 2014

December ovelveion	Ale atmost and control
Reason for exclusion	Abstract only
	· ··································

Footnotes

References to studies

Included studies

Burgio 2003

Burgio, L.; Stevens, A.; Guy, D.; Roth, D. L.; Haley, W. E.. Impact of two psychosocial interventions on white and African American family caregivers of individuals with dementia. The Gerontologist 2003;43(4):568-579. [DOI:]

Chenoweth 2009

Chenoweth, L.; King, M. T.; Jeon, Y. H.; Brodaty, H.; Stein-Parbury, J.; Norman, R.; Haas, M.; Luscombe, G.. Caring for Aged Dementia Care Resident Study (CADRES) of person-centred care, dementia-care mapping, and usual care in dementia: a cluster-randomised trial. The Lancet.Neurology 2009;8(4):317-325. [DOI: 10.1016/S1474-4422(09)70045-6 [doi]]

Cohen Mansfield 2012

Cohen-Mansfield, Jiska; Thein, Khin; Marx, Marcia S.; Dakheel-Ali, Maha; Freedman, Laurence. Efficacy of nonpharmacologic interventions for agitation in advanced dementia: a randomized, placebo-controlled trial. The Journal of clinical psychiatry 2012;73(9):1255-61. [DOI: https://dx.doi.org/10.4088/JCP.12m07918]

Gonyea 2006

Gonyea, J. G.; O'Connor, M. K.; Boyle, P. A.. Project CARE: a randomized controlled trial of a behavioral intervention group for Alzheimer's disease caregivers. The Gerontologist 2006;46(6):827-832. [DOI: 46/6/827 [pii]]

Gormley 2001

Gormley, N.; Lyons, D.; Howard, R.. Behavioural management of aggression in dementia: a randomized controlled trial. Age and Ageing 2001;30(2):141-145. [DOI:]

Huang 2003

Huang, H. L.; Shyu, Y. I.; Chen, M. C.; Chen, S. T.; Lin, L. C.. A pilot study on a home-based caregiver training program for improving caregiver self-efficacy and decreasing the behavioral problems of elders with dementia in Taiwan. International journal of geriatric psychiatry 2003;18(4):337-345. [DOI: 10.1002/gps.835 [doi]]

Kovach 2006

Kovach, C. R.; Logan, B. R.; Noonan, P. E.; Schlidt, A. M.; Smerz, J.; Simpson, M.; Wells, T.. Effects of the Serial Trial Intervention on discomfort and behavior of nursing home residents with dementia. American Journal of Alzheimer's Disease and Other Dementias 2006;21(3):147-155. [DOI: 10.1177/1533317506288949 [doi]]

McCabe 2015

McCabe, M. P.; Bird, M.; Davison, T. E.; Mellor, D.; MacPherson, S.; Hallford, D.; Seedy, M.. An RCT to evaluate the utility of a clinical protocol for staff in the management of behavioral and psychological symptoms of dementia in residential aged-care settings. Aging & mental health 2015;19(9):799-807. [DOI: 10.1080/13607863.2014.967659 [doi]]

Pieper 2016

Pieper, Marjoleine J. C.; Francke, Anneke L.; van der Steen, Jenny T.; Scherder, Erik J. A.; Twisk, Jos W. R.; Kovach, Christine R.; Achterberg, Wilco P.. Effects of a Stepwise Multidisciplinary Intervention for Challenging Behavior in Advanced Dementia: A Cluster Randomized Controlled Trial. Journal of the American Geriatrics Society 2016;64(2):261-269. [DOI: https://dx.doi.org/10.1111/jgs.13868]

Proctor 1999

Proctor, R.; Burns, A.; Powell, H. S.; Tarrier, N.; Faragher, B.; Richardson, G.; Davies, L.; South, B. Behavioural management in nursing and residential homes: a randomised controlled trial. Lancet (London, England) 1999;354(9172):26-29. [DOI: S0140-6736(98)08237-3 [pii]]

Teri 2005a

Teri, L.; Huda, P.; Gibbons, L.; Young, H.; van Leynseele, J.. STAR: a dementia-specific training program for staff in assisted living residences. The Gerontologist 2005;45(5):686-693. [DOI: 45/5/686 [pii]]

Teri 2005b

[Empty]

Excluded studies

Ballard 2016

Ballard, Clive; Orrell, Martin; YongZhong, Sun; Moniz-Cook, Esme; Stafford, Jane; Whittaker, Rhiannon; Woods, Bob; Corbett, Anne; Garrod, Lucy; Khan, Zunera; Woodward-Carlton, Barbara; Wenborn, Jennifer; Fossey, Jane. Impact of antipsychotic review and nonpharmacological intervention on antipsychotic use, neuropsychiatric symptoms, and mortality in people with dementia living in nursing homes: A factorial cluster-randomized controlled trial by the Well-Being and Health for People with Dementia (WHELD) program. The American Journal of Psychiatry 2016;173(3):252-262. [DOI:]

Ballard 2017

Ballard, Clive; Orrell, Martin; Sun, Yongzhong; Moniz-Cook, Esme; Stafford, Jane; Whitaker, Rhiannon; Woods, Bob; Corbett, Anne; Banerjee, Sube; Testad, Ingelin; Garrod, Lucy; Khan, Zunera; Woodward-Carlton, Barbara; Wenborn, Jennifer; Fossey, Jane. Impact of antipsychotic review and non-pharmacological intervention on health-related quality of life in people with dementia living in care homes: WHELD-a factorial cluster randomised controlled trial. International journal of geriatric psychiatry 2017;32(10):1094-1103. [DOI: https://dx.doi.org/10.1002/gps.4572]

Ballard 2017a

Ballard C.; Fossey J.; Corbett A.; Orrell M.; Romeo R.; Moniz-Cook E.; Woods B.; Whitaker R. Impact of wheld intervention on neuropsychiatric symptoms, antipsychotic use and quality of life in people with dementia living in nursing homes: Acluster randomized trial. Alzheimer's and Dementia 2017;13(7):P171. [DOI:]

Dichter 2015

Dichter M.N.; Quasdorf T.; Schwab C.G.G.; Trutschel D.; Haastert B.; Riesner C.; Bartholomeyczik S.; Halek M.. Dementia care mapping: Effects on residents' quality of life and challenging behavior in German nursing homes. A quasi-experimental trial. International Psychogeriatrics 2015;27(11):1875-1892. [DOI: http://dx.doi.org/10.1017/S1041610215000927]

Farran 2004

Farran, CJ; Gilley, DW; McCann, JJ; Bienias, JL; Lindeman, DA; Evans, DA. Psychosocial interventions to reduce depressive symptoms of dementia caregivers: A randomized clinical trial comparing two approaches. Journal of Mental Health and Aging 2004;10(4):337-350. [DOI:]

Fernandez Calvo 2015

Fernandez-Calvo, Bernardino; Contador, Israel; Ramos, Francisco; Olazaran, Javier; Mograbi, Daniel C.; Morris, Robin G.. Effect of unawareness on rehabilitation outcome in a randomised controlled trial of multicomponent intervention for patients with mild Alzheimer's disease. Neuropsychological rehabilitation 2015;25(3):448-77. [DOI: https://dx.doi.org/10.1080/09602011.2014.948461]

Fossey 2006

Fossey, J.; Ballard, C.; Juszczak, E.; James, I.; Alder, N.; Jacoby, R.; Howard, R. Effect of enhanced psychosocial care on antipsychotic use in nursing home residents with severe dementia: cluster randomised trial. BMJ (Clinical research ed.) 2006;332(7544):756-761. [DOI: bmj.38782.575868.7C [pii]]

Gitlin 2003

Gitlin, L. N.; Winter, L.; Corcoran, M.; Dennis, M. P.; Schinfeld, S.; Hauck, W. W.. Effects of the home environmental skill-building program on the caregiver-care recipient dyad: 6-month outcomes from the Philadelphia REACH Initiative. The Gerontologist 2003;43(4):532-546. [DOI:]

Gitlin 2010

Gitlin, L. N.; Winter, L.; Dennis, M. P.; Hodgson, N.; Hauck, W. W.. Targeting and managing behavioral symptoms in individuals with dementia: a randomized trial of a nonpharmacological intervention. Journal of the American Geriatrics Society 2010;58(8):1465-1474. [DOI: 10.1111/j.1532-5415.2010.02971.x [doi]]

Goga 2017

Goga, Joshana K.; Depaolo, Antonio; Khushalani, Sunil; Walters, J. K.; Roca, Robert; Zisselman, Marc; Borleis, Christopher. Lean Methodology Reduces Inappropriate Use of Antipsychotics for Agitation at a Psychiatric Hospital. The Consultant pharmacist: the journal of the American Society of Consultant Pharmacists 2017;32(1):54-62. [DOI: https://dx.doi.org/10.4140/TCP.n.2017.54]

Halek 2013

Halek, Margareta; Dichter, Martin Nikolaus; Quasdorf, Tina; Riesner, Christine; Bartholomeyczik, Sabine. The effects of dementia care mapping on nursing home residents' quality of life and staff attitudes: design of the quasi-experimental study Leben-QD II. BMC geriatrics 2013;13(Journal Article):53. [DOI: https://dx.doi.org/10.1186/1471-2318-13-53]

LosadaBaltar 2004

Losada Baltar, A.; Izal Fernandez de Troconiz, M.; Montorio Cerrato, I.; Marquez Gonzalez, M.; Perez Rojo, G. Differential efficacy of two psychoeducational interventions for dementia family caregivers. Revista de neurologia 2004;38(8):701-708. [DOI: rn2003522 [pii]]

Mador 2004

Mador, J. E.; Giles, L.; Whitehead, C.; Crotty, M.. A randomized controlled trial of a behavior advisory service for hospitalized older patients with confusion. International journal of geriatric psychiatry 2004;19(9):858-863. [DOI: 10.1002/gps.1165 [doi]]

Moniz Cook 2008

Moniz-Cook, E.; Elston, C.; Gardiner, E.; Agar, S.; Silver, M.; Win, T.; Wang, M.. Can training community mental health nurses to support family carers reduce behavioural problems in dementia? An exploratory pragmatic randomised controlled trial. International journal of geriatric psychiatry 2008;23(2):185-191. [DOI: 10.1002/gps.1860 [doi]]

Reisberg 2015

Reisberg B.; Monteiro I.; Torossian C.; Xu J.; Janjua K.; Ghimire S.; Sommese K.; Kenowsky S.. Effects of a comprehensive, individualized person-centered management program on persons with moderately severe Alzheimer's disease: A randomized controlled trial-comprehensive study findings. Alzheimer's and Dementia 2015;11(7):P608-P609. [DOI:]

Sampson 2011

Sampson, Elizabeth L.; Jones, Louise; Thuné-Boyle, Ingela, C.V.; Kukkastenvehmas, Riitta; King, Michael; Leurent, Baptiste; Tookman, Adrian; Blanchard, Martin R.. Palliative assessment and advance care planning in severe dementia: An exploratory randomized controlled trial of a complex intervention. Palliative medicine 2011;25(3):197-209. [DOI: 10.1177/0269216310391691]

Straubmeier 2017

Straubmeier, Melanie; Behrndt, Elisa-Marie; Seidl, Hildegard; Ozbe, Dominik; Luttenberger, Katharina; Graessel, Elmar. Non-Pharmacological Treatment in People With Cognitive Impairment. Deutsches Arzteblatt international 2017;114(48):815-821. [DOI: https://dx.doi.org/10.3238/arztebl.2017.0815]

Teri 2000

Teri, L.; Logsdon, R. G.; Peskind, E.; Raskind, M.; Weiner, M. F.; Tractenberg, R. E.; Foster, N. L.; Schneider, L. S.; Sano, M.; Whitehouse, P.; Tariot, P.; Mellow, A. M.; Auchus, A. P.; Grundman, M.; Thomas, R. G.; Schafer, K.; Thal, L. J.; Alzheimer's Disease Cooperative Study. Treatment of agitation in AD: a randomized, placebo-controlled clinical trial. Neurology 2000;55(9):1271-1278. [DOI:]

Teri 2003

Teri, L.; Gibbons, L. E.; McCurry, S. M.; Logsdon, R. G.; Buchner, D. M.; Barlow, W. E.; Kukull, W. A.; LaCroix, A. Z.; McCormick, W.; Larson, E. B. Exercise plus behavioral management in patients with Alzheimer disease: a randomized controlled trial. Jama 2003;290(15):2015-2022. [DOI: 10.1001/jama.290.15.2015 [doi]]

Thyrian 2017

Thyrian, Jochen Rene; Hertel, Johannes; Wucherer, Diana; Eichler, Tilly; Michalowsky, Bernhard; Dreier-Wolfgramm, Adina; Zwingmann, Ina; Kilimann, Ingo; Teipel, Stefan; Hoffmann, Wolfgang. Effectiveness and Safety of Dementia Care Management in Primary Care: A Randomized Clinical Trial. JAMA psychiatry 2017;74(10):996-1004. [DOI: https://dx.doi.org/10.1001/jamapsychiatry.2017.2124]

van 2013

van, de Ven; Draskovic, Irena; Adang, Eddy M. M.; Donders, Rogier; Zuidema, Sytse U.; Koopmans, Raymond T. C. M.; Vernooij-Dassen, Myrra. Effects of dementia-care mapping on residents and staff of care homes: a pragmatic cluster-randomised controlled trial. PloS one 2013;8(7):e67325. [DOI: https://dx.doi.org/10.1371/journal.pone.0067325]

VanHaitsma 2015

Van Haitsma, Kimberly S.; Curyto, Kimberly; Abbott, Katherine M.; Towsley, Gail L.; Spector, Abby; Kleban, Morton. A Randomized Controlled Trial for an Individualized Positive Psychosocial Intervention for the Affective and Behavioral Symptoms of Dementia in Nursing Home Residents. Journals of Gerontology Series B: Psychological Sciences & Social Sciences 2015;70(1):35-45. [DOI:]

Weiner 2002

Weiner, M. F.; Tractenberg, R. E.; Sano, M.; Logsdon, R.; Teri, L.; Galasko, D.; Gamst, A.; Thomas, R.; Thal, L. J.. No long-term effect of behavioral treatment on psychotropic drug use for agitation in Alzheimer's disease patients. Journal of geriatric psychiatry and neurology 2002;15(2):95-98. [DOI: 10.1177/089198870201500208 [doi]]

Zarit 1987

Zarit, S. H.; Anthony, C. R.; Boutselis, M.. Interventions with care givers of dementia patients: comparison of two approaches. Psychology and aging 1987;2(3):225-232. [DOI:]

Zwijsen 2013

Zwijsen S.; Hertogh C.; Smalbrugge M.; Gerritsen D.; Eefsting J.; Margriet, Pot A.. Grip on challenging behaviour: Effects of a structured multidisciplinary care program for management of challenging behaviour on dementia special care units. International Psychogeriatrics 2013;25(Journal Article):S65-S66. [DOI: http://dx.doi.org/10.1017/S1041610213002147]

Zwijsen 2014

Zwijsen, Sandra A.; Smalbrugge, Martin; Eefsting, Jan A.; Twisk, Jos W. R.; Gerritsen, Debby L.; Pot, Anne Margriet; Hertogh, Cees M. P. M.. Coming to grips with challenging behavior: a cluster randomized controlled trial on the effects of a multidisciplinary care program for challenging behavior in dementia. Journal of the

American Medical Directors Association 2014;15(7):531.e1-531.e10. [DOI: https://dx.doi.org/10.1016/j.jamda.2014.04.007]

Other references

Additional references

Other published versions of this review

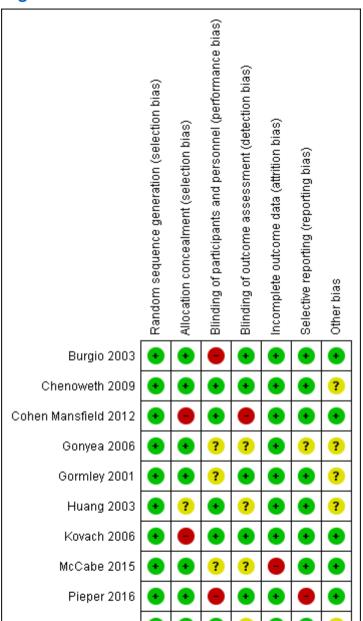
Data and analyses

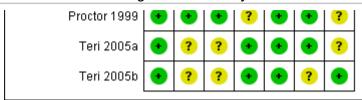
2 Functional analysis vs Control, Longest FU, min 4 wk, max 6 mo

Outcome or Subgroup	Studies	Participants	Statistical Method	Effect Estimate
2.1 Antipsychotic usage	2	388	Risk Ratio (IV, Random, 95% CI)	1.00 [0.55, 1.81]
2.2 BPSD	9	990	Std. Mean Difference (IV, Random, 95% CI)	-0.25 [-0.38, -0.13]
2.3 Restraint	0		Risk Ratio (IV, Fixed, 95% CI)	No totals
2.4 Agitation	5	717	Std. Mean Difference (IV, Random, 95% CI)	-0.43 [-0.74, -0.11]
2.5 Quality of life	2	242	Std. Mean Difference (IV, Random, 95% CI)	-0.06 [-0.32, 0.19]
2.6 ADL	1	105	Mean Difference (IV, Fixed, 95% CI)	-2.20 [-6.41, 2.01]
2.7 Depression	4	542	Std. Mean Difference (IV, Random, 95% CI)	-0.22 [-0.55, 0.12]

Figures

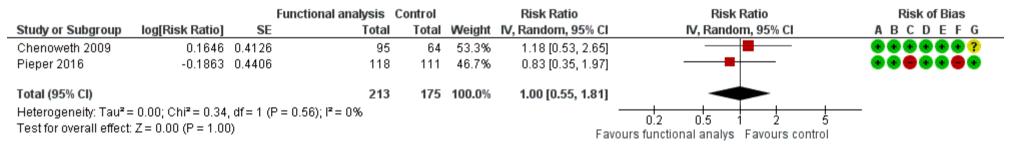
Figure 1





Risk of bias summary: review authors' judgements about each risk of bias item for each included study.

Figure 2 (Analysis 2.1)

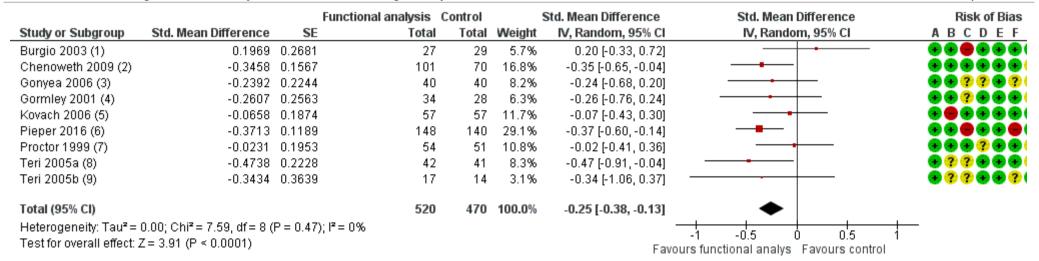


Risk of bias legend

- (A) Random sequence generation (selection bias)
- (B) Allocation concealment (selection bias)
- (C) Blinding of participants and personnel (performance bias)
- (D) Blinding of outcome assessment (detection bias)
- (E) Incomplete outcome data (attrition bias)
- (F) Selective reporting (reporting bias)
- (G) Other bias

Forest plot of comparison: 2 Functional analysis vs Control, Longest FU, min 4 wk, max 6 mo, outcome: 2.1 Antipsychotic usage.

Figure 3 (Analysis 2.2)



Footnotes

- (1) RMBPC
- (2) NPI
- (3) NPI
- (4) BEHAVE-AD
- (5) BEHAVE-AD
- (6) NPI
- (7) Crichton Royal behavioural rating scale
- (8) RMBPC
- (9) NPI

Risk of bias legend

- (A) Random sequence generation (selection bias)
- (B) Allocation concealment (selection bias)
- (C) Blinding of participants and personnel (performance bias)
- (D) Blinding of outcome assessment (detection bias)
- (E) Incomplete outcome data (attrition bias)
- (F) Selective reporting (reporting bias)
- (G) Other bias

Forest plot of comparison: 2 Functional analysis vs Control, Longest FU, min 4 wk, max 6 mo, outcome: 2.2 BPSD.

Figure 4 (Analysis 2.4)

(E) Incomplete outcome data (attrition bias) (F) Selective reporting (reporting bias)

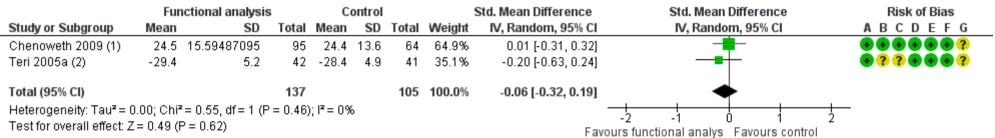
(G) Other bias

			Functional analysis	Control		Std. Mean Difference	Std. Mean Difference	Risk of E
Study or Subgroup	Std. Mean Difference	SE	Total	Total	Weight	IV, Random, 95% CI	IV, Random, 95% CI	ABCDE
Chenoweth 2009 (1)	-0.3436	0.1567	101	70	22.5%	-0.34 [-0.65, -0.04]	-	$\bullet \bullet \bullet \bullet \bullet$
Cohen Mansfield 2012 (2)	-1.0844	0.2095	89	36	19.3%	-1.08 [-1.50, -0.67]		$\bullet \bullet \bullet \bullet \bullet$
Huang 2003 (3)	-0.23	0.2897	24	24	14.9%	-0.23 [-0.80, 0.34]		? • ? •
McCabe 2015 (4)	-0.3573	0.2206	48	37	18.6%	-0.36 [-0.79, 0.08]		??
Pieper 2016 (5)	-0.1567	0.1181	148	140	24.7%	-0.16 [-0.39, 0.07]	 +	$\bullet \bullet \bullet \bullet \bullet$
Total (95% CI)			410	307	100.0%	-0.43 [-0.74, -0.11]	•	
Heterogeneity: Tau² = 0.09; C	chi ² = 15.13, df = 4 (P = 0	0.004); <mark>P</mark>	= 74%					_
Test for overall effect: Z = 2.6	3 (P = 0.009)					Fa	vours fuctional analysi Favours control	
<u>Footnotes</u>							Risk of bias legend	
(1) CMAI							(A) Random sequence generation (selec	tion bias)
(2) ABMI							(B) Allocation concealment (selection bia	s)
(3) CMAI							(C) Blinding of participants and personne	l (performance bia
(4) CMAI							(D) Blinding of outcome assessment (det	ection bias)

Forest plot of comparison: 2 Functional analysis vs Control, Longest FU, min 4 wk, max 6 mo, outcome: 2.4 Agitation.

Figure 5 (Analysis 2.5)

(4) CMAI (5) CMAI



Test for overall effect: Z = 0.49 (P = 0.62)

Footnotes

- (1) QUALID
- (2) QoL-AD proxy reported

Risk of bias legend

- (A) Random sequence generation (selection bias)
- (B) Allocation concealment (selection bias)
- (C) Blinding of participants and personnel (performance bias)
- (D) Blinding of outcome assessment (detection bias)
- (E) Incomplete outcome data (attrition bias)
- (F) Selective reporting (reporting bias)
- (G) Other bias

Forest plot of comparison: 2 Functional analysis vs Control, Longest FU, min 4 wk, max 6 mo, outcome: 2.5 Quality of life.

Figure 6 (Analysis 2.6)

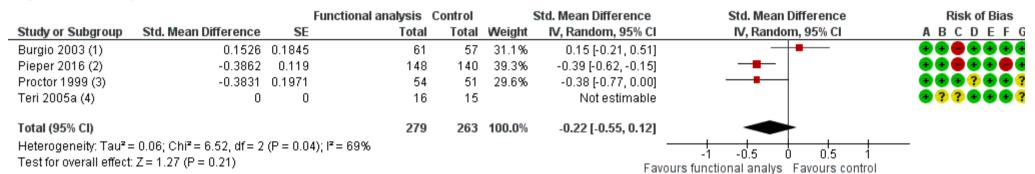
	Fur	nctional analysis	6		Control			Mean Difference		Mean Difference	Risk of Bias
Study or Subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV, Fixed, 95% CI		IV, Fixed, 95% CI	ABCDEFG
Proctor 1999	-13.2	13.87211028	54	-11	7.28717187	51	100.0%	-2.20 [-6.41, 2.01]			• • • ? • • ?
Total (95% CI)			54			51	100.0%	-2.20 [-6.41, 2.01]			
Heterogeneity: Not ap Test for overall effect:	•							Fav	-10 ours fun	-5 0 5	10

Risk of bias legend

- (A) Random sequence generation (selection bias)
- (B) Allocation concealment (selection bias)
- (C) Blinding of participants and personnel (performance bias)
- (D) Blinding of outcome assessment (detection bias)
- (E) Incomplete outcome data (attrition bias)
- (F) Selective reporting (reporting bias)
- (G) Other bias

Forest plot of comparison: 2 Functional analysis vs Control, Longest FU, min 4 wk, max 6 mo, outcome: 2.6 ADL.

Figure 7 (Analysis 2.7)



Footnotes

- (1) CES-D. Data for White and African American have been pooled.
- (2) Cornell (CSDD)
- (3) AGECAT, depression subscale
- (4) GDS

Risk of bias legend

- (A) Random sequence generation (selection bias)
- (B) Allocation concealment (selection bias)
- (C) Blinding of participants and personnel (performance bias)
- (D) Blinding of outcome assessment (detection bias)
- (E) Incomplete outcome data (attrition bias)
- (F) Selective reporting (reporting bias)
- (G) Other bias

Forest plot of comparison: 2 Functional analysis vs Control, Longest FU, min 4 wk, max 6 mo, outcome: 2.7 Depression.