Interventions with potential to reduce sedentary time in adults: systematic review and meta-analysis

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ABSTRACT

Context Time spent in sedentary behaviours (SB) is associated with poor health, irrespective of the level of physical activity. The aim of this study was to evaluate the effect of interventions which included SB as an outcome measure in adults.

Methods Thirteen databases, including The Cochrane Library, MEDLINE and SPORTDiscus, trial registers and reference lists, were searched for randomised controlled trials until January 2014. Study selection, data extraction and quality assessment were performed independently. Primary outcomes included SB, proxy measures of SB and patterns of accumulation of SB. Secondary outcomes were cardiometabolic health, mental health and body composition. Intervention types were categorised as SB only, physical activity (PA) only, PA and SB or lifestyle interventions (PA/SB and diet).

Results Of 8087 records, 51 studies met the inclusion criteria. Meta-analysis of 34/51 studies showed a reduction of 22 min/day in sedentary time in favour of the intervention group (95% CI −35 to −9 min/day, n=5868). Lifestyle interventions reduced SB by 24 min/day (95% CI −41 to −8 min/day, n=3981, moderate quality) and interventions focusing on SB only by 42 min/day (95% CI −79 to −5 min/day, n=62, low quality). There was no evidence of an effect of PA and combined PA/SB interventions on reducing sedentary time.

Conclusions There was evidence that it is possible to intervene to reduce SB in adults. Lifestyle and SB only interventions may be promising approaches. More high quality research is needed to determine if SB interventions are sufficient to produce clinically meaningful and sustainable reductions in sedentary time.

INTRODUCTION

There is growing public health concern about the amount of time spent in sedentary behaviours (SB). SB are defined as behaviours where sitting or lying is the dominant posture and energy expenditure is very low.[1] Sedentary time accumulates daily while commuting, at work, at home and during leisure time.[2] Where studies have controlled for the influence of moderate-to-vigorous physical activity (MVPA), too much time spent in SB is associated with poor health, including elevated cardiometabolic risk markers, type 2 diabetes and premature mortality.[3–9] Where studies have controlled for the influence of total sedentary and moderate-to-vigorous activity time, increased breaks in sedentary time have been shown to be beneficially associated with waist circumference, body mass index (BMI), triglycerides and 2 h plasma glucose.[10] Interventions interrupting extended sitting with frequent short activity breaks have enhanced markers of cardio metabolic health.[11–13]

Recent systematic reviews have summarised the literature in respect to health implications,[14–18] measurement,[19] prevalence,[20] correlates[21] and interventions in young people.[22] To date, only one review of the evidence on interventions to influence total SB in adults has been published.[23] The review concluded that interventions with a specific goal of increasing PA levels and those which combined an increase in PA levels with a decrease in sedentary time resulted in modest reductions in SB, while interventions focusing on SB only resulted in greater reduction of sedentary time. The present systematic review expands this existing evidence[23] in five ways: (1) evaluating intervention effects using more precise categories of interventions; (2) assessing effects on pattern of SB accumulation; (3) conducting subgroup analyses; (4) including only randomised controlled trials (RCTs); and (5) assessing effects on health outcomes.

The primary aim of this review was to evaluate the effect of interventions which included an SB outcome measure in adults. The secondary aim was to determine the effects of interventions, which included an SB outcome, on measures of health.

METHODS

The protocol for this review is available online at the International Prospective Register for Systematic Reviews.[24]

Study selection criteria

Studies were eligible for inclusion if they met the following criteria:

Study design: RCTs

Population: Adults aged 18 years or more who have left school.

Intervention: Any intervention which included an SB outcome measure in free-living adults was eligible; those in clinical settings such as hospitals were excluded. Eligible control conditions were no intervention, waiting list, attention control (eg, general health information), usual care (eg, diabetes treatment involving lifestyle counselling) and alternative treatment conditions (eg, a structured exercise programme).

Outcomes: Studies reporting any of the following outcomes were included:
► Objectively measured SB obtained from accelerometers

- Objectively measured sitting time obtained from inclinometers
- Objectively or self-reported patterns of accumulation of SB
- Self-reported total sitting time
- Self-reported proxy measures of sitting time where it is not certain that people are sitting (eg, screen time and transport time) and proxy measures of overall SB (eg, occupational sitting time)

Other inclusion criteria: Only full text articles published in the English language were included in this review.

Data sources and searches

In January 2014, the Cochrane Central Register of Controlled Trials (Issue 12 of 12 December 2013), MEDLINE (1946-November week 3 2013), EMBASE (1980-week 1 2014), PsycINFO (1806-November week 5 2013), SPORTDiscus (1975-7 January 2014), CINAHL (1937-7 January 2013), Cochrane Database of Systematic Reviews (Issue 1 of 12 January 2014), Database of Health Promotion Research (Biblomap, Issue 4 of 4, October 2013), Database on Obesity and SB Studies (16 January 2014), Conference Proceedings Citation Indexes (Web of Science, 1900 to current), controlled-trials.com (16 January 2014), WHO International Clinical Trial Registry (16 January 2014) and the Networked Digital Library of Theses and Dissertations (1900-current) were searched. The search strategy for MEDLINE is listed in online supplementary 1. Reference lists and citations of relevant studies were examined and experts in the field contacted for details of ongoing and unpublished studies.

Study selection

At least two reviewers independently screened the titles/abstracts (AM, RJ) and full text articles (AM and RJ, CF or DHS). Eligibility disagreements were resolved by a third reviewer (NM).

Data extraction and quality assessment

Duplicate data extraction was performed independently for 10% of the included studies (AM and RJ, CF or DHS) and discrepancies resolved through discussion. The following secondary outcomes for this review were recorded from included studies:

- Biomarkers of cardiometabolic risk including blood glucose levels, blood lipid levels, total cholesterol levels, glycosylated haemoglobin, blood pressure
- Mental health outcomes including depression and anxiety
- Objectively obtained BMI, waist circumference and/or fat mass.

The full list of extracted data items can be obtained from the study protocol.[24]

Quality of all studies was assessed by two reviewers (AM, DHS) using the Tool for Assessing Risk of Bias from the Cochrane Collaboration.[25] Risk of bias was scored as 'high', 'unclear' or 'low' for the following domains: (1) participant selection bias, (2) intervention performance bias, (3) effect detection bias, (4) outcome reporting bias, (5) attrition bias and (6) bias due to comparability of baseline groups.

Publication bias was examined using a funnel plot whenever meta-analyses included 10 or more studies.[25]

Quality of evidence for primary outcomes was assessed using the GRADEpro software developed by the Grading of Recommendations Assessment Development and Evaluation (GRADE) Working Group.[26] An overall quality score is based on the assessment of risk of bias, indirectness, imprecision, inconsistency and publication bias of primary outcomes. The GRADE Working Group grades of evidence are high, moderate, low and very low quality.

Data synthesis and analysis

Studies reporting similar outcome measures were combined in meta-analyses using random effects models to account for intervention heterogeneity. Where suitable data were not reported, efforts were made to obtain the data from study authors. To account for variability between studies, inverse variance was used, giving more weight for studies with less variability. Effect sizes were estimated as mean differences (min/day) between the intervention and control groups. Review Manager 5.2 was used for quantitative analysis.[27]

For cluster RCTs where control of clustering was missing, intervention effects were approximately corrected by reducing the sample size of each trial to its 'effective sample size'. The sample size was divided by the design effect, which is $[1+(M-1)\times ICC]$, where M is the average of cluster size and ICC is the intracluster correlation coefficient.[25] An ICC of 0.01 was used.

Where suitable data were available, studies were combined in a meta-analysis regardless of whether missing data were imputed by authors. Variation in the degree of missing data was considered as a potential source of heterogeneity of results. A sensitivity analysis to examine the effect of inclusion of complete cases on robustness of intervention effects was performed.

Further heterogeneity of findings was assessed by comparing similarity of included studies in terms of study design, participants, interventions, outcomes and study quality. The cause of heterogeneity was evaluated by conducting subgroup and sensitivity analyses. Statistical heterogeneity was assessed by calculating the I^2 statistic indicating the variability of the intervention effect due to heterogeneity. Variability of more than 50% may indicate moderate to substantial heterogeneity of intervention effects according to the Cochrane Handbook.[25]

Subgroup analyses within this review focused on:

- Intervention type (SB, PA/SB or lifestyle which, in addition to PA/SB, also included a dietary/nutrition component)
- Gender (men, women, men and women)
- Intervention duration (<3 months, 3–6 months, >6 months)
- Follow-up duration (<3 months, 3–6 months, 7–12 months, >12 months)
- Intervention setting (work place vs home/community)
- Outcome measurement tool (objective measurement tool, sitting time self-report, proxy measurement tool)
- Study aim (SB as a primary vs secondary study aim)

Sensitivity analyses were used to test the effect of including studies which were cluster designs, used usual care or alternative treatment control groups, or were at 'high risk' of performance and attrition bias.

Included studies lacking data suitable for meta-analysis are described narratively.

RESULTS
Results of the literature search

Figure 1 displays the PRISMA diagram of the literature search. Inclusion criteria were met by 57 records which comprised 51 studies. Thirty-six studies provided adequate data to be included in meta-analyses.

Characteristics of included studies

Study and participant characteristics are summarised in table 1 of the online supplementary material. Of the 51 included studies (18 480 participants), 44 were RCTs[28–70] and seven were cluster RCTs[71–77] conducted in Europe (n=25), the USA (n=18), Australia (n=7) and China (n=1). The majority of studies were carried out in a mixed gender population (n=35); 13 studies

Figure 1 PRISMA diagram of the literature search results.

Figure 1 PRISMA diagram of the literature search results.

[Flow diagram contents:]

8,072 records identified through database searching

15 additional records identified through other sources

2,191 duplicates removed

5,896 records screened

5,668 records excluded

213 full-text records assessed for eligibility

156 full-text records excluded:

No sedentary behavior outcome measure: n=110

Ongoing trials: n= 37

No valid sedentary behavior outcome measure: n=2

Clinical setting: n=1

Acute intervention: n=1

Intervention not independent from outcome: n=2

Cross-sectional assessment: n=2

Insufficient data for decision: n=1

51 studies (57 records) included in narrative synthesis

36 studies included in quantitative synthesis (meta-analysis)

targeted women only[29 42 50 51 56 57 60 61 67 69 71 76] and three studies targeted men only.[29 31 44] Most studies included participants aged between 18–60 years (n=44), while seven studies included participants older than 60 years of age.[33 35 37 38 48 62 72] Twenty-three studies were conducted in overweight or obese adults, five studies in participants with type 2 diabetes mellitus and three studies in participants with high levels of cardiovascular risk factors. Two studies were conducted in pregnant women.

Types of intervention and control conditions varied substantially between included studies (see online supplementary table S1). Three studies employed an intervention specifically to reduce SB,[40 44 63] 16 studies aimed at increasing PA levels,[30 35 36 39 41 46 48 49 55 58–60 64 66 72 78] nine studies combined both approaches of reducing SB and increasing PA levels,[32 43 53 62 65 68 70 76 77] one study assessed the effect of a dietary intervention on SB,[61] and 22 studies (20 reports) applied a multicomponent lifestyle intervention and observed effects on sedentary behaviour (among other outcomes).[29 33 34 37 38 42 45 47 50–52 54 56 57 67 69 71 73 74 75] Twenty studies offered an alternative intervention,[30 36 39–41 45 46 49 52–55 59 61–63 68 72 77] 10 studies the usual/routine care,[29 37 38 42 50 51 67 71 74 75] seven studies used a waiting list control,[29 34 48 64 69 76 78] five studies an attention control,[35 44 56 57 60] and control participants of seven studies received no intervention at all.[32 33 43 47 58 66 70 73]

Risk of bias of included studies

Figure 2 shows each risk of bias item presented as percentages across all included studies.

Selection bias

Correct randomisation was used in 65% of the studies (33/51), and therefore there was low risk of bias in these studies. However, for the remaining studies, insufficient details were reported and

thus assessed as 'unclear'. In nearly 70% (35/51) of the studies, there was lack of reporting on whether or not participants knew in advance their group allocation, and thus there was an unclear risk of bias. For studies that provided information, studies were judged to be at low risk of allocation concealment bias.

Performance bias

It is recognised that in lifestyle interventions it is not possible to blind participants and researchers delivering the intervention to group allocation and this creates high risk of bias. However, 67% (34/51) of included studies were considered at low risk of performance bias because SB was not the primary outcome. A further 31% (16/51) of included studies were judged to be at high risk of performance bias because the participants and researchers delivering the intervention were not blinded to the purpose of the intervention, which was reducing SB. Risk of performance bias was unclear for one study[33] due to insufficient information provided.

Detection bias

Sixty-one per cent of the studies (31/51) assessed SB through self-reports and thus were at high risk for detection bias. The risk of cross-contamination was 'low' in half of the studies and 'unclear' in the other half.

Attrition bias

The issue of incomplete outcome data was sufficiently addressed in 47% (24/51) of the studies, and thus these studies were at low risk of attrition bias. However, 43% (22/51) of the studies did not account for missing data and thus were at high risk of attrition bias. Five studies were at 'unclear' risk of attrition bias.

Table 1 Intervention effects for change of sedentary behaviour by subgroups

| Subgroup	Studies	Participants	Intervention effect (min/day), MD (95% CI, I^2)
Sex*			
Men	2	434	−57.94 (−86.14 to −29.74; 0%)
Women	10	1541	−5.97 (−23.51 to 11.57; 33%)
Men/women	22	3893	−25.32 (−42.94 to −7.69; 83%)
Intervention duration†			
≤3 months	14	1474	−47.51 (−76.57 to −18.46; 81%)
3–6 months	11	2119	−15.20 (−33.08 to 2.68; 67%)
>6 months	9	2275	0.30 (−17.83 to 18.44; 61%)
Follow-up duration‡			
<3 months	17	1954	−42.17 (−67.31 to −17.02; 84%)
3–6 months	13	2489	−22.29 (−41.61 to −2.96; 77%)
7–12 months	11	2327	−26.60 (−45.95 to −7.24; 73%)
>12 months	5	1264	−3.06 (−34.05 to 27.94; 83%)
Intervention setting‡			
Workplace	8	1790	−8.93 (−26.64 to 8.78; 66%)
Other	26	4078	−28.21 (−46.34 to −10.09; 80%)
Assessment tool‡			
*activ*PAL	2	67	−45.37 (−87.99 to −2.74; 76%)
Actigraph	4	334	−27.93 (−70.71 to 14.85; 75%)
Sitting time questionnaire	12	2576	−10.92 (−30.59 to 8.74; 57%)
Proxy measure questionnaire	17	2983	−29.39 (−50.56 to −8.21; 84%)
Intervention aim‡			
SB Primary outcome	14	2258	−24.05 (−45.43 to −2.67; 73%)
SB Secondary outcome	22	3764	−23.17 (−40.02 to −6.32; 80%)

*statistically significant subgroup difference at p<0.01.
†statistically significant subgroup difference at p <0.05.
‡non-significant subgroup difference.
SB, sedentary behaviour.

Comparability of baseline groups
Over 50% (29/51) of the studies were at low risk of bias. Apparent flaws in the randomisation process were found in three studies[53 76 78] and therefore assessed at high risk of bias related to the comparability of baseline groups. For the remaining studies, no formal assessment of the comparability of baseline groups was reported, and thus the risk of bias was 'unclear'.
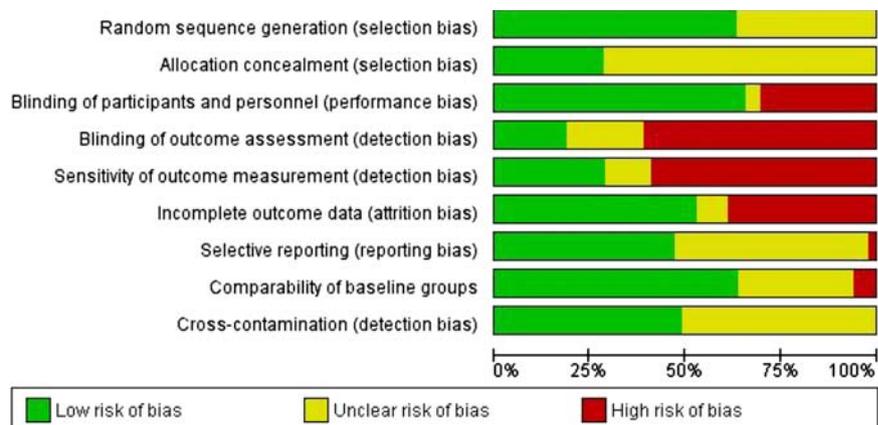
Reporting bias
For half of the studies (26/51), access to a published study protocol or trial register was missing so that the risk of selective reporting was 'unclear'. However, nearly 50% (24/51) of the studies were at low risk of selective outcome reporting. One study did not report all outcomes as stated in the study protocol and thus was at high risk of selective reporting.[70]

Publication bias
Lifestyle interventions were the only category of interventions where at least 10 studies were available and thus suitable for assessment of publication bias using the funnel plot (see online supplementary figure S1). The asymmetric distribution of effect sizes might indicate a publication bias towards studies with beneficial effects for reducing SB. However, an asymmetric funnel plot might be a study size effect.

Effect of interventions
Primary outcomes
The primary outcomes reported were overall time spent in SB as minutes per day (n=49) or percentage of assessed time period (n=3), number of sitting breaks (n=3) and number of prolonged sitting events (n=3).

Online supplementary table S1 summarises the original trial authors' conclusions of study outcomes. Twenty studies indicated a beneficial effect of interventions for reducing SB in favour of the intervention group. Of these, 10 studies employed a lifestyle intervention,[29 33 34 37 38 42 51 52 54 74] six studies targeted increase in PA,[30 41 46 48 64 78] two studies were combined PA/SB interventions[32 68] and two studies were SB interventions.[40 63] Two studies reported a beneficial intervention effect in favour of the control group;[39 60] both studies were PA interventions. Control conditions were attention control[60] and an alternative exercise treatment.[39] Twenty-four studies suggested no evidence of a group difference in SB: 10 lifestyle interventions,[29 45 50 52 56 57 67 71 73 75] seven PA interventions,[35 36 49 55 58 66 72] six PA/SB interventions,[53 62 65 70 76 77] and one SB intervention.[44] Four studies—two lifestyle,[47 69] one PA/SBs,[43] one dietary intervention[61]—did not conclude on SB outcomes despite assessing SB.

A meta-analysis of 34 studies (5868 participants) suggested an overall reduction in sedentary time by mean differences (MD) of −22.34 min/day (95% CI −35.81 to −8.88, p=0.001, I^2=71%) in favour of the intervention group. Figure 3 shows effect sizes of individual studies and pooled results by intervention type. Findings indicated a beneficial effect of interventions specifically targeting the reduction in SB as well as interventions employing a
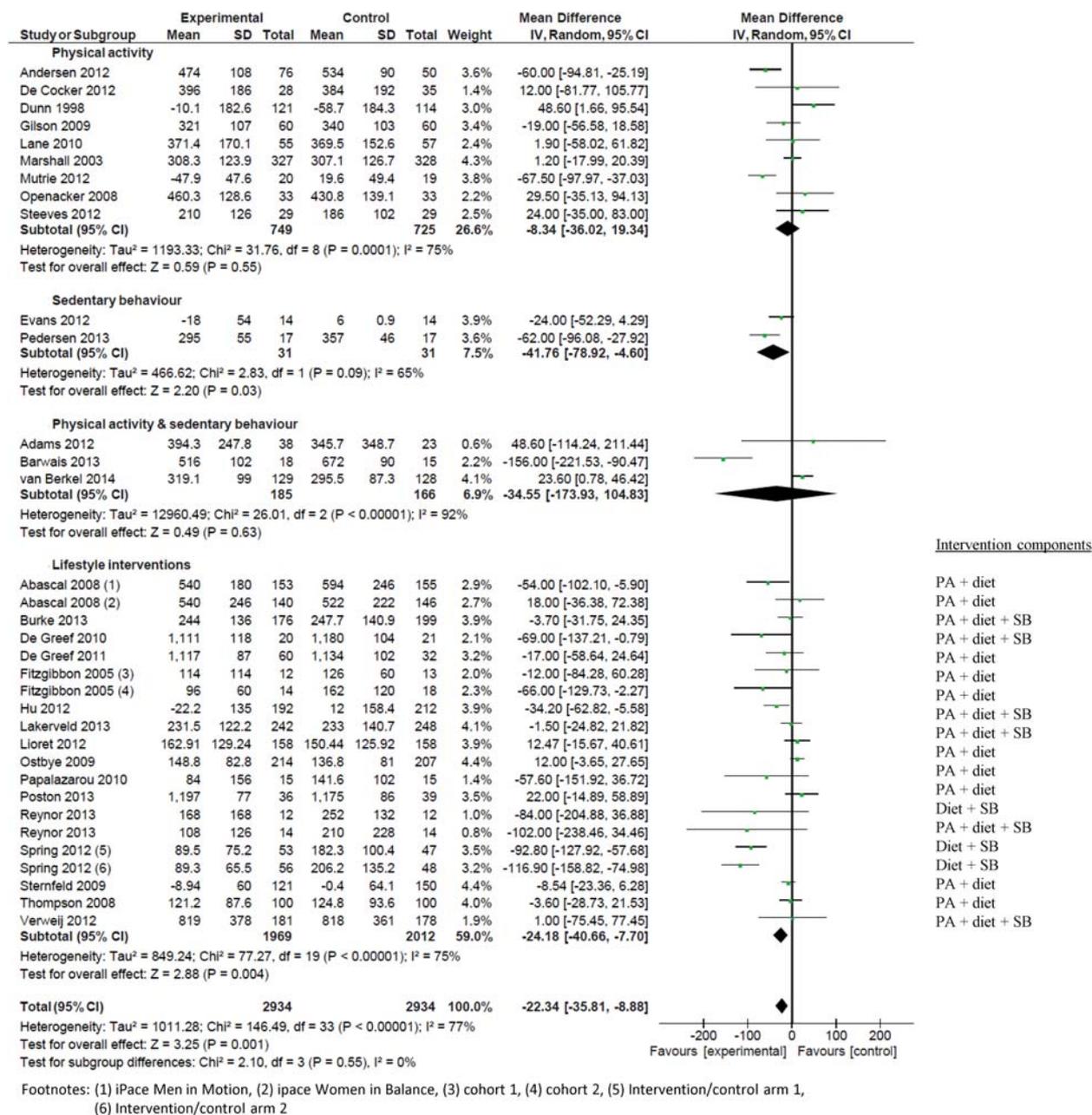
Figure 2 Risk of bias item presented as percentages across all studies.

Study or Subgroup	Experimental Mean	SD	Total	Control Mean	SD	Total	Weight	Mean Difference IV, Random, 95% CI	Intervention components
Physical activity									
Andersen 2012	474	108	76	534	90	50	3.6%	-60.00 [-94.81, -25.19]	
De Cocker 2012	396	186	28	384	192	35	1.4%	12.00 [-81.77, 105.77]	
Dunn 1998	-10.1	182.6	121	-58.7	184.3	114	3.0%	48.60 [1.66, 95.54]	
Gilson 2009	321	107	60	340	103	60	3.4%	-19.00 [-56.58, 18.58]	
Lane 2010	371.4	170.1	55	369.5	152.6	57	2.4%	1.90 [-58.02, 61.82]	
Marshall 2003	308.3	123.9	327	307.1	126.7	328	4.3%	1.20 [-17.99, 20.39]	
Mutrie 2012	-47.9	47.6	20	19.6	49.4	19	3.8%	-67.50 [-97.97, -37.03]	
Openacker 2008	460.3	128.6	33	430.8	139.1	33	2.2%	29.50 [-35.13, 94.13]	
Steeves 2012	210	126	29	186	102	29	2.5%	24.00 [-35.00, 83.00]	
Subtotal (95% CI)			749			725	26.6%	-8.34 [-36.02, 19.34]	
Heterogeneity: Tau² = 1193.33; Chi² = 31.76, df = 8 (P = 0.0001); I² = 75%									
Test for overall effect: Z = 0.59 (P = 0.55)									
Sedentary behaviour									
Evans 2012	-18	54	14	6	0.9	14	3.9%	-24.00 [-52.29, 4.29]	
Pedersen 2013	295	55	17	357	46	17	3.6%	-62.00 [-96.08, -27.92]	
Subtotal (95% CI)			31			31	7.5%	-41.76 [-78.92, -4.60]	
Heterogeneity: Tau² = 466.62; Chi² = 2.83, df = 1 (P = 0.09); I² = 65%									
Test for overall effect: Z = 2.20 (P = 0.03)									
Physical activity & sedentary behaviour									
Adams 2012	394.3	247.8	38	345.7	348.7	23	0.6%	48.60 [-114.24, 211.44]	
Barwais 2013	516	102	18	672	90	15	2.2%	-156.00 [-221.53, -90.47]	
van Berkel 2014	319.1	99	129	295.5	87.3	128	4.1%	23.60 [0.78, 46.42]	
Subtotal (95% CI)			185			166	6.9%	-34.55 [-173.93, 104.83]	
Heterogeneity: Tau² = 12960.49; Chi² = 26.01, df = 2 (P < 0.00001); I² = 92%									
Test for overall effect: Z = 0.49 (P = 0.63)									
Lifestyle interventions									
Abascal 2008 (1)	540	180	153	594	246	155	2.9%	-54.00 [-102.10, -5.90]	PA + diet
Abascal 2008 (2)	540	246	140	522	222	146	2.7%	18.00 [-36.38, 72.38]	PA + diet
Burke 2013	244	136	176	247.7	140.9	199	3.9%	-3.70 [-31.75, 24.35]	PA + diet + SB
De Greef 2010	1,111	118	20	1,180	104	21	2.1%	-69.00 [-137.21, -0.79]	PA + diet + SB
De Greef 2011	1,117	87	60	1,134	102	32	3.2%	-17.00 [-58.64, 24.64]	PA + diet
Fitzgibbon 2005 (3)	114	114	12	126	60	13	2.0%	-12.00 [-84.28, 60.28]	PA + diet
Fitzgibbon 2005 (4)	96	60	14	162	120	18	2.3%	-66.00 [-129.73, -2.27]	PA + diet
Hu 2012	-22.2	135	192	12	158.4	212	3.9%	-34.20 [-62.82, -5.58]	PA + diet + SB
Lakerveld 2013	231.5	122.2	242	233	140.7	248	4.1%	-1.50 [-24.82, 21.82]	PA + diet + SB
Lioret 2012	162.91	129.24	158	150.44	125.92	158	3.9%	12.47 [-15.67, 40.61]	PA + diet
Ostbye 2009	148.8	82.8	214	136.8	81	207	4.4%	12.00 [-3.65, 27.65]	PA + diet
Papalazarou 2010	84	156	15	141.6	102	15	1.4%	-57.60 [-151.92, 36.72]	PA + diet
Poston 2013	1,197	77	36	1,175	86	39	3.5%	22.00 [-14.89, 58.89]	Diet + SB
Reynor 2013	168	168	12	252	132	12	1.0%	-84.00 [-204.88, 36.88]	PA + diet + SB
Reynor 2013	108	168	14	210	228	11	0.8%	-102.00 [-238.46, 34.46]	Diet + SB
Spring 2012 (5)	89.5	75.2	53	182.3	100.4	47	3.5%	-92.80 [-127.92, -57.68]	Diet + SB
Spring 2012 (6)	89.3	65.5	56	206.2	135.2	48	3.2%	-116.90 [-158.82, -74.98]	PA + diet
Sternfeld 2009	-8.94	60	121	-0.4	64.1	150	4.4%	-8.54 [-23.36, 6.28]	PA + diet
Thompson 2008	121.2	87.6	100	124.8	93.6	100	4.0%	-3.60 [-28.73, 21.53]	PA + diet + SB
Verweij 2012	819	378	181	818	361	178	1.9%	1.00 [-75.45, 77.45]	
Subtotal (95% CI)			1969			2012	59.0%	-24.18 [-40.66, -7.70]	
Heterogeneity: Tau² = 849.24; Chi² = 77.27, df = 19 (P < 0.00001); I² = 75%									
Test for overall effect: Z = 2.88 (P = 0.004)									
Total (95% CI)			2934			2934	100.0%	-22.34 [-35.81, -8.88]	
Heterogeneity: Tau² = 1011.28; Chi² = 146.49, df = 33 (P < 0.00001); I² = 77%									
Test for overall effect: Z = 3.25 (P = 0.001)									
Test for subgroup differences: Chi² = 2.10, df = 3 (P = 0.55), I² = 0%									

Favours [experimental]   Favours [control]
(axis: -200 -100   0   100   200)

Footnotes: (1) iPace Men in Motion, (2) ipace Women in Balance, (3) cohort 1, (4) cohort 2, (5) Intervention/control arm 1, (6) Intervention/control arm 2

Figure 3 Forest plot of the intervention effect for reducing sitting time in minutes/day in adults by type of intervention. PA, physical activity; SB, sedentary behavior.

lifestyle intervention approach on reduced SB. Specific SB interventions (n=2, 62 participants) yielded an MD of −41.76 min/day (95% CI −78.92 to −4.60, p=0.003, I²=65%) and lifestyle interventions (n=20, 3881 participants) an MD of −24.18 min/day (95% CI −40.66 to −7.70, p=0.004, I²=75%). There was no evidence of a statistically significant effect of PA interventions or combined PA/SB interventions for reducing SB.

Pooled intervention effects on SB patterns indicated no statistically significant effect for both the number of sitting breaks per hour or the number of prolonged sitting events of more than 30 min.

As indicated by the large I² statistic, the level of statistical heterogeneity between studies was high. Subgroup analyses were conducted (defined a priori) to assess potential reasons for heterogeneity (table 1). A significant subgroup difference between assessed groups was detected for gender and intervention

duration. Studies in men-only (n=2; 434 men), but not women-only (n=10; 1541 women), resulted in significant intervention effects for reduced SB of intervention group participants (MD −57.94 min/day, 95% CI −86.14 to −29.74 min/day, p<0.001). The combined effects of mixed gender studies (n=22; 3393 participants) also showed benefit in favour of the intervention group (MD −25.32 min/day, 95% CI −42.94 to −7.69 min/day, p=0.005). Interventions of up to 3 months resulted in a significant reduction in sedentary time by an MD of −47.51 min/day (95% CI −76.57 to −18.46 min/day, p=0.001, 14 studies, 1474 participants) in favour of the intervention group, whereas longer intervention durations of more than 3 months did not show beneficial intervention effects (table 1). Heterogeneity between studies could not be explained by follow-up duration, intervention setting, type of assessment tool and whether reducing SB was a primary or secondary aim

of the study. However, subgroup analysis revealed that long-term effects of interventions were evident up to 12 months. The beneficial intervention effects attenuated at a follow-up duration of more than 12 months. All intervention settings except workplaces resulted in a significant reduction in SB in favour of the intervention group. Objective assessment of SB using an inclinometer and subjective assessment using proxy measure questionnaires resulted in a detection of a beneficial intervention effect. The overall intervention effect was not influenced by whether SB was a primary or secondary outcome (table 1).

Sensitivity analyses (see online supplementary tables S2–S5) show that results on SB for different types of interventions were not affected by inclusion of cluster RCTs, studies at high risk of attrition and performance bias, and studies with usual care or alternative treatment as the control group.

Secondary outcomes

Studies reported intervention effects on fasting blood glucose concentration,[31 42 56] glycosylated haemoglobin levels,[37 42 69] triglyceride levels,[31 42 56 69] low-density lipoprotein levels,[31 42 56 69] total cholesterol,[37 42 56 69] high-density lipoprotein levels,[31 39 42 56 64 69] blood pressure,[32 38 43 57 59 65 70] BMI,[29 33 36 37 42 55 56 57 58 59 62 64 69 74] waist circumference,[31 42 55–59 62 64 69 74 76] percentage body fat[42 55 56 58 62 64] and mental health outcomes.[29 41 48 49 64 72] Some studies indicated a reduction in these secondary outcomes; however, studies were PA-only or lifestyle interventions and none of the studies were SB-only studies. Therefore, it is not possible to determine the intervention effect of reduced SB on cardiometabolic risk, body composition and mental health outcome. Specific SB studies did not assess the intervention effect on health outcomes. Meta-analysis results for each outcome are not reported here but are available from the authors.

Quality of evidence

Table 2 summarises the quality of evidence for reducing sedentary time by intervention type and duration. Owing to the intention of comparing different types of intervention with various control conditions, which was considered in the sensitivity analyses, the quality of evidence was not downgraded for indirectness or heterogeneity. Many plausible reasons for heterogeneity exist (eg, variation in population age, ethnicity, socioeconomic status).

Lifestyle interventions

The overall quality of evidence for lifestyle interventions was moderate with downgrading of the evidence by one level due to limitations in the design and implementation of the included studies.

PA/SB interventions

The overall quality of evidence of combined PA and SB interventions for reducing SB was moderate. The quality was downgraded by one level for high risk of bias in the majority of included studies.

PA interventions

Overall, the quality of PA intervention was moderate with the majority of studies having a high risk of detection and attrition bias.

SB interventions

The quality of evidence for reducing SB in adults was low based on the two studies available. The quality was downgraded twice for imprecision of results and high risk of performance bias. Participants and personnel were not blinded to the intervention intention.

DISCUSSION

Summary of main findings

There was clear evidence that it is possible to intervene to reduce SB in adults by 22 min/day in favour of the intervention group. Moderate to high-quality evidence on the efficacy of lifestyle interventions for reducing SB suggests that this may be a promising approach. Interventions focusing on SB only resulted in the greatest reduction in sedentary time (42 min/day); however, the quality of evidence was low and restricted to two studies only. Findings suggested that intervention durations up to 3 months and interventions targeting men and mixed genders can produce significant reductions in SB. There was no evidence that PA and combined PA/SB interventions reduced SB. Evidence of intervention effects on changes in patterns of accumulation of SB was limited. Encouragingly, intervention effects were evident up to 12 months. Interventions in any setting except the workplace resulted in a significant reduction in SB in favour of the intervention group.

This systematic review sought to evaluate the evidence of effects of interventions which included SB as an outcome measure on cardiometabolic risk factors, body composition and mental health outcomes. Studies reporting these outcomes were PA or lifestyle interventions, and thus it was unclear whether any intervention effect was due to reduction in SB. Furthermore, the majority of studies that assessed health-related outcomes did not show a reduction in SB. However, improvement of health outcomes due to reduction of SB has been demonstrated in laboratory-based studies[12] and a recently published community-based RCT.[79]

Comparison of the findings with the literature

Prince et al[23] published a systematic review on the effects of interventions for reducing SB in adults. Our findings are consistent with those of Prince et al in relation to the effect of PA/SB interventions and interventions focusing on SB only, despite there being no overlap of included studies in the latter. The SB studies on which Prince et al based their main conclusion were excluded from this review because they either did not report a valid SB outcome measure[80] or the intervention was not independent of the outcome (measuring TV viewing time while blocking TV function).[81] In contrast to Prince et al, we found no evidence of a beneficial effect on SB from interventions focused on increasing PA. This difference in findings may be explained by six studies in our review being classed as lifestyle interventions while Prince et al classed them as PA interventions and one study being classed as a PA/SB intervention while Prince et al classed it as a PA intervention. Authors of future reviews should use precise categories of intervention types to identify the potential of single or multicomponent interventions (eg, lifestyle intervention which, in addition to PA/SB, also included a dietary/nutrition component) to reduce SB.

Other systematic reviews have been conducted with a focus on the effect of workplace interventions for reducing sitting time.[82–84] Some findings are consistent[82] with the findings of this study on the effect of workplace interventions to reduce SB while others were not.[83 84] Inconsistency can be explained by differences in

Table 2 GRADE assessment of quality of evidence

Interventions for reducing sedentary behaviour

Outcomes	Illustrative comparative risks* (95% CI) Corresponding risk Interventions for reducing sedentary behaviour	Number of Participants (studies)	Quality of the evidence (GRADE)
Effect of lifestyle interventions	The mean effect of lifestyle interventions in the intervention groups was 24.18 min/day lower (40.66 to 7.70 lower)	3981 (20 studies)	⊕⊕⊕⊖ moderate†
Intervention duration ≤3 months	The mean effect of lifestyle interventions—intervention duration ≤3 months in the intervention groups was 97.75 min/day lower (121.88 to 73.61 lower)	297 (5 studies)	⊕⊕⊕⊕ high
Intervention duration 3–6 months	The mean effect of lifestyle interventions—intervention duration 3–6 months in the intervention groups was 8.42 min/day lower (19.05 lower to 2.21 higher)	1664 (7 studies)	⊕⊕⊕⊖ moderate‡
Intervention duration >6 months	The mean effect of lifestyle interventions—intervention duration >6 months in the intervention groups was 3.99 min/day lower (21.93 lower to 13.96 higher)	2040 (8 studies)	⊕⊕⊕⊖ moderate†
Effect of physical activity/ sedentary behaviour interventions	The mean effect of physical activity/sedentary behaviour interventions in the intervention groups was 32.51 min/day lower (106.52 lower to 41.50 higher)	471 (4 studies)	⊕⊕⊕⊖ moderate†
Intervention duration ≤3 months	The mean effect of physical activity/sedentary behaviour interventions—intervention duration ≤3 months in the intervention groups was 54.69 min/day lower (166.60 lower to 57.22 higher)	214 (3 studies)	⊕⊖⊖⊖ very low§¶
Intervention duration 3–6 months	The mean effect of physical activity/sedentary behaviour interventions—intervention duration 3–6 months in the intervention groups was 23.60 min/day higher (0.78 higher to 46.42 higher)	257 (1 study)	⊕⊕⊕⊖ moderate**
Intervention duration >6 months	No evidence available	0 (0)	No evidence available
Effect of physical activity interventions	The mean effect of physical activity interventions in the intervention groups was 6.08 min/day lower (38.00 lower to 25.84 higher)	1354 (8 studies)	⊕⊕⊕⊖ moderate††
Intervention duration ≤3 months	The mean effect of physical activity interventions—intervention duration ≤3 months in the intervention groups was 10.43 min/day lower (49.85 lower to 28.98 higher)	935 (5 studies)	⊕⊕⊕⊖ moderate††
Intervention duration 3–6 months	The mean effect of physical activity interventions—intervention duration 3–6 months in the intervention groups was 21.52 min/day lower (103.55 lower to 60.51 higher)	184 (2 studies)	⊕⊕⊕⊖ moderate††
Intervention duration >6 months	The mean effect of physical activity interventions—intervention duration >6 months in the intervention groups was 48.60 min/day higher (1.66 to 95.54 higher)	235 (1 study)	⊕⊕⊕⊖ moderate‡‡
Effect of sedentary behaviour interventions	The mean effect of sedentary behaviour interventions in the intervention groups was 41.76 min/day lower (78.92 to 4.60 lower)	62 (2 studies)	⊕⊕⊖⊖ low§·§§
Intervention duration ≤3 months	The mean effect of sedentary behaviour interventions—intervention duration ≤3 months in the intervention groups was 41.76 min/day lower (78.92 to 4.60 lower)	62 (2 studies)	⊕⊕⊖⊖ low§·§§
Intervention duration 3–6 months	No evidence available	0 (0)	No evidence available
Intervention duration >6 months	No evidence available	0 (0)	No evidence available

GRADE Working Group grades of evidence.
High quality: Further research is very unlikely to change our confidence in the estimate of effect.
Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.
Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.
Very low quality: We are very uncertain about the estimate.
*The basis for the assumed risk (eg, the median control group risk across studies) is provided in footnotes. The corresponding risk (and its 95% CI) is based on the assumed risk in the control group and the relative effect of the intervention (and its 95% CI).
†The majority of studies were of high risk of selection, performance or detection bias.
‡Half of the studies were of high risk for performance bias (no blinding of participants or personnel to the intervention intention).
§The wide CI indicates imprecision of results.
¶All studies were of high risk of performance bias and more than half showed high risk of attrition.
**The study was of high risk of selection bias.
††Studies were of high risk of detection or attrition bias.
‡‡The study was of high risk of detection bias.
§§The studies were of high risk of performance bias, that is, participants and personnel were not blinded.

inclusion criteria, since the majority of studies included in these reviews were not RCTs and thus did not qualify for our review. However, further high-quality RCTs investigating the effect of workplace interventions on sitting time are currently being conducted and publication of new evidence will follow shortly.[85]

Implications for research and practice

Findings from lifestyle interventions and studies focusing on reducing SB are promising. While this is encouraging, SB are health-related behaviours and part of a pathway to better health outcomes. More high-quality research is needed that includes clinical health outcome measures. However, the findings of this review should encourage clinicians and public health practitioners to provide advice on how to reduce total volume of sitting time and breaking up long periods of sitting. This advice should not diminish or replace advice on achieving the recommended levels of MVPA. It is somewhat surprising that interventions that targeted PA alone, or even PA and SB, appeared to be

less effective in reducing SB. This suggests that attention needs to be paid to the ways in which SB are targeted in these interventions. For example, it may be important to improve knowledge about the independent health risks of SB and to highlight the risk of compensatory behaviour (eg, a feeling that you have earned the right to be sedentary because you went for a brisk walk earlier). Given the evidence that increased breaks in SB are associated with improved health status, consensus is needed on the most appropriate SB patterning descriptors to use which are sensitive to intervention (eg, 'breaking rate' or time spent/number of longer sedentary events). New interventions should also be developed around technologies that allow people to monitor their SB in addition to their physical activity to support them in setting goals to reduce their SB and increase PA.

The majority of studies included in the meta-analyses assessed intervention effects using self-report. While self-report measures are pragmatic and may provide contextual information, they have limitations in terms of accuracy. Subgroup analysis revealed that objective assessment of SB using a posture measurement tool such as the *activ*PAL and subjective assessment using proxy measure questionnaires (captures context specific sitting time) resulted in the detection of a beneficial intervention effect. Assessment tools that measure posture might be more valid and reliable in measuring SB and thus detecting intervention effects compared to *estimation of SB* via *accelerometry* (eg, ActiGraph). Therefore, researchers and practitioners should use posture measurement tools and context specific measurement tools which may prompt a reliable cognitive recall of sedentary behaviour.

Heterogeneity between studies was only partly explained by differences of studies in gender and intervention duration. Further work is warranted to identify the 'active ingredients' of the successful interventions and to explore the specific behaviour change techniques employed as well as barriers and facilitators of SB interventions. General principles for development of interventions to reduce SB have been established drawing from behavioural research on physical activity.[86] Examples include evaluating interventions designed for very specific contexts (work environments at home) and using behaviour change theory and associated techniques[87] to systematically understand and change SB in different groups and settings.

Additionally, future studies should consider the influence of gender, given that some cohort studies suggested deleterious relationships of SB with health outcomes to be more pronounced in women than men. However, based on our review evidence, interventions with the potential to reduce SB showed limited effects when targeting women. Limited evidence was available on intervention effects on sedentary time in older adults.

Strengths and limitations

The systematic and transparent methods reported here reduce identification and selection bias. The inclusion criteria used for study designs (only RCTs) meant that the risk of bias was reduced. Overall, the robust methods used in this review ensure that the results and conclusions are likely to be as truly valid and replicable as possible. Subgroup and sensitivity analyses enabled a more nuanced understanding and interpretation of the results, as well as exploring the effect of potentially influential variables. Lastly, our exploration of the clinical outcomes was a strength, and led to the identification of research gaps which should be addressed in future RCTs.

One limitation was that no subgroup analysis for age was undertaken because there were too few studies in older adults.

CONCLUSION

There was evidence that it is possible to intervene to reduce SB in adults by around 22 min/day. Lifestyle interventions and those targeting SB only may be promising approaches, but more high-quality research is needed. More research is also needed to determine if SB interventions are sufficient to produce clinically meaningful and sustainable reductions in sedentary time. Further work is needed to identify the 'active' intervention components.

What are the new findings?

- ▶ Interventions targeting sedentary behaviour (SB) and lifestyle interventions can reduce sedentary time in adults.
- ▶ Interventions targeting an increase in physical activity and interventions combining an increase of physical activity with reducing sedentary behaviour did not reduce sedentary time in adults.
- ▶ We do not yet know if effective interventions for reducing sedentary behaviour result in clinically meaningful and sustained improvements in health outcomes.

How might it impact on clinical and public health practice in the near future?

- ▶ The findings of this study (together with the broader body of relevant evidence) do not point to specific recommendations on the degree of reduction in sitting time required to deliver significant health benefits. Nevertheless, the findings should encourage clinicians and public health practitioners to provide advice about reducing the total volume of sitting time and breaking up long periods of sitting by demonstrating that such advice can be effective. This advice should not diminish or replace advice on achieving recommended levels of physical activity.
- ▶ Interventions with a focus on physical activity should provide additional emphasis on the importance of and barriers to reducing SB. New technologies should be developed to allow self-monitoring and goal setting around SB as well as physical activity.
- ▶ Awareness will be raised on the topic of sedentary behaviour and its impact on health.
- ▶ Interventions that target sedentary behaviour will be developed and tested.
- ▶ Further research is needed to determine the clinical significance of changing patterns of sedentary behaviour.

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