Original Article

Breaking sedentary behaviour has the potential to increase/ maintain function in frail older adults

Juliet A. Harvey, Sebastien F.M. Chastin, Dawn A. Skelton

Institute of Applied Health Research, School of Health and Life Sciences, Glasgow Caledonian University, Glasgow, Scotland, UK

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Abstract

Objectives: This pilot study aimed to evaluate the effect of a sedentary behaviour (SB) reduction intervention (Stomp Out (Prolonged) Sitting (SOS)) in frail older adults. Methods: Participants (>65years) were recruited from sheltered housing complexes and randomized into 2 groups. On weeks 2, 6 and 10 both groups had face-to-face 40min motivational sessions, including feedback on physical function and SB. One group had the addition of real-time tactile feedback on sitting. Total sedentary time and patterns of SB were recorded by activPAL, along with validated measures of function: Timed Up and Go (TUG), Sit-to-Stand (STS) and balance tests. Outcomes were analyzed by intention-to-treat mixed model analysis. Results: Twenty-three participants started the SOS intervention. Health issues led to high attrition in this frail population. TUG (4 seconds faster) and STS (>2 rises more in 30 seconds) scores improved significantly in both groups. There were no significant changes in SB parameters. Conclusion: Motivational interviewing alongside functional test feedback, visual and real-time feedback on SB improved physical function over the study. This pilot study suggests that sit-to-stand transitions to break prolonged sitting time may help reduce frailty and functional decline in people who are often unable to engage in more intense exercise interventions.

Keywords: Sedentary Behavior, Sitting Time, Physical Function, Frailty, Older Adults

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Introduction

The aim of health and social care policy in the UK is to support people to live healthier independent lives at home, which are, as far as possible, focused on prevention and early management^{1,2}. In epidemiological studies, high levels of sitting time have been linked with frailty, falls and poor physical function and those older adults (OA) who break their sitting time more frequently have better physical function³. Frailer OA may potentially have more to gain from reducing sedentary time than younger, or more able adults, if there is the potential for functional gains without the higher energy expenditure of more formal exercise sessions, which are often considered unachievable in this group4.

Sedentary behaviour (SB) is defined by both posture (sitting or reclining) and low energy expenditure (<1.5 Metabolic Equivalent of Task (METS)) during waking hours⁵. OA spend 65-80% of the waking day sedentary⁶ which has a deleterious effect on metabolism, health and physical function^{7,8}. Not all sitting is detrimental, therefore it is important to aim SB interventions at times in the day when SB is passive or non-purposeful and in the context of other tasks of the 24 hour day9.

Interventions that are primarily aimed at changing sitting time, rather than combined with increasing physical activity, are thought to be the most successful at reducing sitting time¹⁰. Reducing SB, along with having its own benefits, may be an "entry level" intervention to engage OA in moving more often and towards their optimal physical activity potential by embedding small bouts of functional training within their daily routine. To date, interventions designed for OA aimed at reducing SB have asked the participants to set goals 11-15

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Corresponding author: Dr Juliet Harvey, Institute of Applied Health Research, Glasgow Caledonian University, 70 Cowcaddens Road, Glasgow, Scotland, UK, G4 OBA

E-mail: Juliet.Harvey@gcu.ac.uk Edited by: Yannis Dionyssiotis Accepted 17 February 2018

or provided an information booklet with tips and tasks¹⁶ over a period of 12 weeks or less. SB has been reported using objective measures (activPAL (AP)¹²⁻¹⁴, ActiGraph¹¹ or Sense WearPro¹⁵) and self-report¹⁵⁻¹⁶. Results are mixed and only two studies observed functional outcomes.

Rosenberg et al. (2015) found a small positive effect on gait speed as a result of an 8 week programme to reduce SB in community dwelling older adults¹³. Barone Gibbs et al. (2017) found a small positive effect on Chair Sit-to-Stand Test (STS), but not on objectively measured SB¹⁵. Frail OA are underserved in SB intervention studies. Only Matei et al. (2015)¹⁶ included older adults living in sheltered housing and this group was unable to positively change their sitting time, recorded by self-report.

It is recognised that interventions to change behaviour are complex and there is no clear approach to achieving behaviour change in OA¹⁷. Behavioural feedback is useful as it allow the participant to reflect on their behaviour. The activPAL-VTaP (VT) (PAL Technologies: www.paltechnologies.com) is an AP activity monitor unit with the additional facility of being able to provide real-time tactile feedback to the user on their sitting time by way of a vibration prompt. It was not known whether real-time feedback could have an effect on reducing SB in older adults. The aim of this randomized pilot trial was to examine the use of both real-time and retrospective SB feedback alongside functional feedback during motivational sessions and determine its effect on objectively measured SB and physical function in frailer older adults living in sheltered housing.

Materials and methods

Study design

The study was a two-arm randomized pilot study. Participants were recruited from Sheltered Housing Complexes in Glasgow (8 sites managed by 3 Sheltered Housing Associations and participants from Glasgow Caledonian University (GCU) Older Adults Database). Participants were randomized (random number table) before baseline assessments into either the AP or VT group. Data was collected between April 2013 and June 2014. The GCU School of Health and Life Sciences Ethical Committee gave approval (HLS12/59) and all participants gave written informed consent. Inclusion and exclusion criteria can be found in Table 1. The study is presented according to the Template for Intervention Description and Replication (TIDieR) guidelines (Appendix 1)18. A target of 25 participants was set as this was estimated to detect a 4% reduction in sitting time with (80%, alpha 0.05), to allow for attrition we aimed to recruit 35 participants.

Intervention: Motivational session and retrospective feedback (AP)

During the study all participants received three 40 min face-to-face sessions with retrospective feedback every 4

Inclusion	Exclusion
65+ years of age	Allergy to adhesive tape (for fixing the AP/VT to the thigh)
Medically Stable	Any acute or uninvestigated medical conditions.
Living in Sheltered Housing	
Independently Ambulatory (with or without a walking aid)	
Score <7 in the 6-CIT (Cognitive Impairment Test) [‡]	
Able to achieve 100 degrees of hip flexion in both hips from a sitting and lying position (required to turn off VT equipment).	
Have intact vibro-tactile sensation on the thigh (assessed by neurothesiometer).	

*Brooke P, Bullock R: Validation of a 6 item cognitive impairment test with a view to primary care usage. Int J Geriatr Psychiatry 1999: 14 (11): 936-40.

Table 1. Inclusion and exclusion criteria.

weeks. The intervention was based on: the Transtheoretical Model¹⁹; previous intervention studies^{11,20}; discussion with Sheltered Housing Support Officers; lifelogging contextual work and qualitative interviews with older adults^{9,21}. A support pack was produced to ensure each participant received the same basic information and could refer to the pack between visits (Appendix 2).

Intervention: Addition of real-time feedback (VT)

The VT group, in addition to motivational sessions and retrospective feedback, received real-time feedback. VT parameters were set to meet each individual participants needs, relating to collaborative goals set during the intervention programme. Therefore, the time between the prompts (vibrations) differed from individual to individual along with the reset time. The reset time defines the time in upright activity required to restart the sitting timer. The participants could turn off the reminder at any time. It was also possible for the participants to choose to change prompt parameters.

Data collection

The study took place over a 14 week period including: baseline; 3 motivational sessions (Weeks 2, 6, 10); 4 weeks follow-up (Week 14), where the vibration was removed from the VT units. One researcher (an experienced Physiotherapist) provided the intervention and collected the data at the participant's home. From week 1 of the study

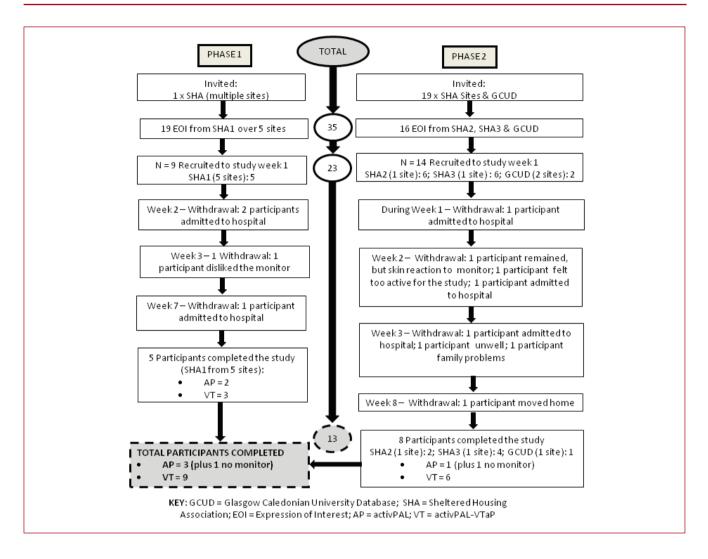


Figure 1. Consort Diagram for SOS Study.

onwards, participants were asked to wear the AP/VT activity monitor 24 hours a day. The monitor was applied by the researcher on day one of the study. After 7 days the first set of data was retrieved and the second monitor was applied to the opposite leg. At baseline the participants completed the functional tests and questionnaires about their health, well-being, physical activity and SB^{11,22}. These outcomes were repeated post study and at follow-up by the researcher un-blinded to the intervention group.

Functional performance testing

Validated functional tests were employed: Chair Sitand-Reach for flexibility²³; 30 secs STS for lower limb strength²⁴; Timed Up and Go (TUG)²⁵ for mobility and falls risk; balance was assessed by a Balance Screening Tool²⁶. All test are commonly used in Physiotherapy practice and have published 'normal values', this allowed the participants to compare their results to age related normal values during session 1. Where it was thought that the test was beyond the participant's ability, or they declined to take part, that element of testing was omitted.

Sedentary behaviour measurement by activity monitoring

The AP inclinometer is a thigh mounted objective monitor shown to be a highly accurate method of measuring sitting posture in OA^{27,28}. The AP/VT monitors were covered in medical grade waterproof packets. On the visits the monitors were applied to the mid–thigh using an adhesive pad ('activPAL Stickie'), this was then covered with a hypoallergenic tape (Opsite Fexifix). The participants were instructed to remove the monitor and telephone the researcher if any adverse event occurred. The monitor also displayed the wording 'activity monitor' and a contact

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Baseline Test	VT Group	N	AP Group	N
Sex (% female)	67	12	67	11
Age (years, Mean (SD))	74.0±6.1	12	83.2±7.7	11
BMI (kg/m², Mean (SD))	28.3±7.2	11	26.3±4.9	10
No. of Medications	5.3 ⁺⁺	12	6.5 ^{††}	11
No. educated >HS	3	12	0	11
Timed Up & Go (sec, Mean (SD))	26.6±7.1 ⁺⁺	11	29.8±16.1 ⁺⁺	5
Employed	0	12	0	11
EuroQoL(score/5)†	1.6	12	1.5	11
>1 Fall /last year (%)	50 ^{††}	12	63.6 ^{††}	11
Widow (%)	16.7	12	54.5	11
Single (%)	16.7	12	36.4	11
Divorced (%)	33.3	12	0	11
Married (%)	33.3	12	9.1	11

[†] = The European Quality of Life Questionnaire (EQ-5D-5L) average scores of self-reported: mobility, self-care, daily activities, pain, anxiety and depression collated; ^{††} = indication of frailty; BMI = Body Mass Index from self-reported height/weight; HS = High School.

Table 2. Descriptive statistics at baseline.

number in case the monitor was lost or the user was in a situation where they were unable to explain its purpose.

Data processing

The SB data was analyzed using the data from the activity monitors following the protocol of Chastin and Granat (2010)²⁹. The variables analyzed were total sedentary time and patterns of SB. To avoid a daily participant burden of keeping a sleep diary over the study period, waking day SB was assessed by the following protocol: the start of the day was defined as the first stand/step after O6.30 hrs; the end of the day was defined by the step/stand before the first sit/lie after 22:00 hrs; if there was no sit/lie commencing between 22:00 hrs and 23:59 hrs the end of day will be defined at the point of the last stand/step before 22:00 hrs. To determine validity of this protocol this was compared to sleep diaries self-reported by the participants at the baseline week.

Statistical analysis

Data were analyzed using IBM SPSS 22 software. Total sitting time is expressed as a percentage of the waking day (SB Ratio). Fragmentation is a measure of patterns of sitting, where the number of bouts of sitting time per hour of the day is divided into total sitting time (b/hr). We analyzed the data as intention-to-treat (ITT) mixed model analysis (MMA) to assess the effectiveness of the intervention to: a) improve physical function and b) to change SB. All participants that started week one were included in the analysis. In each model there was a repeated measurement of the

participants' outcomes. The model takes into consideration the factor of the group and the covariant of time for each of the dependent variables (SB or functional outcomes). As others authors had reported a diminished intervention effect with time^{11,13,16}, we developed a MMA to examine the effect of day of the week i.e. day 1 being the first day following the intervention period, and so on, along with week of the month in order to understand the longitudinal effect.

Results

Recruitment & retention of participants

Recruitment was difficult and retention was low, reflecting the frailty of the older adults recruited in the study. As can be seen from Figure 1, there were 2 phases of recruitment and the most common reason for withdrawal was participants' health status.

Descriptive statistics

Twenty-three participants (17 female; mean age: 78.8 ± 8.1 ; 12 VT group) were recruited and started week 1 (Figure 1). The majority of participants were female and despite randomization there were baseline differences in the groups (Table 2). The average waking day recording period was 14.0 ± 0.2 hrs, this correlated well with the finding of the baseline sleep diaries (mean sleep reported= 9.5 ± 0.9 hrs, therefore 14.5 hrs remain as the waking period, N=15). Thirteen participants remained to the end of the study (one without AP monitor), representing 56.5% retention.

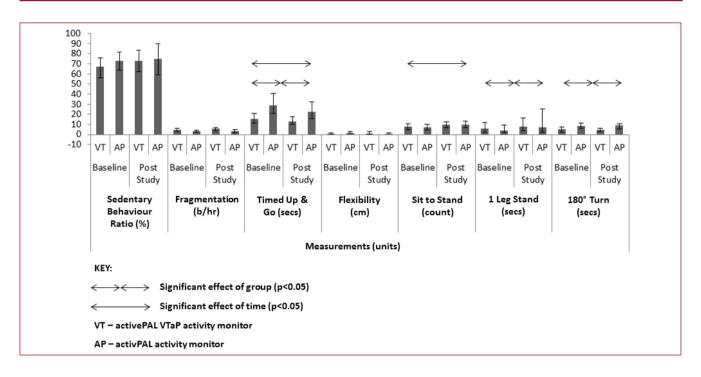


Figure 2. Marginal Mean (CI) of sedentary behaviour and function outcomes (N=23) as intention-to-treat mixed model analysis.

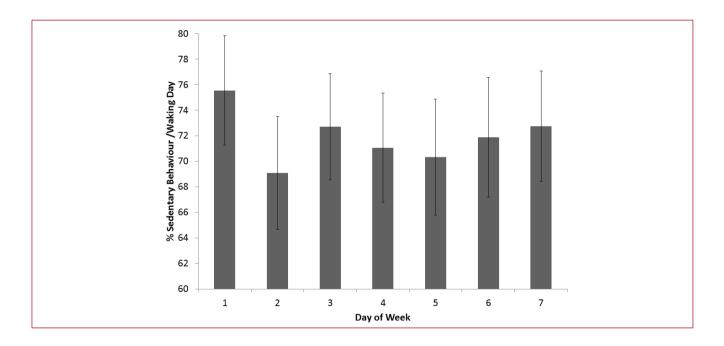


Figure 3. Marginal mean (CI) sedentary ratio by day (N=23). The figure examines the week after the motivational session has been delivered and combines each of the post-session days over the period of the study.

Intention-to-treat

When the ITT MMA is applied to the physical function data (Figure 2), the VT groups had better physical function throughout the study. There were changes over time for both

AP and VT groups. There was a significant effect of group and time for TUG test and a significant effect of time only was found in the STS test. The 1 leg stand test and 180° turn test had a significant group effect. As can be seen from Figure 2, there was no significant effect found for group or time with SB ratio, SB fragmentation or flexibility.

When the ITT MMA was applied to time periods, there was a significant effect of day (number of days after the intervention). As can be seen from Figure 3, there is a pattern of sedentary accumulation where an initial post intervention drop is observed (after each motivation session) followed by increase, then a second smaller drop. There was no significant effect of group, week or day and week, with sedentary ratio data. When the MMA of fragmentation of SB was examined, the only significant effect was group with the VT group showing more fragmentation. There was no significant effect of day, week, or day and week on fragmentation.

Adverse reactions

One participant (AP group) had a skin irritation under the area of the activity monitor and was unable to continue wearing the monitor from week 2, but remained in the study (minus activity monitor feedback).

Discussion

An intervention aimed at decreasing and breaking up long periods of SB, thus encouraging small bouts of functional training (i.e. rising from a chair) little, often, daily, maybe more risk averse and acceptable to frailer OA than formal exercise training. The older adults recruited for the SOS study were frail evidenced by the health-related drop-out, fall history, daily medication use and baseline TUG scores. Post study, both groups made favorable and clinically important gains in physical function suggesting the SOS intervention increased the participants' ability to mobilize from a chair (TUG 4 secs faster and 2 more STS in 30 secs).

These findings are comparable to other studies of younger OA with better physical function at baseline where a small, but significant, positive effect was found in gait speed¹³ and STS score¹⁵. Barone Gibbs et al.¹⁵ study compared a "Sit Less" group to a "Get Active" group. Interestingly, the "Sit Less" group made functional gains that the "Get Active" group did not make despite a significant increase in their moderate to vigorous intensity physical activity (MVPA). The functional tests used have a ceiling effect, therefore, the frailer the start point, the more gain there is available to make. Regularly breaking up sitting is an activity intervention in itself, with the true effect of exercise by stealth through repeated sitto-stands, rather than exercising only in 'sessions' and potentially sitting more in between.

Although there were no significant changes in sedentary time, fragmentation tended to improve for the VT group over time. It would seem that breaking up periods of prolonged sitting, more than total sedentary time, was easier for the older adults to gain and sustain change overtime, but only with the use of the vibrational prompts. There is also diminished effect as time progresses from the point of intervention. Both this study and Gardiner et al. noted weekly behavioural patterns where OA appear to make

positive changes in the first few post intervention days, followed by a negative change in the subsequent days, followed by a second, smaller, move towards positive change. This patterning may be due to fatigue or compensation and merits further investigation to both ensure the interventions are appropriate and sustainable³⁰.

The SOS study was a long-term intervention study with face-to-face monthly interventions, where individuals activity monitor and physical function feedback was discussed and appropriate collaborative goals were set. Other intervention studies aimed at reducing SB, tend to focus on the outcomes of SB and/or physical activity 11-16 and, to the authors knowledge, have not considered discussing participants own physical function as a motivational tool, this is therefore a novel component of the SOS study worth considering in future studies. Interestingly, a pilot study of the Functional Fitness MOT where functional normative data feedback and physical activity advice was given, found that OA expressed more interest in breaking up prolonged sitting, than in seeking out physical activity opportunities³¹. Frailer OA valued the faceto-face intervention of the SOS study32 and it would appear that information booklets only are not sufficient to instigate self-reported sedentary change in those living in sheltered

The SOS pilot study is limited by its small sample size, high attrition rates in this frail group, so was consequently underpowered to detect changes in SB. The study assessments were carried out by one researcher, not blinded to group allocation as the observations formed part of the intervention. A large RCT that addresses these limitations, with a longer follow up, is warranted.

For OA living in sheltered housing, it is appropriate to provide tailored advice to recommend reducing SB, as this will improve their sit-to-stand transfer and ability to mobilize. This study provided participants feedback on, and assessed, their physical function, along with their SB. It is recommended that future intervention studies should measure effects on function in this frail group and consider using functional feedback as a motivational tool. The novel use of the real-time feedback is also worth further investigation, although the SOS study did not unequivocally demonstrate its benefit across all outcomes.

In conclusion, it is hard to recruit and retain frailer OA to long-term RCT studies, but this vulnerable group may have the most to gain from reducing SB, thus is worth perseverance of the research and clinical communities to engage these individuals. Even in this frailer group, with a large health related drop-out, there was a significant improvement in functional ability, suggesting overall the SOS intervention had positive functional effects on frailer OA, despite no measurable change in SB.

Acknowledgements

With thanks to the residents and staff of the sheltered housing complexes that participated in this study

Ethical Approval

Glasgow Caledonian University Ethical Committee (Reference Number: HLS12/59).

Funding

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Contributors

All authors participated in the conception of this study, interpretation of the results and reviewed the manuscript. JH carried out recruitment, the intervention and data collection, analysis and writing of the manuscript.

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Appendix 1. The TIDieR Checklist (page numbers relate to the originally submitted manuscript).

	The TiDieR (Template for Intervention Description and Replic				
Information to include when describing an intervention and the location of the information					
ltem	Item	Where lo	ocated **		
number		Primary paper (page or appendix number)	Other†(details		
	BRIEF NAME	P1			
1.	Provide the name or a phrase that describes the intervention.				
	WHY	P1			
2.	Describe any rationale, theory, or goal of the elements essential to the intervention.				
	WHAT	P4-P7			
3.	Materials: Describe any physical or informational materials used in the intervention, including those				
	provided to participants or used in intervention delivery or in training of intervention providers.				
	Provide information on where the materials can be accessed (e.g. online appendix, URL).	P4-P7			
4.	Procedures: Describe each of the procedures, activities, and/or processes used in the intervention,				
•	including any enabling or support activities.				
	WHO PROVIDED	P5			
5.	For each category of intervention provider (e.g. psychologist, nursing assistant), describe their	13			
э.					
	expertise, background and any specific training given.	D.F.			
	HOW	P5			
6.	Describe the modes of delivery (e.g. face-to-face or by some other mechanism, such as internet or				
	telephone) of the intervention and whether it was provided individually or in a group.				
	WHERE	P5			
7.	Describe the type(s) of location(s) where the intervention occurred, including any necessary				
	infrastructure or relevant features.				
	WHEN and HOW MUCH	P5			
8.	Describe the number of times the intervention was delivered and over what period of time including				
0.	the number of sessions, their schedule, and their duration, intensity or dose.				
	TAILORING	P5			
9.	If the intervention was planned to be personalised, titrated or adapted, then describe what, why,				
	when, and how.				
	MODIFICATIONS	P9			
10. *	If the intervention was modified during the course of the study, describe the changes (what, why,				
	when, and how).				
	HOW WELL				
11.	Planned: If intervention adherence or fidelity was assessed, describe how and by whom, and if any	N/A			
	strategies were used to maintain or improve fidelity, describe them.				
12.‡	Actual: If intervention adherence or fidelity was assessed, describe the extent to which the	N/A			
	intervention was delivered as planned.	- 100 1			

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^{**} Authors - use N/A if an item is not applicable for the intervention being described. Reviewers - use '?' if information about the element is not reported/not sufficiently reported.

[†] If the information is not provided in the primary paper, give details of where this information is available. This may include locations such as a published protocol or other published papers (provide citation details) or a website (provide the URL).

[‡] If completing the TIDieR checklist for a protocol, these items are not relevant to the protocol and cannot be described until the study is complete.

^{*} We strongly recommend using this checklist in conjunction with the TIDieR guide (see BMJ 2014;348:£1687) which contains an explanation and elaboration for each item.

^{*} The focus of TIDIER is on reporting details of the intervention elements (and where relevant, comparison elements) of a study. Other elements and methodological features of studies are covered by other reporting statements and checklists and have not been duplicated as part of the TIDIER checklist. When a randomised trial is being reported, the TIDIER checklist should be used in conjunction with the CONSORT statement (see www.consort-statement.org) as an extension of Item 5 of the CONSORT 2010 Statement. When a clinical trial protocol is being reported, the TIDIER checklist should be used in conjunction with the SPIRIT statement as an extension of Item 11 of the SPIRIT 2013 Statement (see www.spirit-statement.org). For alternate study designs, TIDIER can be used in conjunction with the appropriate checklist for that study design (see www.equator-network.org).

Appendix 2. The support pack was personalized to each individual and included self-completing sections, thus allowing the participants to have a sense of responsibility and control over their behaviour (the intervention pack is available here: https://sedentaryblethering.files.wordpress.com/2017/07/sos-intervention-pack-juliet-harvey.pdf).

