The development of an intervention for reducing sitting time in the workplace

Submission date 21/11/2016	Recruitment status No longer recruiting	Prospectively registered[X] Protocol
Registration date 02/12/2016	Overall study status Completed	[_] Statistical analysis plan[X] Results
Last Edited 07/08/2020	Condition category Other	Individual participant data

Plain English summary of protocol

Background and study aims

Long periods of sitting down (e.g. during the working day) increase a person's risk of heart disease and diabetes. This has been demonstrated even for people who exercise regularly. Unfortunately with the rise of computer working more and more people are sitting for long periods of time during the day, and so this represents an urgent public health concern. This study aims to develop a new approach to helping people reduce their sitting time based on established psychological strategies for changing behaviour.

Who can participate?

Office workers aged 18 or over at King's College London

What does the study involve?

In the first week of the study participants are monitored using inclinometer devices which record when they are sitting or standing. Participants also keep track of the tasks they do at work in order to see whether they sit or stand more during particular types of task. After this 'monitoring' period participants are visited by the researcher for a session where they are given feedback on their sitting patterns during the monitoring week. Participants are then provided with 'sit-stand' desks which allow them to easily change between a sitting and standing position during their working day. Participants also choose from a range of strategies to change their sitting behaviour patterns, and importantly, to ensure that their change in behaviour is maintained in the long term. Participants use the desks for 12 weeks during which time they are monitored with the inclinometer devices for one week at a time 1 week later, 5 weeks later and 11 weeks later. After each of these monitoring weeks the researcher meets with the participants and interviews them to find out about their experiences of using both the sit-stand desk and the strategies for reducing their sitting.

What are the possible benefits and risks of participating?

The benefits of the study include the use of the sit-stand desk for 3 months, the opportunity to work with the researcher to tailor a range of strategies to their needs to help them reduce their sitting, and finally the receipt of a £100 Amazon voucher upon completion of every aspect of the study. The risks of taking part in the study surround the use of the sit-stand desk itself. Prolonged standing can lead to discomfort and even injury, particularly if there is a pre-existing

condition. Recommendations are provided throughout the study to reduce the chance of this occurring.

Where is the study run from? King's College London (UK)

When is the study starting and how long is it expected to run for? September 2016 to June 2017

Who is funding the study? Medical Research Council (UK)

Who is the main contact? Dr Stephen Dewitt

Contact information

Type(s) Scientific

Contact name Dr Stephen Dewitt

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers N/A

Study information

Scientific Title

Development and piloting of an intervention to reduce workplace sitting time: the REducing SItting Time Study (the RESIT study)

Acronym RESIT

Study objectives

The aim of the current study is to gain insight into the most efficacious methods of reducing sitting time in the workplace for desk-based workers in order to develop an intervention protocol.

Ethics approval required Old ethics approval format

Ethics approval(s)

Psychiatry, Nursing & Midwifery Research Ethics Panel (King's College London), 29/09/2016, ref: LRS-16/17-3718

Study design Uncontrolled pre-post design

Primary study design Interventional

Secondary study design Non randomised study

Study setting(s) Other

Study type(s) Other

Participant information sheet

Not available in web format, please use contact details to request a participant information sheet

Health condition(s) or problem(s) studied

Sedentary behaviour

Interventions

The study uses an uncontrolled, pre-post design among a sample of 30 KCL office workers, to codesign and assess the acceptability of a workplace sitting reduction intervention, with three main follow-up points.

This is a pilot study with only one condition (the intervention).

The intervention comprises three key 'ingredients': firstly, a behaviour and cue-monitoring phase, with feedback; secondly, the provision of a height-adjustable desk for 3 months; and thirdly, a 'menu' of behaviour change techniques from which participants can select to customise the intervention to their needs.

In the cue monitoring phase, participants are asked firstly to wear an activPAL accelerometerinclinometer device for one week and secondly to keep a record of the tasks they undertake at work during that week. At the end of this week they will also be asked to provide a subjective measure of the amount of time spent sitting during work for that week.

Ten days later, in the intervention session, participants will be first provided feedback on the cue monitoring week. Their subjective measure of sitting time will be compared to the objectively measured time, to raise awareness of their sitting behaviour. Any emergent relationship between sitting duration and time of day, day of the week, or task type will be presented to participants. Following this participants will be provided with a VariDesk Pro Plus 30 'sit-stand' desk and will be provided guidance and tips on its use.

In the same session participants' opportunity, motivation and capability to reduce their sitting time will be assessed using the COM-B model (Michie, Atkins & West, 2014). Based on participants' answers to these questions they will be offered a range of behaviour change techniques including a range of goal setting, action planning, habit formation, problem solving, habit disruption and motivational quotes from other sit-stand desk users. Following the intervention session all participants will be sent a summary of the information provided.

Follow up interviews will be undertaken 1, 6 and 12 weeks after the intervention session. In each of these sessions the researcher will collect the accelerometer (fitted one week prior in each case) and run a semi-structured interview. The interview schedule at week one only will include questions on the participant's initial motivation to participate in the study, motivation to reduce sitting time and their initial expectations about reducing sitting time. All three interview schedules will then include questions on the participant's experiences of standing since the previous meeting, their perceptions of their capability, motivation on opportunity to stand over that period, questions on the conduciveness of the physical and social office environment to reducing sitting. Finally they will be asked questions related to their specific choice of behaviour change interventions to gain insight into their experience of employing these techniques. At the final session (week 12) the sit-stand workstation will be removed.

Intervention Type

Behavioural

Primary outcome measure

 Periods of, and transitions between, sitting and standing, measured using accelerometerinclinometer devices over one week at 1, 5 and 11 weeks after the intervention session
 Qualitative data from responses to open-ended interview questions across three sessions 1, 6 and 12 weeks after the intervention session

Secondary outcome measures

1. Participant records of tasks undertaken during monitoring week from Day 1 to Day 8 2. Participants' subjective measure of sitting time on Day 10

Overall study start date 18/09/2016

Completion date 01/06/2017

Eligibility

Key inclusion criteria

1. Office- and desk-based KCL employees whose job requires them to sit at a dedicated workstation (i.e. not a 'hot-desker') for the majority of their working day and to follow this working day pattern at least 3 days per week

2. Aged 18 or over (there is no upper age limit)

3. Able to stand at work (i.e., no physical impairment precluding standing in the workplace)

Participant type(s)

Healthy volunteer

Age group

Adult

Lower age limit

18 Years

Sex

Both

Target number of participants 30

Total final enrolment

29

Key exclusion criteria

 Participants must not have taken part in similar 'standing while working' research previously or used a sit-stand desk at their work station for two or more days in a row
 They also must not have plans to leave KCL or plans to take an absence for longer than 10 consecutive work days for the duration of the study period (October 2016 to June 2017)

Date of first enrolment

15/10/2016

Date of final enrolment 31/05/2017

Locations

Countries of recruitment England

United Kingdom

Study participating centre King's College London James Clerk Maxwell Building 57 Waterloo Road

London United Kingdom SE1 8WA

Sponsor information

Organisation King's College London

Sponsor details Psychology Department, IoPPN Henry Wellcome Building De Crespigny Park Denmark Hill London England United Kingdom SE5 8AF +44 (0)20 3228 3084 psychology@kcl.ac.uk

Sponsor type University/education

ROR https://ror.org/0220mzb33

Funder(s)

Funder type Research council

Funder Name Medical Research Council

Alternative Name(s) Medical Research Council (United Kingdom), UK Medical Research Council, MRC

Funding Body Type Government organisation

Funding Body Subtype National government

Results and Publications

Publication and dissemination plan

1. The protocol for the study is intended to be published prior to completion of recruitment 2. Upon completion the study will be published in a peer-reviewed journal, will be presented at conferences and will be written in a plain English form and published in blog format. Further details are to be confirmed at a later date

Intention to publish date

01/06/2018

Individual participant data (IPD) sharing plan

The datasets generated during and/or analysed during the current study are/will be available upon request from Stephen Dewitt or Benjamin Gardner

IPD sharing plan summary

Available on request

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Protocol article	protocol	28/11/2017		Yes	No
<u>Results article</u>	results	01/12/2019	07/08/2020	Yes	No