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Booster Breaks: health promoting work breaks

Submission date
16/02/2015

Recruitment status
No longer recruiting

Registration date
26/03/2015

Overall study status
Completed

Last Edited
25/09/2017

Condition category
Other

[_] Prospectively registered
[_] Protocol
[_] Statistical analysis plan
[X] Results
[_] Individual participant data

Plain English summary of protocol

Background and study aims
The workplace is an important potential setting for physical activity (PA) interventions (i.e. programmes). However, few workplace PA interventions take full advantage of work breaks and result in limited-to-modest success. Booster Breaks are organized work breaks that are designed to improve employees physical and psychological health, increase job satisfaction and improve work productivity. Booster Breaks have been developed to alleviate work-related stress and encourage more PA behavior in employees that may otherwise have sedentary jobs. Examples of Booster Breaks include short exercise or meditation sessions. The aim of this study is to test whether Booster Breaks increase PA among sedentary employees, compared with individualized PA work breaks and typical work breaks.

Who can participate?
Employees with sedentary office jobs from four workplaces in a large, urban southwestern U.S. city.

What does the study involve?
Participants are randomly allocated into one of three groups. Those in group 1 are assigned to the Usual Break (control) group. Those in group 2 are assigned to the Computer Prompt (individualized PA work breaks) group. Those in group 3 are assigned to the Booster Break group. The Usual Break condition includes usual or typical work break practices and behaviors. The two interventions are developed to be consistent with the WHO Healthy Workplace Framework and Model. The individualized Computer Prompt condition is designed to interrupt prolonged sitting time by introducing 3-minute breaks, 5 times per day. The group-based Booster Break condition is a peer-led, once-a-day, 15-minute PA session that guides employees through a series of stretching, strengthening, and aerobic movements followed by a brief meditation. Based on a participation rate threshold of 70%, workplaces were classified either as consistent or inconsistent implementers of the intervention. A number of physiological and behavioral measures are assessed before the study starts and 6 months later.

What are the possible benefits and risks of participating?
Not provided at time of registration

Where is the study run from?
Four different work places in a large, urban southwestern U.S. city

When is the study starting and how long is it expected to run for?
January 2009 to January 2013

Who is funding the study?
National Institutes of Health (USA)

Who is the main contact?
Dr Wendell Taylor

Contact information

Type(s)
Scientific

Contact name
Dr Wendell Taylor

Contact details
The University of Texas Health Science Center at Houston
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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
NIH (Grant No. R03 NR010291)

Study information

Scientific Title
Impact of Booster Breaks on physical activity among sedentary employees: a cluster randomized controlled trial

Study objectives
It was hypothesized that, compared with non-Booster Break participants, Booster Break participants will have significant improvements in:
1. Physiological measures (i.e., blood pressure, fasting lipids, triglycerides, and anthropometrics)

2. PA (increase) and sedentary behavior (decrease)
3. PA mediators
4. Employee and organizational psychosocial constructs

Ethics approval required
Old ethics approval format

Ethics approval(s)
This study was approved by the appropriate IRBs and the University Committee for the Protection of Human Subjects

Study design
Randomized controlled trial

Primary study design
Interventional

Secondary study design
Randomised controlled trial

Study setting(s)
Community

Study type(s)
Prevention

Participant information sheet
Not available in web format, please use the contact details to request a patient information sheet

Health condition(s) or problem(s) studied
Physical inactivity, sedentary behavior, obesity

Interventions
Participants were randomized to the Usual Break (control group), Computer Prompt, or Booster Break condition. The Usual Break condition included usual or typical work break practices and behaviors. The two interventions were developed to be consistent with the WHO Healthy Workplace Framework and Model. The individualized Computer Prompt condition was designed to interrupt prolonged sitting time by introducing 3-minute breaks, 5 times per day. The group-based Booster Break condition was a peer-led, once-a-day, 15-minute PA session that guides employees through a series of stretching, strengthening, and aerobic movements followed by a brief meditation.

Intervention Type
Behavioural

Primary outcome measure
1. Lipid profile
2. Blood pressure
3. Height
4. Weight

5. International Physical Activity Questionnaire (IPAQ)
6. Pedometer readings

Measures were taken at baseline and immediately after the intervention (i.e after 6 months)

Secondary outcome measures
Physical activity mediators and employee and organizational psychosocial constructs: self-report assessments at baseline and immediately after the intervention (i.e after 6 months)

Overall study start date
09/01/2009

Completion date
09/01/2013

Eligibility

Key inclusion criteria
1. Participants' jobs required sitting for at least 5 hours per day
2. English proficiency
3. Full-time employment [35–40 hours/week]
4. Age >17 years
5. No physician-limited physical activity

Participant type(s)
Healthy volunteer

Age group
Adult

Sex
Both

Target number of participants
3 clusters, 5 departments in each cluster, 5 to 15 individuals in each department

Key exclusion criteria
1. Physician prohibited physical activity
2. Part-time employees

Date of first enrolment
09/01/2009

Date of final enrolment
09/01/2012

Locations

Countries of recruitment
United States of America

Study participating centre
The University of Texas Health Science Center at Houston
School of Public Health
CHPPR
7000 Fannin Street
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Sponsor information

Organisation
The University of Texas Health Science Center at Houston

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Sponsor type
University/education

ROR
https://ror.org/03gds6c39

Funder(s)

Funder type
Government

Funder Name
National Institutes of Health

Alternative Name(s)
Institutos Nacionales de la Salud, US National Institutes of Health, NIH

Funding Body Type
Government organisation

Funding Body Subtype
National government

Location
United States of America

Results and Publications

Publication and dissemination plan

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not expected to be made available

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	feasibility study results	01/08/2010		Yes	No
Other publications	participants' perspectives	01/06/2013		Yes	No
Results article	results	17/11/2016		Yes	No