

# Booster Breaks: health promoting work breaks

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<b>Registration date</b> 26/03/2015	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
<b>Last Edited</b> 25/09/2017	<b>Condition category</b> Other	<input type="checkbox"/> 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

**Protocol serial number**  
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

## **Study type(s)**

Prevention

## **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(s)**

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)

## **Key secondary outcome(s)**

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

## **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

**Healthy volunteers allowed**

No

**Age group**

Adult

**Sex**

All

**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

Suite 2670

Houston

United States of America

TX 77030

**Sponsor information**

## Organisation

The University of Texas Health Science Center at Houston

## ROR

<https://ror.org/03gds6c39>

## Funder(s)

### Funder type

Government

### Funder Name

National Institutes of Health

### Alternative Name(s)

US National Institutes of Health, Institutos Nacionales de la Salud, NIH, USNIH

### Funding Body Type

Government organisation

### Funding Body Subtype

National government

### Location

United States of America

## Results and Publications

### 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?
<a href="#">Results article</a>	feasibility study results	01/08/2010		Yes	No
<a href="#">Results article</a>	results	17/11/2016		Yes	No
<a href="#">Other publications</a>	participants' perspectives	01/06/2013		Yes	No