CARDIOMETABOLIC RISK OF PROLONGED SITTING IN YOUNG ADULTS * A PILOT STUDY

Published: 03-08-2011 Last updated: 29-04-2024

Primary Objective:To investigate the feasibility of studying of the effects of 6 days of predominantly prolonged sitting on cardiometabolic health in young adults. Secondary

Objective: To collect preliminary data on the effects of 6 days of...

Ethical review Approved WMO **Status** Recruitment stopped

Health condition type Glucose metabolism disorders (incl diabetes mellitus)

Study type Observational invasive

Summary

ID

NL-OMON38241

Source

ToetsingOnline

Brief title

CARDIOMETABOLIC RISK OF SITTING - A PILOT

Condition

• Glucose metabolism disorders (incl diabetes mellitus)

Synonym

Indicators of dysfunction of glucose and triglyceride metabolism: elevated blood plasma levels of glucose, insulin and triglycerides.

Research involving

Human

Sponsors and support

Primary sponsor: Vrije Universiteit Medisch Centrum

Source(s) of monetary or material Support: Ministerie van OC&W

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Intervention

Keyword: BLOOD, CARDIOMETABOLIC RISK, PROLONGED SITTING, YOUNG ADULTS

Outcome measures

Primary outcome

The main outcome measures of this pilot study are blood pressure and plasma

levels of glucose, insulin, triglycerides and NEFA in all blood samples.

Secondary outcome of this pilot study is the autonomic nerve

Secondary outcome

N.A.

Study description

Background summary

Recent preliminary experimental findings in obese adults on the acute effects of prolonged sitting show a detrimental increase in triglycerides prolonged sitting during a 7-hr sitting day compared to a non-sitting day. Breaking up the 7-hr sitting every 20min can attenuate these detrimental acute effects on glucose metabolism. In the first part of our pilot study we additionally demonstrated in healthy young adults, that hourly interruptions in sitting can also attenuate these detrimental acute effects. However, no evidence exists on the deterimental effects of repeated uninterrupted sitting. This information is highly necessary to obtain insight in the detrimental health effects of a sedentary lifestyle, e.g. reflecting daily life.

Study objective

Primary Objective:

To investigate the feasibility of studying of the effects of 6 days of predominantly prolonged sitting on cardiometabolic health in young adults.

Secondary Objective:

To collect preliminary data on the effects of 6 days of predominantly prolonged sitting on cardiometabolic health in young adults.

Study design

Pilot study involving two laboratory sitting days, similar to the prolonged sitting condition in the first part of this pilot study. In addition, each participants will complete 6 days of 'lifestyle sitting', consisting of reducing the interruptions in sitting time as much as possible, in between the two sitting days.

Study burden and risks

Participants will visit the laboratory on two occasions (two laboratory sitting days), having fasted for at leas 10 hours. Anthropometrics (e.g. weight, height, waist- and hip circumference, body fat percentage) will be measured at baseline. Venous blood will be collected at baseline and hourly during the 8-hour laboratory days (9 samples in total for each sitting day), using an in-dwelling catheter. At similar time points blood pressure will be measured using an automatic oscillometric method. Participants will complete 6 days of lifestyle sitting in between the two laboratory sitting days. After completion of this pilot study, participant will be motivated to minimize their time spent on prolonged sitting and increase their time spent on physical activity. They will receive a pedometer, which they can keep, in order to increase motivation. There is no risk associated with participants in this pilot experiment.

Contacts

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Trial sites

Listed location countries

Netherlands

Eligibility criteria

Age

Adults (18-64 years) Elderly (65 years and older)

Inclusion criteria

Being apparently healthy, having an active lifestyle (i.e. at least 30 minutes moderate to vigorous physical activity at each day of the week), aged 18-23, Dutch or English speaking, and signed informed consent.

Exclusion criteria

Known physical activity contraindications, major illness/injury (acute or chronic) or physical problems that may limit the ability to perform the experiment.

Study design

Design

Study type: Observational invasive

Intervention model: Other

Allocation: Non-randomized controlled trial

Masking: Open (masking not used)

Control: Active

Primary purpose: Basic science

Recruitment

NL

Recruitment status: Recruitment stopped

Start date (anticipated): 02-12-2011

Enrollment: 10

Type: Actual

Ethics review

Approved WMO

Date: 03-08-2011

Application type: First submission

Review commission: METC Amsterdam UMC

Approved WMO

Date: 10-01-2012

Application type: Amendment

Review commission: METC Amsterdam UMC

Approved WMO

Date: 20-04-2012

Application type: Amendment

Review commission: METC Amsterdam UMC

Study registrations

Followed up by the following (possibly more current) registration

No registrations found.

Other (possibly less up-to-date) registrations in this register

No registrations found.

In other registers

Register ID

CCMO NL37053.029.11