

Neural effects of exercise, diet and sleep

Submission date 10/11/2015	Recruitment status No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 16/11/2015	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 30/12/2022	Condition category Mental and Behavioural Disorders	<input type="checkbox"/> Individual participant data <input type="checkbox"/> Record updated in last year

Plain English summary of protocol

Background and study aims

The brain of a child and that of an adult are organised very differently. In adolescence the brain undergoes extensive remodelling, maturing and building connections. It is during this time that the cerebral cortex (the folded, outer part of the brain) undergoes extensive changes, marking the development of complex thinking processes, increasing a persons' capacity for learning. There are many known benefits of taking part in regular physical activity for the body, but there is a good deal of evidence that it is also valuable for healthy cerebral development. Despite this, teenagers are becoming increasingly inactive, many becoming overweight or obese. Many studies have shown that cognitive function (mental abilities such as thinking, reasoning, memory and attention) is improved after taking part in exercise in children, boosting academic performance. The aim of this study is to find out whether taking part in high-intensity exercise will help to improve cognitive function and learning in adolescents.

Who can participate?

Healthy adolescents aged between 15 and 19 who attend a participating high school or vocational school.

What does the study involve?

Participants are randomly allocated to one of two groups. Those in the first group take part in three 20 minute exercise sessions every week during school hours. In these sessions, participants complete a warm up (low intensity), and then high intensity interval running exercise such as running or circuit training, followed by a short low intensity cool down. Those in the second group continue with their usual exercise regime and have no extra training sessions. At the start of the study and then again after one months, participants in both groups have a scan to test their brain activity, as well as completing a number of tests to find out if there has been any change to their cognitive function. A subgroup of the study population also have a scan to test their brain activity before and after a single high intensity interval exercise to study the effects of acute exercise on brain functioning.

What are the possible benefits and risks of participating?

Participants will benefit from taking part in the study as they will be able to receive detailed information about their health and fitness levels, as well as feedback about any changes they could make to improve. There is a small risk of injury during the exercise training, although measures will be taken to prevent this.

Where is the study run from?
University of Jyväskylä (Finland)

When is the study starting and how long is it expected to run for?
January 2015 to December 2018

Who is funding the study?
Jenny and Antti Wihuri Foundation (Finland)

Who is the main contact?
Dr Eero Haapala

Contact information

Type(s)
Scientific

Contact name
Dr Eero Haapala

ORCID ID
<https://orcid.org/0000-0001-5096-851X>

Contact details
Sport and Exercise Medicine
Faculty of Sport and Health Sciences
PO-Box 35 (VIV)
University of Jyväskylä
Jyväskylä
Finland
40014

Additional identifiers

Protocol serial number
N/A

Study information

Scientific Title
High-intensity exercise training intervention study to improve cognitive functions and learning in adolescents

Acronym
NEEDS

Study objectives
High-intensity and low-volume exercise intervention will improve cognitive function and learning in the study group compared to the control group during the three month intervention period.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Ethics committee of University of Jyväskylä, 25/08/2015

Study design

Current study design:

Single-centre single-blind parallel controlled cross-over study

Previous study design:

Single-centre single-blind cluster-randomized controlled trial

Primary study design

Interventional

Study type(s)

Other

Health condition(s) or problem(s) studied

Cognition, learning, brain function

Interventions

Current Interventions as of 17/03/2017:

Participants are allocated to the intervention group or the control group.

Intervention group: Participants take part in high-intensity and low volume exercise training during school days. The aim of the exercise intervention is to provide three short (approximately 20 minute) exercise sessions weekly and to improve cardio-respiratory fitness, totaling in all 12 sessions over the 1-month intervention period. As a training mode high intensity running will be adopted. The protocol includes 3 minute warm-up at light to moderate intensity and eight to twelve repeated bouts of 0.5 to 1 minute at 90% to 100% of the capacity of the participants interspersed by 75–90 second recovery at light intensity, followed by 3 minute recovery period at light to moderate intensity.

In the sub-study, the participants participate in three brain scans. First scan is performed two weeks before second visit to the laboratory. During the second visit, the participants perform a brain scan before and after the acute high intensity interval exercise. The exercise is performed using mechanically braked cycle ergometer. The exercise protocol include eight 30-second maximal intensity exercise bouts interspersed by 90-second active recovery.

Control Group: Participants will continue their usual physical activity behavior without supervised exercise by the study.

There is no follow-up planned.

Previous Interventions:

Participants are randomly allocated to the intervention group or the control group.

Intervention group: Participants will take part in high-intensity and low volume exercise training during school days. The aim of the exercise intervention is to provide three short (approximately

20 minute) exercise sessions weekly and to improve cardio-respiratory fitness and motor performance, totaling in all 24 sessions over the 2-month intervention period. As a training mode high intensity running and circuit training will be adopted. The protocol includes 3 minute warm-up at light to moderate intensity and eight to twelve repeated bouts of 1 minute at 90% to 100% of the capacity of the participants interspersed by 75–90 second recovery at light intensity, followed by 3 minute recovery period at light to moderate intensity.

Control Group: Participants will continue their usual physical activity behavior without supervised exercise by the study.

After the intervention period, the participants will be followed-up until the end of vocational and high school for approximately for two years (from baseline) but no intervention is provided during follow-up.

Intervention Type

Behavioural

Primary outcome(s)

Current primary outcome measures as of 17/03/2017:

1. Attention, working memory, associative learning, processing speed and executive function is measured using the computerized CogState battery at the baseline and at the 3 months
2. Brain functions by magnetoencephalography (MEG) in a subsample of participants taking part in sub study investigating the effects of acute high intensity interval training on cognition and brain functioning.

Previous primary outcome measures:

1. Attention, working memory, associative learning, processing speed and executive function is measured using the computerized CogState battery at the baseline and at the 3 months
2. Reading and arithmetic skills are measured using pseudo word reading and KTLT tests, respectively, at the baseline and at the 3 months
3. Brain functions by magnetoencephalography (MEG) at baseline and 3 months

Key secondary outcome(s)

Current secondary outcome measures as of 17/03/2017:

1. Cardiorespiratory fitness is measured using maximal cycle ergometer test with respiratory gas analyses at baseline and 3 months
2. Physical activity is measured using PA-3D accelerometer, the Youth Physical Activity Questionnaire and the Youth Sedentary Behaviour questionnaire at baseline and 3-months
3. Sleep length and quality is measured using the Epworth sleepiness scale and Basic Nordic Sleep Questionnaire at baseline and 3 months
4. Arterial stiffness is measured using an Arteriograph at baseline and 3 months
5. Motivation is measured using a questionnaire developed by Professor Niemivirta in the 2002 study "Motivation and performance in context: The influence of goal orientations and instructional setting on situational appraisal and task performance" at baseline and 3 months
6. Self-efficacy is measured using a questionnaire developed by Professor Niemivirta in the 2002 study "Motivation and performance in context: The influence of goal orientations and instructional setting on situational appraisal and task performance" at baseline and 3 months
7. Depression is measured using the 21 item Beck Depression Scale at baseline and 3 months

Previous secondary outcome measures:

1. Cardiorespiratory fitness is measured using maximal cycle ergometer test with respiratory gas

analyses at baseline and 3 months

2. Physical activity is measured using PA-3D accelerometer, the Youth Physical Activity

Questionnaire and the Youth Sedentary Behaviour questionnaire at baseline and 3-months

3. Sleep length and quality is measured using the Epworth sleepiness scale and Basic Nordic Sleep Questionnaire at baseline and 3 months

4. Arterial stiffness is measured using an Arteriograph at baseline and 3 months

5. Adiposity is measured using dual-energy x-ray absorptiometry at baseline and using bio-impedance device at baseline and 3 months

6. Motivation is measured using a questionnaire developed by Professor Niemivirta in the 2002 study "Motivation and performance in context: The influence of goal orientations and instructional setting on situational appraisal and task performance" at baseline and 3 months

7. Self-efficacy is measured using a questionnaire developed by Professor Niemivirta in the 2002 study "Motivation and performance in context: The influence of goal orientations and instructional setting on situational appraisal and task performance" at baseline and 3 months

8. Depression is measured using the 21 item Beck Depression Scale at baseline and 3 months

Completion date

30/12/2018

Eligibility

Key inclusion criteria

Current inclusion criteria as of 17/03/2017:

1. Aged 15-19 years of age
2. Attending high school and vocational schools

Previous inclusion criteria:

1. Aged 15-17 years of age
2. Attending high school and vocational schools

Participant type(s)

Healthy volunteer

Healthy volunteers allowed

No

Age group

Child

Lower age limit

15 years

Upper age limit

19 years

Sex

All

Key exclusion criteria

1. Heart disease
2. Untreated or poorly controlled type 1 diabetes
3. Musculoskeletal disease or trauma
4. Severe depression or anxiety

Date of first enrolment

01/12/2015

Date of final enrolment

30/03/2017

Locations

Countries of recruitment

Finland

Study participating centre

University of Jyväskylä

Department of Biology of Physical Activity

Jyväskylä

Finland

FI-40014

Sponsor information

Organisation

University of Jyväskylä

ROR

<https://ror.org/05n3dz165>

Funder(s)

Funder type

Charity

Funder Name

Jenny ja Antti Wihurin Rahasto

Alternative Name(s)

Jenny JA Antti Wihurin Rahasto sr, Jenny and Antti Wihuri Foundation

Funding Body Type

Private sector organisation

Funding Body Subtype

Trusts, charities, foundations (both public and private)

Location

Finland

Funder Name

Päivikki and Sakari Sohlberg Foundation

Results and Publications

Individual participant data (IPD) sharing plan

Not provided at time of registration

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Other publications		01/09/2020	30/12/2022	Yes	No
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes