

# The health promoting potential of nature experiences in virtual reality

<b>Submission date</b> 11/08/2020	<b>Recruitment status</b> No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 28/08/2020	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
<b>Last Edited</b> 06/09/2022	<b>Condition category</b> Other	<input type="checkbox"/> Individual participant data

## Plain English summary of protocol

Background and study aims.

Green exercise (any physical activity in the presence of nature) is proposed as a key cost-effective nature-based solution to promote health and well-being. However, green exercise is often subjected to a number of barriers, such as weather conditions and accessibility to quality natural environments. Virtual Reality (VR), a computer-generated virtual environment, can in this context provide a valuable supplement to integrate nature exposure and green exercise.

The specific purposes of this project are:

1. To establish how to best develop a high-quality GreenVR;
2. To study how healthy adults perceive and respond to different GreenVRs, in order to establish which type of GreenVR provides the best mental health outcomes;
3. To evaluate the risk of our GreenVR to induce cyber-sickness, identifying factors that can cause and prevent it.
4. To evaluate the potential of GreenVR technology as a strategy to promote more participation in real green exercise.

Who can participate?

Healthy adults with normal or corrected to normal vision and limited experience with virtual reality (less than monthly use).

What does the study involve?

Participants will be required to meet at the sport-physiology laboratory at INN University and watch a short video clip validated to induce feelings of sadness. The participant will then be randomly assigned to either walk on a treadmill while watching a virtual walk in nature or walk on a treadmill without the virtual experience. Two virtual scenarios will be tested, both showing the exact same environment, but developed using different techniques: one will be developed using commercial 360° video-camera, while the other will be developed using video-games technology. To avoid 'placebo effects', the participants won't be told what condition they will undergo. Questionnaires and measures of heart rate and blood pressure will be administered before, during, and after the treadmill walk.

What are the possible benefits and risks of participating?

All participants will receive a gift certificate of 100 NKR. Some may experience improved mood

and increased motivation to exercise outdoors. On the other hand, some may experience minor negative side effects associated with cyber-sickness, such as dizziness and general discomfort.

Where is the study run from?

INN University, Elverum, Norway.

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

April 2020 to December 2020

Who is funding the study?

Inland Norway University of Applied Sciences and RFF Innlandet

Who is the main contact?

Prof Giovanna Calogiuri, giovanna.calogiuri@inn.no)

### **Study website**

<https://app.cristin.no/projects/show.jsf?id=624906>

## **Contact information**

### **Type(s)**

Scientific

### **Contact name**

Prof Giovanna Calogiuri

### **ORCID ID**

<http://orcid.org/0000-0003-1289-1026>

### **Contact details**

Inland Norway University of Applied Sciences (INN University)

Postboks 400

Elverum

Norway

2418

+47 62430245

[giovanna.calogiuri@inn.no](mailto:giovanna.calogiuri@inn.no)

### **Type(s)**

Scientific

### **Contact name**

Ms Sigbjørn Litleskare

### **ORCID ID**

<http://orcid.org/0000-0001-8322-2837>

### **Contact details**

Inland Norway University of Applied Sciences (INN University)

Postboks 400

Elverum  
Norway  
2418  
+47 62 43 02 18  
sigbjorn.litleskare@inn.no

## Additional identifiers

### EudraCT/CTIS number

Nil known

### IRAS number

### ClinicalTrials.gov number

Nil known

### Secondary identifying numbers

Cristin-Project-ID: 624906

## Study information

### Scientific Title

GreenVR - Developing and validating health-promoting VR systems

### Acronym

GreenVR 2020

### Study objectives

H1 – virtual green exercise will be characterized by lower effort relative to exercise intensity compared to regular indoor exercise, with no difference between a 360° video and a 3D-model  
H2 – virtual green exercise will be perceived as a more positive experience compared to regular indoor exercise, with no difference between a 360° video and a 3D-model  
H3 – virtual green exercise will have a more positive impact on measures of psychophysiological health compared to regular indoor exercise, with a superior effect for 3D-model compared to 360° video for affective responses  
H4: Virtual green exercise will lead to increased ratings of connectedness with nature and future green exercise intention, with no difference between a 360° video and a 3D-model  
H5: The increased ratings of future green exercise intention will be associated with the participants' changes in connectedness with nature, affective responses (enjoyment and changes in positive affect, tranquillity, negative affect, and fatigue) and perceived environmental restorativeness (fascination, being away, extent, and compatibility)

### Ethics approval required

Old ethics approval format

### Ethics approval(s)

Approved 13/08/2020, Regional committee for medical and health-related research (REK) (Gullhaugveien 1-3, 0484 Oslo, Norway; +47 22 84 55 11; rek-sorost@medisin.uio.no), ref:134663

### Study design

Single-blinded randomized controlled trial with parallel groups and repeated measurements

### **Primary study design**

Interventional

### **Secondary study design**

Randomised controlled trial

### **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**

Wellbeing

### **Interventions**

Treatment conditions:

The participants will undergo a 10-minutes self-paced walk on a manually driven treadmill whilst being exposed to either a 360° video or a 3D-model representing a natural environment. The exposure to the treatments and control conditions will be preceded by a preliminary elicitation of sadness using a short-film.

1. Treatment 1: Treadmill walk, in combination with exposure to a 3D model IVE
2. Treatment 2: Treadmill walk, in combination with exposure to a 360' video IVE
3. Control: Treadmill walk only with no exposure to any IVE

IVE= Immersive Virtual Environments

Measurements will be taken at three time-points:

1. Baseline, i.e. right before starting the experimentation
2. After watching the short-video (pre-exposure)
3. After exposure to the treatments or control conditions

### **Intervention Type**

Device

### **Phase**

Not Specified

### **Drug/device/biological/vaccine name(s)**

Immersive Virtual Environments

### **Primary outcome measure**

Affect (Physical activity affect scale, PAAS) measured at pre-exposure and post-exposure

### **Secondary outcome measures**

1. LTEQ: Leisure time exercise questionnaire measured at baseline
2. GE-INT: Future green exercise intention measured at baseline and post-exposure
3. N-connect: Connectedness with nature scale measured at baseline and post-exposure
4. PAAS: Physical activity affect scale measured at baseline, pre-exposure and post-exposure
5. Blood pressure (BP, measured using a semi-automatic sphygmomanometer) for 5 min at baseline, pre-exposure and post-exposure
6. Enjoyment (Single-item scale) measured at post-exposure
7. Perceived exertion (Ratings of perceived exertion, RPE) measured at post-exposure
8. Cyber sickness (Simulator-sickness questionnaire) measured at post-exposure
9. Presence (Presence see scale) measured at post-exposure
10. Perceived environmental restorativeness (Perceived restorativeness scale, PRS) measured at post-exposure
11. Walking speed (measured by the treadmill) for a 10 min period during exposure
12. Heart rate (continuous monitoring during walking) for a 10 min period during exposure

**Overall study start date**

01/04/2020

**Completion date**

04/12/2020

## Eligibility

**Key inclusion criteria**

1. Age >18 years old
2. Normal or corrected to normal vision
3. Limited experience with VR (less than monthly use)
4. No previous diagnosis of balance impairments

**Participant type(s)**

Healthy volunteer

**Age group**

Adult

**Lower age limit**

18 Years

**Sex**

Both

**Target number of participants**

60 participants

**Total final enrolment**

60

**Key exclusion criteria**

Limited understanding of Norwegian language

**Date of first enrolment**

01/09/2020

**Date of final enrolment**

30/12/2020

## Locations

**Countries of recruitment**

Norway

**Study participating centre**

**Inland Norway University of Applied Sciences**

Postboks 400

2418 Elverum

Norway

Elverum

Norway

2418

## Sponsor information

**Organisation**

Inland Norway University of Applied Sciences

**Sponsor details**

Faculty of Social and Health Sciences

INN University

Postboks 400

2418 Elverum

Norway

Elverum

Norway

2418

+47 62 43 00 00

postmottak@inn.no

**Sponsor type**

University/education

**Website**

<https://eng.inn.no>

**ROR**

<https://ror.org/02dx4dc92>

# Funder(s)

## Funder type

Government

## Funder Name

Regional Research Fund Inland Norway (RFF - innlandet) - Project code: 292761

# Results and Publications

## Publication and dissemination plan

Current publication and dissemination plan as of 17/12/2021:

The plan is to publish two articles. One for H1-H3 (study 1) and one for H4 with focus on Connectedness with Nature (study 2). A third study with focus on Future green exercise intention (H4) will be written as a "multi-study" article along with data from previous studies. Findings relative to H5 will not be published in journal articles, but presented at conferences and internal meetings and used as an explorative foundation for future studies. Preliminary results will be presented at various research conferences, at national and international level.

Previous publication and dissemination plan:

The plan is to publish two articles. One for H1-H3 (study 1) and one for H4-H5 (study 2). Study 2 will be written as a "multi-study" article along with data from previous studies. Preliminary results will be presented at various research conferences, at national and international level.

## Intention to publish date

30/06/2022

## Individual participant data (IPD) sharing plan

The datasets generated during and/or analysed during the current study are not expected to be made available due to consent not being given. Sample-level data (e.g. means and standard deviations) will be made available upon request from Giovanna Calogiuri (giovanna.calogiuri@usn.no) or Sigbjørn Litleskare (sigbjorn.litleskare@inn.no)

## IPD sharing plan summary

Available on request

## Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Basic results</a>		26/01/2022	26/01/2022	No	No
<a href="#">Results article</a>		06/07/2022	06/09/2022	Yes	No
<a href="#">Results article</a>		05/04/2022	06/09/2022	Yes	No