The health promoting potential of nature experiences in virtual reality

Submission date	Recruitment status	[X] Prospectively registered		
11/08/2020	No longer recruiting	Protocol		
Registration date	Overall study status	Statistical analysis plan		
28/08/2020	Completed	[X] Results		
Last Edited	Condition category	[] Individual participant data		
06/09/2022	Other			

Plain English summary of protocol

Background and study aims.

Green exercise (any physical activity in the presence of nature) is proposed as a key costeffective 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

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

Sponsor type

University/education

postmottak@inn.no

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 asof 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 Natute (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?
Basic results		26/01/2022	26/01/2022	No	No
Results article		06/07/2022	06/09/2022	Yes	No
Results article		05/04/2022	06/09/2022	Yes	No