

# Virtual reality mediated deep breathing and relaxation training in pediatric and psychiatric care

<b>Submission date</b>	<b>Recruitment status</b>	<input type="checkbox"/> Prospectively registered
13/05/2025	No longer recruiting	<input checked="" type="checkbox"/> Protocol
<b>Registration date</b>	<b>Overall study status</b>	<input type="checkbox"/> Statistical analysis plan
23/05/2025	Completed	<input type="checkbox"/> Results
<b>Last Edited</b>	<b>Condition category</b>	<input type="checkbox"/> Individual participant data
14/10/2025	Mental and Behavioural Disorders	<input checked="" type="checkbox"/> Record updated in last year

## Plain English summary of protocol

### Background and study aims

Virtual reality techniques have been used for various purposes in healthcare to find novel ways to treat different medical conditions and to prepare patients for unpleasant procedures. The study aims to create virtual reality applications for children for learning deep breathing and relaxation, and to study their effectiveness, safety and usability in different child patient groups. The main goal is to learn if virtual reality-mediated deep breathing and relaxation training can reduce experiences of stress and anxiety via autonomic nervous system stimulation.

### Who can participate?

The participants are child patients aged 8-17 years.

### What does the study involve?

The participants perform either guided deep breathing or relaxation exercises using a virtual reality headset. The exercise is guided by a avatar character providing verbal instructions in a virtual nature environment.

### What are the possible benefits and risks of participating?

The study patients receive the benefit of a learning experience of the guided deep breathing or relaxation exercise and the possible stress and anxiety reducing effects of these exercises. The possible risk is having nausea or vertigo, which are occasionally reported during the use of a virtual reality headset.

### Where is the study run from?

Tampere University Hospital (Finland)

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

May 2022 to June 2024

### Who is funding the study?

1. Tampere University (Finland)

2. The Finnish Foundation for Pediatric Research (Finland)
3. Finnish Brain Foundation (Finland)
4. Päivikki and Sakari Sohlberg Foundation (Finland)

Who is the main contact?  
MD, PhD Sauli Palmu, [sauli.palmu@tuni.fi](mailto:sauli.palmu@tuni.fi)

## Contact information

### Type(s)

Public, Scientific

### Contact name

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Principal investigator

### Contact name

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## Additional identifiers

### Clinical Trials Information System (CTIS)

Nil known

**ClinicalTrials.gov (NCT)**

Nil known

**Protocol serial number**

Nil known

## Study information

**Scientific Title**

The effect of virtual reality mediated deep breathing and relaxation training on stress and anxiety in child patients: a pilot trial

**Acronym**

VirNE

**Study objectives**

Deep breathing and relaxation training reduce stress and anxiety in child patients.

**Ethics approval required**

Ethics approval required

**Ethics approval(s)**

approved 15/06/2021, The Ethics Committee of the Wellbeing Services County of Pirkanmaa (PL 2000, Tampere, 33521, Finland; +358 50 347 0251, +358 50 329 5667; toimikunta.eettinen@pirha.fi), ref: RL21070L

**Study design**

Multicenter interventional non-randomized controlled pilot trial

**Primary study design**

Interventional

**Study type(s)**

Treatment

**Health condition(s) or problem(s) studied**

Reducing anxiety and stress in child patients

**Interventions**

Current interventions as of 14/10/2025:

The patients are allocated to either the intervention or the control group in numerical order. The intervention is virtual reality-mediated deep breathing or relaxation training. The study patients perform a 6-minute deep breathing or relaxation exercise with a virtual reality headset in the pediatric unit, observing a 360-degree virtual Finnish nature environment and an avatar character providing the patient with verbal guidance throughout the exercise. The study patients will perform the exercise once (in a distraction group) or four times (in a training group; one session per week in a 4-week period). During the exercise, heart rate variability is measured using a heart rate sensor belt (a chest strap). The control condition is treatment as usual.

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## Previous interventions:

The patients are allocated to either the intervention or the control group in numerical order. The intervention is virtual reality-mediated deep breathing or relaxation training. The study patients perform a 6-minute deep breathing or relaxation exercise with a virtual reality headset in the pediatric unit, observing a 360-degree virtual Finnish nature environment and a bot figure providing the patient with verbal guidance throughout the exercise. The study patients will perform the exercise once (in a distraction group) or four times (in a training group; one session per week in a 4-week period). During the exercise, heart rate variability is measured using a heart rate sensor belt (a chest strap). The control intervention is no treatment.

### **Intervention Type**

Device

### **Phase**

Not Specified

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

Oculus Quest 2 - virtual reality headset

### **Primary outcome(s)**

Heart rate variability measured using the virtual reality headset during the research situation

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

Heart rate and user feedback measured using the virtual reality headset during the research situation

### **Completion date**

30/06/2024

## **Eligibility**

### **Key inclusion criteria**

1. Child patients aged 8-17 years
2. Literate
3. Fluent in Finnish

### **Participant type(s)**

Patient

### **Healthy volunteers allowed**

No

### **Age group**

Child

### **Lower age limit**

8 years

**Upper age limit**

17 years

**Sex**

All

### **Key exclusion criteria**

1. Unstable heart condition
2. Anatomical facial features preventing the use of the headset
3. Hearing impairment or cochlear implant
4. Vision impairment
5. Tendency for vertigo
6. Clinically unevaluated seizures
7. Epilepsy
8. Migraine
9. Bulimia nervosa
10. Relative exclusion criteria: motion sickness, underweight

**Date of first enrolment**

04/05/2022

**Date of final enrolment**

30/06/2024

## **Locations**

**Countries of recruitment**

Finland

**Study participating centre**

**Tampere University Hospital**

Elämänaukio 2

Tampere

Finland

33520

**Study participating centre**

**Päijät-Häme Central Hospital**

Keskussairaalankatu 7

Lahti

Finland

15850

## **Sponsor information**

**Organisation**  
Tampere University

**ROR**  
<https://ror.org/033003e23>

## Funder(s)

**Funder type**  
University/education

**Funder Name**  
Tampere University

**Funder Name**  
Finnish Foundation for Pediatric Research

**Funder Name**  
Finnish Brain Foundation

**Funder Name**  
Päivikki and Sakari Sohlberg Foundation

## Results and Publications

### Individual participant data (IPD) sharing plan

The datasets generated and/or analyzed during the current study are not publicly available due ethical and data privacy issues in healthcare, but are partly available from Elina Karppa (elina.karppa@tuni.fi) upon reasonable request. Consent from the participants was required and obtained; data is anonymised with ID codes.

**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="#">Participant information sheet</a>	Participant information sheet	11/11/2025	11/11/2025	No	Yes

[Protocol file](#)

version 3

14/05/2025 23/05/2025 No

No