

# Audio unconscious persuasion for weight loss in overweight adults

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<b>Registration date</b> 27/02/2020	<b>Overall study status</b> Completed	<input type="checkbox"/> Protocol
<b>Last Edited</b> 23/09/2021	<b>Condition category</b> Nutritional, Metabolic, Endocrine	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data

## Plain English summary of protocol

Background and study aims

Weight loss interventions using audio unconscious persuasion are increasingly popular. However, no robust trials have been conducted to assess their effectiveness.

The aim of this trial is to assess the effectiveness of an audio unconscious persuasion weight loss intervention incorporating techniques from NLP and hypnosis (Slimpod®) compared to audio relaxation (control) on weight loss and other measures.

Who can participate?

Healthy adult volunteers with BMI > 25 kg/m<sup>2</sup>

What does the study involve?

Participants will be randomly allocated to the intervention or control groups.

The intervention is Slimpod®, a nine-minute long digital audio recording focused on changing behaviours and attitudes related to diet and weight. Participants listen to the recording once a day for 24 weeks at a time and place of their choosing. Recordings are available in two formats: digital file (.MP3) and compact disc (CD); participants can select their preferred format.

The control group are asked to listen to nine-minute long recording informed by the principles of relaxation therapy once a day for 24 weeks. The control recording does not include a focus on behaviour change.

What are the possible benefits and risks of participating?

Benefits of participating: The potential to lose weight with the need for willpower or conscious effort such as conscious calorie restriction. There is no risk in participating.

Where is the study run from?

City University London (UK)

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

October 2013 to July 2014

Who is funding the study?

Investigator initiated and funded

Who is the main contact?  
Christopher Roycroft-Davis  
chris@thinkingslimmer.com

## Contact information

**Type(s)**  
Scientific

**Contact name**  
Mr Christopher Roycroft-Davis

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## Additional identifiers

**Protocol serial number**  
PSYETH (UPTD) 12/13 71

## Study information

**Scientific Title**  
Revolutionising participants' health and wellbeing through Neuro-reprogramming via Slimpod® app: a RCT

**Study objectives**  
The aim of this pilot randomised controlled trial is to assess the effectiveness of an audio unconscious persuasion weight loss intervention incorporating techniques from NLP and hypnosis (Slimpod®) compared to audio relaxation (control) on weight (primary outcome), eating self-efficacy, exercise confidence, and quality of life (secondary outcomes).

**Ethics approval required**  
Old ethics approval format

**Ethics approval(s)**  
Approved 09/10/2013, Research and ethics committee of the department of psychology of City University London (Jay Leighton, Executive Assistant, School of Arts & Social Sciences, D217, Rhind Building, City, University of London, Northampton Square, London, EC1V 0HB, UK; +44 (0) 20 7040 3362; jay.leighton@city.ac.uk), ref: PSYETH (UPTD) 12/13 71

## **Study design**

Pilot randomized controlled trial

## **Primary study design**

Interventional

## **Study type(s)**

Other

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

Obesity

## **Interventions**

The intervention is Slimpod®, a nine-minute long digital audio recording focused on changing behaviours and attitudes related to diet and weight. Participants listen to the recording once a day for 24 weeks at a time and place of their choosing. Recordings are available in two formats: digital file (.MP3) and compact disc (CD); participants can select their preferred format. The control group are asked to listen to nine-minute long recording informed by the principles of relaxation therapy once a day for 24 weeks. The control recording does not include a focus on behaviour change.

To assess the effectiveness of an audio unconscious persuasion weight loss intervention called Slimpod® compared to audio relaxation (control), overweight adults (i.e., age < 18 years, self-reported Body Mass Index < 25.0 kg/m<sup>2</sup>) will be randomized to intervention and control groups. Weight (primary outcome) will be assessed at trial commencement, mid-trial (12 weeks) and trial end (24 weeks). Secondary outcomes are assessed using the Eating Self-Efficacy Scale (ESES), Exercise Confidence Scale (ECS) and Quality of Life Index Generic Version III (QLI-G3) at the start and end of the trial. The nine-minute Slimpod audio recording and the nine-minute control audio are provided to participants as MP3 files to be listened to daily every day for 24 weeks.

Simple randomization is used to allocate participants to the intervention or control groups using a 1:1 allocation ratio. A computer programme operated by the study researcher is used to generate a random number to assign each participant to a trial arm. The trial statistician is not involved in the process of recruitment, randomization, or group assignment. Blinding It is not possible to blind participants to their group allocation. Participants became aware of their allocation on first listening to the audio recording they had received as the recording for those in the intervention group included weight loss messages, whereas the control group recording did not.

## **Statistical methods**

Data analysis is conducted using R version 3.4.3. Baseline characteristics are reported as mean and standard deviation (SD) for continuous data and n (%) for categorical data. For each measure, baseline is defined as the value collected prior to the commencement of randomised therapy. The two therapy arms are compared using analysis of covariance (ANCOVA) using the change from baseline at each post-baseline assessment separately, with the baseline fitted as a covariate; interaction between baseline and treatment are assessed but removed from the model as not statistically significant.

## **Intervention Type**

Behavioural

**Primary outcome(s)**

Weight (kg) at baseline, 12, and 24-weeks

**Key secondary outcome(s)**

At baseline and 24-weeks:

1. Eating Self-Efficacy Scale (ESES)
2. Exercise Confidence Scale (ECS)
3. Quality of Life Index Generic Version III (QLI-G3)

**Completion date**

01/07/2014

## Eligibility

**Key inclusion criteria**

1. BMI above 25 kg/m<sup>2</sup>
2. Aged 18 years or above

**Participant type(s)**

Healthy volunteer

**Healthy volunteers allowed**

No

**Age group**

Adult

**Lower age limit**

18 years

**Sex**

All

**Total final enrolment**

82

**Key exclusion criteria**

1. Pregnant
2. Diagnosed with mental health condition
3. Seeking medical support for a medical diagnosis unless prior consent was obtained from their medical practitioner

**Date of first enrolment**

05/10/2013

**Date of final enrolment**

01/01/2014

## Locations

## Countries of recruitment

United Kingdom

England

## Study participating centre

City University London

Northampton Square

Clerkenwell

London

United Kingdom

EC1V 0HB

## Sponsor information

### Organisation

ThinkingSlimmer Ltd

## Funder(s)

### Funder type

Other

### Funder Name

Investigator initiated and funded

## Results and Publications

### Individual participant data (IPD) sharing plan

The datasets generated during and/or analysed during the current study are available from the corresponding author on reasonable request, data will be available from April 2, 2020 indefinitely.

### IPD sharing plan summary

Available on request

### Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Abstract results</a>		18/06/2020	23/09/2021	No	No