







# Rethinking approaches to excess weight in adolescents (RENEWAL)

<b>Submission date</b> 10/02/2023	<b>Recruitment status</b> Recruiting	 Retrospectively registered
<b>Registration date</b> 28/02/2023	<b>Overall study status</b> Ongoing	 Protocol not yet added
<b>Last Edited</b> 25/04/2024	<b>Condition category</b> Nutritional, Metabolic, Endocrine	 SAP not yet added
		 Results not yet expected
		 Raw data not yet expected
		 Record updated in last year

## Plain English Summary

### Background and study aims

Approximately one in three adolescents aged 13 to 17 years is above a healthy weight. Excess weight in adolescents can lead to both physical and mental health concerns and often persists into adulthood. Behavioural weight management interventions are currently regarded as best practice in obesity treatment for children and adolescents. Unfortunately success on these programmes is limited and often non-existent when delivered at scale or over the long term. Therefore a new and novel approach is needed to treat excess weight in adolescents. Digital interventions are interventions delivered virtually either through a website, computer programme, virtual coach or mobile app. Studies in other fields have shown promise results when using digital interventions for work with adolescents and preliminary studies using digital interventions for weight loss also show promise. Despite this there are no studies based in the United Kingdom showing the use of a digital intervention for weight management in adolescents.

Our study aims to test whether it is possible to use a mobile app for treating excess weight in overweight and obese adolescents. We will monitor the impact of the intervention on weight, nutrition, physical activity and self-esteem as well as monitoring engagement levels and satisfaction. If successful, we will progress to a bigger study to investigate whether this intervention can support adolescents to reach a healthy weight.

### Who can participate?

Any young person between the ages of 13 and 17 years who is deemed above a healthy weight by a health care professional. Participants must have access to a smart phone or tablet and must not have any other significant physical or mental illnesses.

### What does the study involve?

Young people will be randomised to either the intervention or the control groups (2:1). The control group will receive a 1:1 with a dietitian where they will review current behaviours, set smart goals and be signposted to further resource. The intervention group will gain access to the Second Nature app and be encouraged to use it for 12 weeks with dietitian check-ins at weeks 3 and 7. Both groups will undergo pre and post-intervention assessment and post-intervention progress, or lack thereof, will be reported to the referring HCP.

What are the possible benefits and risks of participating?

Whilst participants may lose weight and increase their health by taking part in this study, we do not know what the outcome will be which is why we are conducting this research. Participants will learn more about healthy eating, exercise and sleep for weight loss and be able to discuss their needs with a dietitian. It is hoped that the results of this study will support young people to better manage their weight in the future. This is a low-risk public health intervention therefore the disadvantages and risks to taking part are minimal. However all participants will be monitored for signs of disordered eating and mental wellbeing and will be referred back to local health services should the need arise.

Where is the study run from?

University of Oxford (UK)

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

January 2022 to July 2025

Who is funding the study?

National Institute for Health and Care Research Applied Research Collaboration Oxford and Thames Valley (UK)

Who is the main contact?

Melissa Little, [melissa.little@phc.ox.ac.uk](mailto:melissa.little@phc.ox.ac.uk)

## Contact information

**Type(s)**

Scientific

**Contact name**

Ms Melissa Little

**ORCID ID**

<http://orcid.org/0000-0001-5619-2679>

**Contact details**

University of Oxford  
Nuffield Department of Primary Care  
Radcliffe Observatory Quarter  
Woodstock Road  
Oxford  
United Kingdom  
OX2 6GG  
+44 7557 360624  
[melissa.little@phc.ox.ac.uk](mailto:melissa.little@phc.ox.ac.uk)

## Additional identifiers

**EudraCT/CTIS number**

Nil known

**IRAS number**

316058

**ClinicalTrials.gov number**

Nil known

**Protocol/serial number**

CPMS 54848, IRAS 316058

## Study information

**Scientific Title**

REthiNking Approaches to Excess Weight in AdoLescents (RENEWAL): a randomised feasibility trial to assess a digital Intervention for managing excess weight in adolescents with overweight and obesity

**Acronym**

RENEWAL

**Study hypothesis**

When compared to a brief weight management intervention, the digital intervention will promote a reduction in BMI Z-score, among adolescents with excess weight.

**Ethics approval required**

Old ethics approval format

**Ethics approval(s)**

Approved 22/11/2022, Health Research Authority and Health and Care Research Wales (Skipton House, 80 London Road, London, SE1 6LH, UK; +44 (0)20 7972 2545; hra.approval@nhs.net), ref: 22/NS/0143

**Study design**

Interventional randomized controlled trial

**Primary study design**

Interventional

**Secondary study design**

Randomised controlled trial

**Study setting(s)**

Home, Internet/virtual

**Study type(s)**

Treatment

**Participant information sheet**

See additional files

**Condition**

## Obesity

### Interventions

**Summary:** this is an individually randomised controlled feasibility trial. Participants, recruited via hospitals, local authorities and NHS trusts, will be randomised 2:1 to intervention group or control group. The control group will receive a brief intervention for weight management and the intervention group will receive the use of the digital intervention (app) for 12 weeks. Most study procedures will be conducted online or remotely (by telephone) including: eligibility assessment, informed consent, baseline assessment, randomisation, engagement with the app-based intervention (or brief intervention for the control group) and 2 dietitian check-in appointments. The final appointment, a post-intervention assessment, will be done by the research team in person at a mutually acceptable location.

**Baseline Assessment:** Once consent has been obtained a member of the research team will support the participant virtually to complete the baseline assessments. These will include three questionnaires (Adult Eating Behaviour Questionnaire, Adolescent Food Habits Checklist, Rosenberg Self-Esteem Questionnaire) as well as obtaining basic demographic data. Participants will only be randomised once they have fully completed the questionnaires.

**Randomisation:** Once consent and baseline assessments have been completed, all eligible participants will be individually randomised to one of two arms in a 2:1 ratio: Intervention (12-week, app-based support), or control (brief intervention on weight management). The randomisation will be conducted using an online randomisation programme such as OxMar trying to balance gender and deprivation level across the two groups. Due to the nature of the intervention, it will not be possible to blind participants or the research team.

**Intervention group:** Participants randomised to the intervention group will receive an email explaining how to download and access the app-based intervention, and a voucher code that will provide access to the programme for 12 weeks. The intervention, provided by Second Nature, involves diet, activity and behaviour change components. It comprises a 3-month remote behavioural change programme with mentoring from a registered dietitian or nutritionist (health coach), peer group support, structured education articles and activity tracking technology. These elements are accessed via a smartphone or web-based application. Participants will also be asked to complete a virtual 'check-in' with the dietitian at 3 and 7 weeks. The purpose of these sessions will be to review the participants' progress, provide support, answer any questions regarding the intervention and troubleshoot any issues.

**Control Group:** Participants randomised to the control group will be informed of their weight status and provided with brief advice. Participants and the dietitian will work together to suggest healthy behaviour changes and set some SMART goals to work on including signposting to online resources such as the NHS Healthier Families website and Change4Life resources. The control group will not be contacted again until the end of the 12-week trial period.

**Post-intervention Assessment:** All participants will be asked to attend a post-intervention assessment appointment. The appointment will be done in-person at a mutually acceptable location where weight and height will be recorded by the research team. Participants will be asked to complete the same questionnaires that they completed at the baseline assessment.

**Interviews:** After the 12-week post intervention visit the research team will conduct qualitative interviews with all participants in the intervention group to learn more about their views on the

intervention. Interviews will last approximately 30 minutes. With participants consent this will be audio-recorded. The interview will be conducted by a trained member of the study team and will be audio-recorded and then transcribed.

## **Intervention Type**

Behavioural

## **Primary outcome measure**

1. Engagement (user contact with the intervention) measured using amalgamated number of contact points with the app throughout the Intervention.
2. Retention (user completion of the intervention) measured using proportion of people attending their 12 week post-intervention appointment at the end of the intervention.
3. Recruitment (length of time to complete recruitment for the study) measured using Average number of people recruited to the intervention per month during the recruitment phase.

## **Secondary outcome measures**

Measurements at baseline and follow-up:

1. BMI Z-score measured using mean difference in BMI z-score between follow-up and baseline.
2. Eating behaviours measured using mean difference in score on the Adult Eating Behaviours questionnaire between follow-up and baseline and mean difference in score on the Adolescent Food Habits Checklist between follow-up and baseline.
3. Physical activity levels measured using mean difference in steps between follow-up and baseline and number of self-record exercises done through app.
4. Self-Esteem measured using mean difference in score on the Rosenberg Self Esteem Scale between follow-up and baseline.

## **Overall study start date**

15/01/2022

## **Overall study end date**

01/07/2025

# **Eligibility**

## **Participant inclusion criteria**

1. Young people aged 13-17 years living in the United Kingdom
2. BMI z-score >91st centile for BMI using the UK 1990 growth reference.
3. Participant has access to an appropriate device to install and use the app.

## **Participant type(s)**

Healthy volunteer

## **Age group**

Child

## **Lower age limit**

13 Years

## **Upper age limit**

17 Years

**Sex**

Both

**Target number of participants**

Planned Sample Size: 30; UK Sample Size: 30

**Participant exclusion criteria**

1. Participants with a mental illness or significant mental health problems that would interfere with adherence to the intervention or trial protocol.
2. Participants with a learning delay that would interfere with adherence to the intervention or trial protocol.
3. Participants with any other illness that would interfere with adherence to the intervention or trial protocol.

**Recruitment start date**

01/02/2023

**Recruitment end date**

01/12/2024

**Locations****Countries of recruitment**

England

United Kingdom

**Study participating centre**

**NIHR CRN: Thames Valley and South Midlands**

Nuffield Department of Primary Care

Oxford

United Kingdom

OX3 9DU

**Sponsor information****Organisation**

University of Oxford

**Sponsor details**

Joint Research Office 1st floor

Boundary Brook House

Churchill Drive

Headington

Oxford

England  
United Kingdom  
OX3 7GB  
-  
ctrng@admin.ox.ac.uk

**Sponsor type**  
University/education

## **Funder(s)**

**Funder type**  
Government

**Funder Name**  
National Institute for Health and Care Research Applied Research Collaboration Oxford and Thames Valley

**Alternative Name(s)**  
NIHR ARC Oxford and Thames Valley, NIHR Applied Research Collaboration Oxford and Thames Valley, Oxford and Thames Valley NIHR Applied Research Collaboration, NIHR Oxford and Thames Valley Applied Research Collaborative, National Institute for Health and Care Research (NIHR) Oxford and Thames Valley Applied Research Collaboration, NIHR Applied Research Collaboration (ARC) for Oxford and the Thames Valley, ARC OTV, OTV ARC, NIHR ARC OTV, NIHR ARC-OxTV, ARC OxTV

**Funding Body Type**  
Government organisation

**Funding Body Subtype**  
National government

**Location**  
United Kingdom

## **Results and Publications**

**Publication and dissemination plan**  
Planned publication in a high-impact peer-reviewed journal

**Intention to publish date**  
01/04/2025

**Individual participant data (IPD) sharing plan**  
The current data sharing plans for this study are unknown and will be available at a later date

## IPD sharing plan summary

Data sharing statement to be made available at a later date

### Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Participant information sheet</a>	Competent youth version 1.1	15/11/2022	24/02/2023	No	Yes
<a href="#">Participant information sheet</a>	Parent version 1.1	15/11/2022	24/02/2023	No	Yes
<a href="#">Participant information sheet</a>	Young person version 1.1	15/11/2022	24/02/2023	No	Yes
<a href="#">HRA research summary</a>			28/06/2023	No	No