

# Feasibility of providing digital physical activity promotion materials for young people with cystic fibrosis

<b>Submission date</b>	<b>Recruitment status</b>	<input type="checkbox"/> Prospectively registered
06/10/2022	Stopped	<input type="checkbox"/> Protocol
<b>Registration date</b>	<b>Overall study status</b>	<input type="checkbox"/> Statistical analysis plan
07/11/2022	Stopped	<input type="checkbox"/> Results
<b>Last Edited</b>	<b>Condition category</b>	<input type="checkbox"/> Individual participant data
03/05/2023	Respiratory	<input type="checkbox"/> Record updated in last year

## Plain English summary of protocol

### Background and study aims

This is a feasibility trial of a digital physical activity intervention for young people living with cystic fibrosis (CF). CF is the most common inherited condition in the UK and is caused by a faulty gene. Physical activity is recommended in managing CF to counter the physical and psychological effects of the condition. Despite the benefits of regular physical activity, it is often reported that engagement in physical activity is poor. Physical activity levels among those with CF decrease during adolescence and remain low during adulthood, which can have long-term health consequences for people with CF impacting their quality of life. Intervening during adolescence is therefore critical.

We have developed a digital physical activity intervention that considers unique barriers experienced by young people with CF to motivate and support engagement in physical activity and to increase their quality of life. The aim of the current study is to help establish whether digitally delivering (e.g., online) physical activity promotion material is accessible, usable, and engaging in supporting young people with CF to be more physically active.

### Who can participate?

Young people aged between 12 and 18 years old with CF

### What does the study involve?

The total time commitment for participation is 6 months. Access to digitally delivered physical activity promotion material will be provided throughout this period. The trial will involve completing questionnaires, taking part in interviews (optional), and wearing activity monitors at different times over the course of the trial.

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

Access to digital physical activity promotion material may lead to participants increasing their levels of physical activity. The type of activity undertaken is of the participant's own choosing (i.

e., non-prescriptive). While increased levels of physical activity may lead to several health benefits, one issue is that it may also lead to increased injury risk. This is viewed as a minor risk of participating.

Where is the study run from?

This trial is being sponsored by the Royal Devon & Exeter NHS Foundation Trust (UK) and will be delivered by researchers based at the University of Exeter (UK) and the University of Bristol (UK)

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

July 2020 to August 2023

Who is funding the study?

National Institute of Health and Care Research (NIHR) Research for Patient Benefit (RFpB) (UK)

Who is the main contact?

Dr Robert Mann

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## Contact information

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## Additional identifiers

#### Clinical Trials Information System (CTIS)

Nil known

#### Integrated Research Application System (IRAS)

308638

#### ClinicalTrials.gov (NCT)

Nil known

#### Protocol serial number

CPMS 53342, IRAS 308638

## Study information

**Scientific Title**

Feasibility randomised controlled trial of digital physical activity promotion materials for young people with Cystic Fibrosis

**Acronym**

Digi PA for Youth CF – Feasibility

**Study objectives**

We expect to learn about:

1. The feasibility of conducting a full-scale randomised controlled trial
2. The feasibility and acceptability of digital physical activity promotion materials
3. Potential mechanisms of action
4. Moderators to be investigated as part of a (future) full-scale randomised controlled trial

**Ethics approval required**

Old ethics approval format

**Ethics approval(s)**

Approved 15/09/2022, West Midlands - Solihull Research Ethics Committee (Meeting held by video-conference via Zoom; +44 (0)207 104 8191, (0)207 104 8269; solihull.rec@hra.nhs.uk), ref: 22/WM/0156

**Study design**

Feasibility randomized controlled trial

**Primary study design**

Interventional

**Study type(s)**

Treatment

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

Respiratory

**Interventions**

The study design is a two-arm feasibility randomised controlled trial with nested mixed-methods process evaluation. Participants will be randomly allocated in a 1:1 ratio to receive one of two types of digital physical activity promotion materials. Both groups will also receive usual care as per clinical practice guidelines for the treatment of patients with CF. Data is collected at baseline, 1-month, and 6-month follow-up. All data collection and access to digital PA materials will take place online.

Having enrolled in the trial (i.e., completed the screening process and returned consent/assent forms), all participants will be asked to provide data at three time points (baseline, 1-month follow-up, and 6-month follow-up). All data collection will take place remotely. The assessments that participants will complete at each timepoint are as follows:

**BASELINE:**

All participants will need to complete a set of questionnaires related to: (1) health and demographic information; (2) health-related quality of life; (3) physical activity behaviour; and (4) health and social care resource use and economic impact. All participants will also be required to wear an activity monitor for a period of 7-days, alongside completing a short log about their

activity and a physical activity questionnaire.

#### 1-MONTH FOLLOW-UP

All participants will need to complete a set of questionnaires related to: (1) health-related quality of life and (2) physical activity behaviour. All participants will also be required to wear an activity monitor for a period of 7-days, alongside completing a short log about their activity and a physical activity questionnaire. All participants in the intervention group will also need to complete a questionnaire about the usability and acceptability of the intervention. A sub-sample of participants will also be invited to take part in a semi-structured interview.

#### 6-MONTHS FOLLOW-UP

All participants will need to complete a set of questionnaires related to: (1) health-related quality of life; (2) physical activity behaviour; and (3) health and social care resource use and economic impact. All participants will also be required to wear an activity monitor for a period of 7-days, alongside completing a short log about their activity and a physical activity questionnaire. All participants will also need to complete a questionnaire about their trial satisfaction. A sub-sample of participants will also be invited to take part in a semi-structured interview.

In addition to the data provided by participants, data related to trial feasibility (e.g., participant enrolment, willingness to be randomised, participant retention) and usual care (e.g., understanding level of physical activity advice provided by participant's usual CF care team) will also be collected. Trial feasibility data will also be recorded as part of trial procedures. Usual care data will be reported by a relevant member of each participant's CF clinical care team.

In combination, these data will allow for the completion of a nested mixed-methods process evaluation and economic evaluation. The process evaluation will focus on assessing the feasibility and acceptability of the digital physical activity promotion materials. The economic evaluation will focus on establishing: (1) the digital physical activity promotion material delivery costs; (2) the most appropriate way to collect health economics data in a full-scale trial; and (3) to set out an appropriate framework for health economic evaluation as part of a full-scale trial.

### **Intervention Type**

Behavioural

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

Feasibility of enrolling participants measured using patient logs throughout the trial:

1. The average number of participants enrolled into the trial per month, including details of their recruitment pathway/route
2. The eligible population, as indicated by the number of eligible participants invited
3. Recruitment (where possible), indicated by the number of participants who submit an Expression of Interest
4. Enrolment/uptake, indicated by the number of participants enrolled

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

1. Retention rates measured using a Trial Process Log throughout the 6-month trial recruitment and follow-up data collection periods
2. Trial assessment completion rates measured as part of the Trial Process Log for each participant at each of the trial data collection timepoints
3. Drop-out rates measured using a Trial Process Log throughout the 6-month trial recruitment and follow-up data collection periods
4. Usage of digital physical activity promotion materials measured using online metrics (e.g., total time spent using digital physical activity materials) throughout the 6-month follow-up data collection period
5. Feasibility of use and trial satisfaction measured using the User Version of the Mobile

Application Rating Scale (uMARS) questionnaire after 1-month follow-up and qualitative interviews after 1-month and 6-months of follow-up in a sub-sample of participants. Trial satisfaction will also be recorded via the Trial Satisfaction Questionnaire (at 6-month follow-up) or Trial Withdrawal Questionnaire (at the point of trial withdrawal).

6. Describing usual care in relation to physical activity will be measured by the completion of an online form by each participant's CF Care Team at the baseline data collection timepoint

### **Completion date**

24/08/2023

### **Reason abandoned (if study stopped)**

Participant recruitment issue

## **Eligibility**

### **Key inclusion criteria**

1. Confirmed diagnosis of cystic fibrosis
2. Aged between 12 and 18 years old
3. Able to access the internet via any electronic device

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

Patient

### **Healthy volunteers allowed**

No

### **Age group**

Child

### **Lower age limit**

12 years

### **Upper age limit**

18 years

### **Sex**

All

### **Total final enrolment**

0

### **Key exclusion criteria**

1. Presence of an existing severe co-morbidity limiting mobilisation and/or participation in physical activity (e.g., orthopaedic, cardiac, or neurological condition)
2. Current (i.e., within the last month) lung function (forced expiratory volume in one second) is below 40% predicted
3. Currently pregnant
4. Inability to understand verbal and/or written English
5. Participation in a concurrent study for which eligibility precludes them from engaging in another research study and/or study focused on physical activity

6. Participation in a Patient and Public Involvement Advisory Group and/or research study that has directly informed the design of the present feasibility trial

**Date of first enrolment**

24/10/2022

**Date of final enrolment**

09/05/2023

## Locations

**Countries of recruitment**

United Kingdom

England

**Study participating centre**

Royal Devon and Exeter Hospital  
Barrack Road  
Exeter  
United Kingdom  
EX2 5DW

## Sponsor information

**Organisation**

Royal Devon University Healthcare NHS Foundation Trust

## Funder(s)

**Funder type**

Government

**Funder Name**

National Institute for Health and Care Research

**Alternative Name(s)**

National Institute for Health Research, NIHR Research, NIHRresearch, NIHR - National Institute for Health Research, NIHR (The National Institute for Health and Care Research), NIHR

**Funding Body Type**

Government organisation

**Funding Body Subtype**

National government

**Location**

United Kingdom

## Results and Publications

**Individual participant data (IPD) sharing plan**

Some of the data collected and created (e.g., anonymised questionnaire data) will be suitable for sharing for research and teaching. Other trial data (e.g., fidelity data) will be trial specific and not suitable for sharing. Only fully anonymised data will be shared at the end of the project.

Potential users will find out about data availability/access via our initial publications and a trial-specific University of Exeter website. Initially, data will be embargoed (i.e., not publicly accessible via the open-access data repository) until the publication of all intended academic papers/reports. Until then, the Trial Management Group will have exclusive use of the data.

Following this period, data will be made available via a suitable open-access data repository (e.g., Open Science Framework). Please note that access to this data will require the approval of the PIs in each instance and will require adherence to a strict licence that includes a confidentiality agreement.

**IPD sharing plan summary**

Other

**Study outputs**

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">HRA research summary</a>		28/06/2023	No		No
<a href="#">Participant information sheet</a>	Participant information sheet	11/11/2025	11/11/2025	No	Yes