

# A study on the impact of the effect of mobile health on airway clearance in patients with primary ciliary dyskinesia

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<b>Registration date</b> 24/10/2024	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
<b>Last Edited</b> 09/10/2024	<b>Condition category</b> Respiratory	<input type="checkbox"/> Individual participant data <input type="checkbox"/> Record updated in last year

## Plain English summary of protocol

### Background and study aims

Primary Ciliary Dyskinesia (PCD) is a rare genetic condition where the cilia, hair-like structures lining the airways, do not function properly. Normally, cilia move mucus and trapped pollutants out of the airways, but in PCD, this process is impaired, leading to mucus buildup and recurrent respiratory infections. These infections can cause irreversible lung damage and affect the ears, sinuses, and lungs. PCD is also linked to sub/infertility, congenital heart defects, and organ laterality defects. There is no bespoke treatment for PCD. Current treatments focus on preventing infections and managing symptoms through Airway Clearance Techniques (ACTs) and antibiotics. Many different types of ACTs are used by patients, most of which involve repeated breathing exercises to move mucus through the airways. Some patients with PCD will use inhalers or nebulisers (e.g., hypertonic saline) before their ACT to loosen the mucus in the airways. Upper airway hygiene involves nasal rinsing with hypertonic saline and/or topical nasal corticosteroids and antibiotics to manage sinus and nasal symptoms. Regular exercise may be also recommended to increase mucus clearance and improve factors that can affect respiratory health (such as cardiovascular fitness, weight, and lung function). Good engagement with ACTs may result in improved respiratory outcomes. However, engaging in a daily routine of ACTs is time-consuming and may be burdensome for PCD patients, leading to performing less efficient techniques or ultimately discontinuing the routine. Furthermore, some PCD patients, particularly young patients, view the performance of ACTs as a visible sign of 'being ill' and may be reluctant to carry them out systematically. This study aims to show that a mobile health app and a wrist-worn wearable device can support PCD patients in their daily ACT routines, improving their quality of life and reducing exacerbations.

### Who can participate?

Adolescents (from age 14 years old upwards) and adults diagnosed with PCD who are prescribed daily ACT routines

What does the study involve?

The study will use a mobile health app and a wrist-worn wearable device to support daily ACT routines. Engagement and wearable data will be collected to track exercise and physical activity. The impact on patients' quality of life and the occurrence of exacerbations will be assessed.

What are the possible benefits and risks of participating?

Benefits include improved engagement with ACT routines, better respiratory outcomes and quality of life, as well as reduced occurrence of exacerbations.

Risks are minimal, as the intervention involves non-invasive technology.

Where is the study run from?

Aparito Limited

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

October 2021 to May 2024

Who is funding the study?

1. Aparito Limited
2. PCD Support UK

Who is the main contact?

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## Contact information

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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**

Randomized controlled study to investigate the impact of mobile health (mobile app and wearables) on the engagement with prescribed airway clearance techniques in patients with primary ciliary dyskinesia (PCD)

### **Acronym**

PCD-ENGAGE

### **Study objectives**

It is hypothesized that the use of mobile health in the form of a bespoke patient-facing app and wearables with behaviour change features tailored to patients' preferences, may improve engagement with prescribed airway clearance daily routine in PCD patients, which may result in improved quality of life and reduced exacerbation occurrence.

### **Ethics approval required**

Ethics approval required

### **Ethics approval(s)**

approved 30/11/2022, Reading Independent Ethics Committee (93 Reading Road, Woodley, Reading, RG5 3AE, United Kingdom; +44 (0)1189691022; m.l.arnott@btinternet.com), ref: None available

## **Study design**

Fully decentralised randomized controlled trial

## **Primary study design**

Interventional

## **Study type(s)**

Other, Quality of life

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

Primary ciliary dyskinesia

## **Interventions**

The study involves a behavioural change intervention via a mobile health app and wrist-worn wearables to support engagement with airway clearance techniques (ACT).

Participants were randomised via an algorithm built into the study platform (Atom5™) following screening, e-consent and eligibility confirmation.

Control group arm: Participants used the app to self-report their engagement with ACTs and only received notifications to prompt logging of requested ACT data regularly to ensure enough engagement with the technology to collect the study data. Participants were also prompted to complete some patient-reported outcomes (PROs) at baseline, 6 weeks, and 12 weeks. These were: QOL-PCD, Patient Health Engagement scale (PHQ), Exacerbation events questionnaire and Spirometry readings. The total follow-up time of patients within the study was 12 weeks. Once the 12-week study period was finished, all participants in the control arm were invited to voluntarily cross over to the intervention group, to receive the activity tracker and were followed for up to 24 weeks (including answering the PROs at weeks 18 and 24). Participants crossing over from the control group provided consent (and assent was applicable) to continue in the extension phase.

Intervention arm group: Same as above with the addition of a consumer-grade CE-marked Garmin Vivosmart® 5 wearable activity tracker. The wearable device was synchronised to the Atom5™ app and passively collected data such as heart rate (HR), activity levels (step count), motion type and intensity, time spent in moderate-to-vigorous activity, and sleep. This data was visualised in real time on a study-dedicated clinician dashboard.

This arm also included the following behaviour change features:

- Reminders - to sync the wearable, and log physiotherapy data
- Compliance reports to notify participants of weekly ACT engagement.
- Weekly physical activity reports to notify participants based on activity data collected through the Garmin watch.
- Rewards: bronze, silver and gold badges were to be issued upon reception of ACT e-diaries for 3, 7 and 14 days in a row, respectively.

## **Intervention Type**

Behavioural

**Primary outcome(s)**

Change in engagement with prescribed airway clearance techniques (ACTs) measured using data collected in a daily self-reported ACT diary to the end of the study period (week 12)

**Key secondary outcome(s)**

Health-related quality of life (HRQoL) of people with PCD measured using the PCD-specific QoL instrument: QOL-PCD adult and adolescent (13-17) versions at baseline, weeks 6, and 12. Additionally, weeks 16 and 24 for those participating in the extension.

**Completion date**

30/05/2024

**Eligibility**

**Key inclusion criteria**

1. Adolescents (from 14 upwards) and adults diagnosed with PCD (written proof such as a letter of diagnosis from their reference specialist centre will be required, and/or genetic information, if known) and performing regular airway clearance techniques (ACTs)
2. Able and willing to provide consent or, if appropriate, participants having an acceptable individual capable of providing consent on their behalf, e.g., parent or guardian of a child under 18 years of age
3. Able to speak and understand English
4. Able and willing to engage with the technology
5. Access to an internet-connected smartphone with iOS or Android system

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Mixed

**Lower age limit**

14 years

**Upper age limit**

100 years

**Sex**

All

**Total final enrolment**

19

**Key exclusion criteria**

1. Severe/acute disease, including:
  - 1.1. Severe haemoptysis 6 months prior study start
  - 1.2. Congenital heart defects, other than dextrocardia
  - 1.3. Cardiac disease
  - 1.4. Ongoing exacerbation. Once their exacerbation is resolved, they could be considered eligible for the study.

**Date of first enrolment**

17/04/2023

**Date of final enrolment**

10/11/2023

## **Locations**

**Countries of recruitment**

United Kingdom

England

Wales

**Study participating centre**

**Aparito Limited**

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## **Sponsor information**

**Organisation**

Aparito Limited

## **Funder(s)**

**Funder type**

Industry

**Funder Name**

Aparito Limited

**Funder Name**  
PCD Support UK

## Results and Publications

### Individual participant data (IPD) sharing plan

The data sharing plans for the current study are unknown and will be made 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>	version 1.5	21/11/2023	09/10/2024	No	Yes
<a href="#">Protocol file</a>	version 4.0	21/11/2023	09/10/2024	No	No