

# An eight week pilot study to investigate the effect of oscillation device on breathlessness in patients with chronic obstructive pulmonary disease

<b>Submission date</b> 21/03/2016	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 24/05/2016	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
<b>Last Edited</b> 24/01/2019	<b>Condition category</b> Respiratory	<input type="checkbox"/> Individual participant data

## Plain English summary of protocol

### Background and study aims

Chronic obstructive pulmonary disease (COPD) is the name used to refer to a number of progressive devastating and debilitating lung diseases, which includes chronic bronchitis, emphysema and chronic obstructive airways disease. People that have COPD typically feel breathless after physical activity, have a persistent cough with phlegm and suffer frequently from chest infections. There is no cure for the condition, but making lifestyle changes (such as stopping smoking) and taking medications (inhalers and/or tablets) can alleviate symptoms. This study aims to investigate the effect of a high frequency airway oscillatory (HFAO) device on breathlessness in patients with COPD. The effects of this device will be measured using a series of questionnaires, exercise testing and measurement of respiratory muscle strength.

### Who can participate?

Adult aged at least 40 and diagnosed with COPD.

### What does the study involve?

The study involves two visits to hospital. At the first visit, the participants are assessed to see whether they are eligible to take part. They are given a number of questionnaires to complete that assess, for example, quality of life, breathlessness and anxiety and depression. They also undergo lung function tests, fitness tests and tests on respiratory muscles strength. The participants are then taught how to use the HFAO device and are asked to use it three times a day for five minutes at a time over the next eight weeks. They are contacted by telephone weekly to discuss any issues with the device and to see whether they are still using it (checking compliance). They are also asked to fill in a daily diary of their use of the device. After the eight weeks, all participants undergo the same assessments as they did at the start of the study.

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

The risks associated with the device are minimal. Benefits include being provided with the Aerosure device and manual to keep after the study has ended. Participants will also have the

opportunity to discuss their condition with a health care professional. The information collected from this study may help in to reducing breathlessness in patients with COPD in the future. Participants will be reimbursed for travel and parking costs.

Where is the study run from?

University Hospitals of Leicester (UK)

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

April 2016 to December 2016

Who is funding the study?

Actegy LTD

Who is the main contact?

Miss Enya Daynes

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## Contact information

**Type(s)**

Public

**Contact name**

Miss Enya Daynes

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**Contact details**

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## Additional identifiers

**EudraCT/CTIS number**

**IRAS number**

**ClinicalTrials.gov number**

**Secondary identifying numbers**

201676

## Study information

**Scientific Title**

An eight week pilot study to investigate the effect of High-Frequency Airway Oscillation on breathlessness in patients with Chronic Obstructive Pulmonary Disease: a single arm interventional trial

**Acronym**

HFAO in COPD- Pilot

**Study objectives**

1. To test the hypothesis that patients training with a HFAO device for 8 weeks will have reduction in dyspnoea compared to baseline.
2. To test the hypothesis that patients training with a HFAO device for 8 weeks will show improvements in exercise capacity, cough frequency and intensity and dyspnoea at rest and upon exertion.

**Ethics approval required**

Old ethics approval format

**Ethics approval(s)**

London Central Research Ethics Committee, 24/05/2016, ref: 16/LO/0924

**Study design**

Single centre single arm trial design administering medical device intervention

**Primary study design**

Interventional

**Secondary study design**

Non randomised study

**Study setting(s)**

Hospital

**Study type(s)**

Treatment

**Participant information sheet**

Not available in web format, please use contact details below to request a participant information sheet.

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

Chronic obstructive pulmonary disease (COPD)

**Interventions**

An Aerosure device will be issued to all participants in the trial. This is a high frequency airway oscillatory device to be used up to three times per day in order to gain an insight into the training effect to reduce breathlessness and improve quality of life.

The study involves two visits to hospital. The initial visit will determine eligibility and conduct baseline measurements which will include: Chronic respiratory questionnaire, Hospital anxiety and depression score, London Chest Activity of Daily living Score, Leicester Cough

Questionnaire, COPD Assessment Test, Incremental Shuttle Walk Test, Endurance Shuttle Walk Test, Spirometry and Inspiratory and expiratory max testing. Participants will be taught how to use the device and required to use it for 8 weeks in duration, three times a day for 5 minutes at a time or equivalent. Participants will be contacted by telephone weekly to discuss any issues with the device and record compliance. They will be required to retain a daily diary of usage. Participants will return after 8 weeks to repeat baseline measurements. At this stage they will have completed the trial. They will have the option to attend a dissemination event to share the results and gain feedback on the trial.

## **Intervention Type**

Device

## **Primary outcome measure**

Breathlessness at baseline and after 8 week of HFAO, measured using the Chronic respiratory Questionnaire (CRQ) Dyspnoea domain

## **Secondary outcome measures**

1. Exercise endurance at baseline and after 8 week of HFAO, measured using the incremental and endurance shuttle walk test
2. Quality of life at baseline and after 8 week of HFAO, measured using the CRQ fatigue, mastery and emotion domains, COPD Assessment Test (CAT), Leicester Cough Questionnaire, London Chest Activities of Daily Living Questionnaire
3. Breathlessness pre and post exercise testing at baseline and 8 weeks of HFAO, measured using the Borg Breathlessness scale
4. Anxiety and depression at baseline and after 8 week of HFAO, measured using the Hospital Anxiety and Depression Scale
5. Inspiratory and expiratory muscle strength at baseline and after 8 week of HFAO, measured using lung function testing
6. Patient experience of trial measured in dissemination event discussion at the end of the trial (estimated October/November 2016)
7. Adherence of device use measured using daily diaries

## **Overall study start date**

20/04/2016

## **Completion date**

31/12/2016

# **Eligibility**

## **Key inclusion criteria**

1. Participant is willing and able to give informed consent for participation in the study
2. Male or Female, aged 40 years or above
3. Confirmed diagnosed of COPD
4. MRC Score of 3 or more. (walk slower than people of the same age on the level or stops for breath when walking at own pace on the level)
5. Able to read and write in English

## **Participant type(s)**

Patient

**Age group**

Adult

**Sex**

Both

**Target number of participants**

24

**Key exclusion criteria**

1. Significant disease (other than COPD) that could cause dyspnoea or exercise limitation
2. Contraindications for exercise (unstable cardiovascular disease; hypertension etc, a full list is described by the American College of Sports Medicine and is routine deployed)
3. Inability/unwillingness to use the device
4. Contraindications to using HFAO device (including severe right heart failure with hypotension), current severe haemoptysis, ineffective cough, rib fractures, pregnancy, current or recent pneumothorax, epilepsy, current pulmonary embolism, oesophageal varices, recent thoracic, upper gastro-intestinal tract or facial surgery)
5. Previously engaged in exercised based research or pulmonary rehab in the last 6 months.
6. Inability to secure informed consent
7. Those unable to communicate in full English will be excluded as the user manual is only available in English

**Date of first enrolment**

01/05/2016

**Date of final enrolment**

13/09/2016

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

England

United Kingdom

**Study participating centre**

University Hospitals of Leicester

Groby Road

Leicester

United Kingdom

LE3 9QP

**Sponsor information**

## Organisation

University Hospitals of Leicester

## Sponsor details

Trust HQ, Level 3, Balmoral Building  
Leicester Royal Infirmary  
Infirmary Square  
Leicester  
England  
United Kingdom  
LE1 5WW

## Sponsor type

Hospital/treatment centre

## ROR

<https://ror.org/02fha3693>

## Funder(s)

### Funder type

Industry

### Funder Name

Actegy LTD

## Results and Publications

### Publication and dissemination plan

The results of this study is intended to be published in international journals and presented at conferences- to be confirmed. The results will also inform the main study which will commence after the completion of the pilot study.

### Intention to publish date

31/01/2017

### Individual participant data (IPD) sharing plan

The current 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?
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<a href="#">Results article</a>	results	01/05/2018	24/01/2019	Yes	No
<a href="#">HRA research summary</a>			28/06/2023	No	No