







CELEB: Lung volume reduction in COPD - surgery

Submission date 23/05/2016	Recruitment status No longer recruiting	 Prospectively registered
		 Protocol added
Registration date 23/05/2016	Overall study status Completed	 SAP not yet added
		 Results added
Last Edited 03/05/2023	Condition category Respiratory	 Raw data not yet added
		 Study completed

Plain English Summary

Background and study aims

Chronic obstructive pulmonary disease (COPD) is the collective name for a group of diseases which affect the lungs. One of the diseases is emphysema, a lung condition in which the tiny air sacs (alveoli) in the lungs become damaged over time. In emphysema, the alveoli become overfilled with air, causing the walls to weaken and eventually burst, creating one larger air space instead of many small ones. This means that the sufferer is unable to breathe out fully, leading to breathlessness which becomes progressively worse. Lung volume reduction surgery (LVRS) is an operation in which the worst affected part of the lung is removed, making more space for the remaining healthier lung to function. There is good evidence that in properly selected individuals this can improve breathlessness and increase life expectancy. It is a major operation and does carry a risk of complications however. A more recent approach is to use a fiberoptic camera (bronchoscope) to place valves into the airways of the lung. These stop air from entering the worst-affected section of the lung causing it to collapse, leading to similar benefits of LVRS. The aim of this study is to compare the effectiveness of these two techniques in the treatment of patients with emphysema.

Who can participate?

Adults with a heterogeneous pattern of emphysema (isolated to certain areas of the lungs and to a varying extent between segments of the lungs)

What does the study involve?

Participants are randomly allocated to one of two groups. Those in the first group undergo lung volume reduction surgery. Those in the second group have endobronchial valves placed into their airways using a bronchoscope while they are sedated. One year after the procedures, participants in both groups undergo a number of physical tests to find out if their lung function has improved as well as their overall quality of life.

What are the possible benefits and risks of participating?

Not provided at time of registration

Where is the study run from?

1. Royal Brompton Hospital (UK)
2. Glenfield Hospital (UK)

3. St Bartholomew's Hospital (UK)
4. Northern General Hospital (UK)
5. Golden Jubilee National Hospital (UK)

When is the study starting and how long is it expected to run for?
April 2016 to September 2020

Who is funding the study?
National Institute for Health Research (UK)

Who is the main contact?
Sara Buttery
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Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Protocol/serial number
30523

Study information

Scientific Title
The CELEB trial: Comparative Effectiveness of Lung volume reduction surgery for Emphysema and

Acronym
CELEB

Study hypothesis

The aim of this study is to evaluate the relative effectiveness and value of two options currently available for the treatment of COPD patients with a heterogeneous pattern of emphysema. These are lung volume reduction surgery (LVRS) and the bronchoscopic placement of endobronchial valves (BLVR).

Ethics approval required

Old ethics approval format

Ethics approval(s)

London - Fulham Research Ethics Committee, 18/02/2016, ref: 16/LO/0286

Study design

Interventional; Design type: Treatment, Device, Surgery

Primary study design

Intentional

Secondary study design

Randomised parallel trial

Study setting(s)

Hospital

Study type(s)

Treatment

Participant information sheet

Not available in web format, please use the contact details to request a patient information sheet

Condition

Emphysema

Interventions

Patients will be randomised to either unilateral video assisted thoracoscopic (VATS) lung volume reduction surgery (LVRS) or endobronchial valves (BLVR) placed to achieve lobar occlusion.

Intervention Type

Device

Primary outcome measure

Change in iBODE score (a composite of BMI, FEV1, MRC dyspnoea score and shuttle walk test distance) one year post procedure.

Secondary outcome measures

1. Health related quality of life is measured using the COPD assessment test score (CAT)
2. Physical activity level
3. Change in residual volume (RV)
4. Fat free mass

Overall study start date

01/04/2016

Overall study end date

30/09/2020

Eligibility

Participant inclusion criteria

1. Adults (aged 18 or over) with COPD
2. FEV1 <60% predicted
3. Significant hyperinflation (TLC>100% predicted, RV>170% predicted)
4. Ex-smoker >3 months
5. MRC dyspnoea score 3 or more
6. CT scan assessed at MDT to have intact interlobar fissures (>90%) and heterogeneous emphysema
7. Provision of informed consent to participate

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex

Both

Target number of participants

Planned Sample Size: 76; UK Sample Size: 76

Total final enrolment

88

Participant exclusion criteria

1. Smoking within 3 months
2. CT scan shows interlobar fissures are not intact
3. Major comorbidity limiting survival
4. Significant pulmonary fibrosis
5. FEV1 and TLco <20%
6. PaO2 < 7.0kPa
7. PaO2 > 7kPa
8. Presence of interlobar collateral ventilation assessed by Chartis system

Recruitment start date

01/06/2016

Recruitment end date

01/06/2019

Locations**Countries of recruitment**

England

Scotland

United Kingdom

Study participating centre**Royal Brompton Hospital**

Fulham Road

London

United Kingdom

SW3 6NP

Study participating centre**Glenfield Hospital**

University Hospitals Leicester

United Kingdom

LE3 9QP

Study participating centre**St Bartholomew's Hospital**

Barts Health NHS Foundation Trust

United Kingdom

EC1A 7BE

Study participating centre**Northern General Hospital**

Sheffield

United Kingdom

S5 7AU

Study participating centre

Golden Jubilee National Hospital

Glasgow

United Kingdom

G81 4DY

Sponsor information

Organisation

Royal Brompton and Harefield NHS Foundation Trust

Sponsor details

Fulham Road

London

England

United Kingdom

SW3 6NP

Sponsor type

Hospital/treatment centre

ROR

<https://ror.org/02218z997>

Funder(s)

Funder type

Government

Funder Name

National Institute for Health 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

Publication and dissemination plan

Not provided at time of registration

Intention to publish date

30/09/2021

Individual participant data (IPD) sharing plan

Not provided at time of registration

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?
Protocol article	protocol	17/10/2018	30/10/2019	Yes	No
Results article		27/04/2023	03/05/2023	Yes	No
HRA research summary			28/06/2023	No	No