Ambulatory oxygen improves the effectiveness of pulmonary rehabilitation in selected patients

Submission date Recruitment status Prospectively registered 26/04/2010 No longer recruiting [] Protocol [] Statistical analysis plan Registration date Overall study status 13/05/2010 Completed [X] Results [] Individual participant data **Last Edited** Condition category 29/07/2013 Respiratory

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Ms Julia Bott

Contact details

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Additional identifiers

Protocol serial number TC08/5

Study information

Scientific Title

The effectiveness of ambulatory oxygen versus room air during pulmonary rehabilitation in patients who are hypoxaemic on exertion and respond to supplemental oxygen at baseline

Study objectives

The aims of this prospective randomised controlled trial is to determine whether ambulatory oxygen during pulmonary rehabilitation provides additional benefit, compared with room air, for patients hypoxaemic only on exertion and who respond to supplemental oxygen at baseline. Additionally, to determine whether the degree of acute response to ambulatory oxygen predicted any additional benefit.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Surrey Ethics Committee approved on the 7th August 2007 (ref: 07/H1109/89)

Study design

Prospective multicentre single blind randomised controlled trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Chronic respiratory disease

Interventions

Patients randomised into two groups:

- 1. Ambulatory oxygen group exercise within pulmonary rehabilitation and at home using ambulatory oxygen
- 2. Room air group exercise within pulmonary rehabilitation and at home on room air

Secondary sponsor details: Trevor Clay Memorial Fund (UK) British Lung Foundation 73 - 75 Goswell Road London EC1V 7ER

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

Exercise tolerance measured using the Endurance Shuttle Walk Test pre- and post-pulmonary rehabilitation

Key secondary outcome(s))

- 1. Quality of life measured using the Self-Report Chronic Respiratory Questionnaire
- 2. Anxiety and depression measured using the Hospital Anxiety and Depression Scale
- 3. Function measured using the Surrey Information on Function Tool

All are questionnaires completed pre- and post-pulmonary rehabilitation.

Completion date

30/09/2009

Eligibility

Key inclusion criteria

Patients (aged 50 - 87 years, either sex) with chronic respiratory disease enrolled in pulmonary rehabilitation who have resting saturations greater than 92%, but desaturate by greater than 4% and to less than 90% on exertion and who then walk greater than 10% further with ambulatory oxygen during field exercise testing at baseline.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

Key exclusion criteria

- 1. Already on oxygen therapy (long-term or ambulatory)
- 2. Those who do not desaturate significantly on exertion
- 3. Those who do desaturate but do not demonstrate benefit using ambulatory oxygen at baseline

Date of first enrolment

01/09/2007

Date of final enrolment

30/09/2009

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

The Respiratory Care Team

Chertsey United Kingdom KT16 0QA

Sponsor information

Organisation

Surrey Community Health (UK)

Funder(s)

Funder type

Charity

Funder Name

Surrey Community Health (UK)

Funder Name

British Lung Foundation (UK) - Trevor Clay Memorial Fund research grant

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration

Study outputs

| Output type | Details | Date created Date added | Peer reviewed? | Patient-facing? |
|-------------------------------|-------------------------------|-------------------------|----------------|-----------------|
| Results article | results | 01/05/2012 | Yes | No |
| Participant information sheet | Participant information sheet | 11/11/2025 11/11/2025 | No | Yes |