Inspiratory muscle training and exercise goal setting to improve functional ability and relieve the symptoms of dyspnoea in advanced cancer patients

Submission date	Recruitment status	Prospectively registered
18/12/2013	No longer recruiting	Protocol
Registration date	Overall study status	Statistical analysis plan
11/04/2014	Completed	Results
Last Edited	Condition category	Individual participant data
13/04/2017	Cancer	Record updated in last year

Plain English summary of protocol

http://www.cancerresearchuk.org/cancer-help/trials/a-study-looking-breathing-exercises-relieve-breathlessness-caused-by-advanced-cancer

Contact information

Type(s)

Scientific

Contact name

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Additional identifiers

Protocol serial number EHSDYSPNOEA

Study information

Scientific Title

A pilot study to investigate whether inspiratory muscle training and exercise goal setting can improve functional ability and relieve the symptoms of dyspnoea in advanced cancer patients

Study objectives

Inspiratory muscle training alongside exercise goal setting will relieve symptoms of dyspnoea for advanced cancer patients.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Pilot randomised controlled trial

Primary study design

Interventional

Study type(s)

Quality of life

Health condition(s) or problem(s) studied

Cancer/dyspnoea

Interventions

Participants are randomised to two groups:

1. Inspiratory muscle training (IMT) and goal setting diary:

An 8 week course of self-directed IMT and exercise. Measurements at weeks 1 and 8: 6MWT, inspiratory muscle pressure, spirometry and a questionnaire relating to dyspnoea symptoms.

2. Control group: Do not receive IMT. Offered breathing control techniques and goal setting advice after they have completed their 8 week measurements.

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

- 1. Functional ability as assessed by the 6-minute walk test (6MWT)
- 2. Relief of dyspnea symptoms and health-related quality of life (HRQoL) as assessed by the St. Georges Respiratory questionnaire (SGRQ)

These are measured at week 1 and week 8.

Key secondary outcome(s))

- 1. Change in spirometry measurements
- 2. Maximal inspiratory mouth pressure (MIP) can be assessed using the IMT devices or using a

handheld respiratory pressure meter

3. Feasibility of the intervention will be assessed by monitoring compliance with the self-directed IMT/exercise logged in the weekly diaries and from any pertinent qualitative data gathered from focus groups

These are measured at week 1 and week 8.

Completion date

01/07/2014

Eligibility

Key inclusion criteria

- 1. Aged 16 or above
- 2. Diagnosed with any type of cancer for which they are having palliative treatment or have ended treatment
- 3. Showing symptoms of dyspnea (including exertional) as assessed by their physician
- 4. Ability to engage in gentle exercise

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

- 1. Patients who have had recent thoracic surgery
- 2. Patients with mental incapacity unable to give informed consent
- 3. Patients unable to understand verbal and written information in English
- 4. Patients who have co-existing moderate or severe chronic obstructive pulmonary disease (COPD)
- 5. Patients on active treatment for cancer
- 6. Patients undergoing radical treatment with curative intent

Date of first enrolment

01/03/2014

Date of final enrolment

01/07/2014

Locations

Countries of recruitment

United Kingdom

Study participating centre EHS Faculty Derby United Kingdom DE221GB

Sponsor information

Organisation

University of Derby (UK)

ROR

https://ror.org/02yhrrk59

Funder(s)

Funder type

University/education

Funder Name

University of Derby (UK)

Funder Name

Royal Derby Hospital (UK) - Lung cancer and mesothelioma Derby account

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type