

An assessment of patient outcomes following a rehabilitation programme for patients who have received heart/lung or lung transplant - a randomised controlled pilot trial

Submission date 30/09/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 30/09/2004	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 28/02/2018	Condition category Surgery	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0201139562

Study information

Scientific Title

An assessment of patient outcomes following a rehabilitation programme for patients who have received heart/lung or lung transplant - a randomised controlled pilot trial

Study objectives

Does the undertaking of a multidisciplinary-led programme of rehabilitation facilitate a better Quality of Life than a document-based rehabilitation programme in lung and or heart/lung transplant out-patients?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Quality of life

Participant information sheet

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

Health condition(s) or problem(s) studied

Surgery: Heart/lung transplant

Interventions

Some patients will receive document packs while some will receive a multidisciplinary-led programme of rehabilitation.

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/07/2004

Completion date

30/06/2005

Eligibility**Key inclusion criteria**

20 patients

Participant type(s)

Patient

Age group

Not Specified

Sex

Both

Target number of participants

20

Key exclusion criteria

Does not meet inclusion criteria

Date of first enrolment

01/07/2004

Date of final enrolment

30/06/2005

Locations**Countries of recruitment**

England

United Kingdom

Study participating centre
Harefield Hospital
Harefield
United Kingdom
UB9 6JH

Sponsor information

Organisation
Department of Health

Sponsor details
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

Website
<http://www.dh.gov.uk/Home/fs/en>

Funder(s)

Funder type
Hospital/treatment centre

Funder Name
Royal Brompton and Harefield NHS Trust (UK)

Results and Publications

Publication and dissemination plan
Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration