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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

Registration date
30/09/2004

Overall study status
Completed

Last Edited
28/02/2018

Condition category
Surgery

[ ] Prospectively registered
[ ] Protocol
[ ] Statistical analysis plan
[ ] Results
[ ] Individual participant data
[ ] Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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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