

Cost effective rehabilitation after shoulder surgery

Submission date 12/09/2003	Recruitment status Stopped	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 12/09/2003	Overall study status Stopped	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 19/10/2011	Condition category Surgery	<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

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0436061262

Study information

Scientific Title

Study objectives

To determine whether a self directed, hospital supervised rehabilitation program after shoulder surgery can equal or improve upon the results obtained with a hospital administered program, changing the role of the physiotherapist from service provider to supervisor.

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)

Not specified

Study type(s)

Not Specified

Participant information sheet

Health condition(s) or problem(s) studied

Surgery: Shoulder

Interventions

1. Supervised programme
2. Non-supervised programme

Trial stopped in 2003, no eligible patients.

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome measure

Cost reduction in rehabilitation after shoulder surgery.

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/10/2000

Completion date

01/10/2003

Reason abandoned (if study stopped)

No eligible patients

Eligibility

Key inclusion criteria

50 patients per year requiring rehabilitation after shoulder surgery.

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

50

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/10/2000

Date of final enrolment

01/10/2003

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Department of Orthopaedic Surgery
Leeds
United Kingdom
LS9 7TF

Sponsor information

Organisation

Department of Health (UK)

Sponsor details

Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type

Government

Website

<http://www.doh.gov.uk>

Funder(s)

Funder type

Government

Funder Name

Leeds Teaching Hospitals 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