Arthroscopic Capsular Release of Frozen Shoulder. Post-operative rehabilitation Programme. Immediate mobilisation versus immediate mobilisation plus external rotation splint. A Randomised Pilot Study

Submission date	Recruitment status	Prospectively registered
28/09/2007	No longer recruiting	Protocol
Registration date	Overall study status	Statistical analysis plan
28/09/2007	Completed	Results
Last Edited	Condition category	[] Individual participant data
14/10/2014	Surgery	Record updated in last year

Plain English summary of protocolNot 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

N0207182917

Study information

Scientific Title

Study objectives

Not provided at time of registration

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised controlled pilot study

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Surgery: Arthroscopy

Interventions

- 1. Patients receive standard regime no sling and mobilise immediate post-operative
- 2. Patients have their treated arm placed in an external rotation splint, which is used at rest and night for 10 days.

Intervention Type

Procedure/Surgery

Phase

Not Applicable

Primary outcome measure

- 1. Symptoms and Functions Pain & Range of movement
- 2. One main subjective measurement Constant Score
- 3. Two main subjective scores VAS pain, Oxford shoulder questionnaire

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/12/2005

Completion date

01/12/2007

Eligibility

Key inclusion criteria

Not provided at time of registration

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

15 patients in each group

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/12/2005

Date of final enrolment

01/12/2007

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Orthopaedic Surgery

Liverpool United Kingdom L7 8XP

Sponsor information

Organisation

Record Provided by the NHSTCT Register - 2007 Update - Department of Health

Sponsor details

The Department of Health, Richmond House, 79 Whitehall London United Kingdom SW1A 2NL +44 (0)20 7307 2622 dhmail@doh.gsi.org.uk

Sponsor type

Government

Website

http://www.dh.gov.uk/Home/fs/en

Funder(s)

Funder type

Government

Funder Name

Royal Liverpool and Broadgreen University Hospitals Trust (UK)

Funder Name

NHS R&D Support Funding (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 summaryNot provided at time of registration