ISRCTN76627520 https://doi.org/10.1186/ISRCTN76627520

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 28/09/2007	**Recruitment status** No longer recruiting	[ ] Prospectively registered [ ] Protocol
Registration date 28/09/2007	**Overall study status** Completed	[ ] Statistical analysis plan [ ] Results
Last Edited 14/10/2014	**Condition category** Surgery	[ ] 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
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
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Sponsor information

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

Sponsor details
The Department of Health, Richmond House, 79 Whitehall
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Sponsor type
Government

Website
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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 summary
Not provided at time of registration