ISRCTN48254181 https://doi.org/10.1186/ISRCTN48254181

A comparison study of internal versus external rotation immobilisation for primary anterior shoulder dislocation

Submission date
29/09/2006

Recruitment status
No longer recruiting

Registration date
29/09/2006

Overall study status
Completed

Last Edited
18/08/2015

Condition category
Musculoskeletal Diseases

[ ] 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

Protocol serial number
N0523163139

Study information

Scientific Title
A comparison study of internal versus external rotation immobilisation for primary anterior shoulder dislocation

Study objectives
Not provided at time of registration

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Primary study design
Interventional

Study design
Randomised controlled trial

Study type(s)
Treatment

Health condition(s) or problem(s) studied
Musculoskeletal Diseases: Shoulder dislocation

Interventions
Arm [A] external rotation
Arm [B] internal rotation

Intervention Type
Procedure/Surgery

Primary outcome(s)
To assess rate of re dislocation per technique of splinting

Key secondary outcome(s)
Not provided at time of registration

Completion date
01/05/2008

Eligibility

Key inclusion criteria
1. Age 20-40 years old
2. Traumatic dislocations
3. First time dislocations
4. No associated fractures of the region
5. No associated rotator cuff tears
6. No co morbidity that affect general health

Participant type(s)
Patient

Healthy volunteers allowed
No

Age group
Adult

Sex
All

Key exclusion criteria
1. Recurrent disclocation
2. Unable to co operate for physiotherapy and/or follow up
3. Co morbidity
4. Associated fracture or rotator cuff injury
5. Significant ipsilateral injury
6. Open injuries
7. Neurological disease
8. Collagen disease
9. Pregnancy
10. Unfit for surgery

Date of first enrolment
01/01/2005

Date of final enrolment
01/05/2008

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
Good Hope Hospital NHS Trust
Birmingham
United Kingdom
B75 7RR

Sponsor information

Organisation
Record Provided by the NHSTCT Register - 2006 Update - Department of Health (UK)

Funder(s)

Funder type
Government

Funder Name
Good Hope Hospital NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration