

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

Submission date 29/09/2006	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 29/09/2006	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 18/08/2015	Condition category Musculoskeletal Diseases	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<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

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 dislocation
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