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

Submission date	Recruitment status	[ ] Prospectively registered
29/09/2006	No longer recruiting	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
29/09/2006	Completed	[ ] Results
Last Edited	**Condition category**	[ ] Individual participant data
18/08/2015	Musculoskeletal Diseases	[ ] 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
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

Study design
Randomised controlled trial

Primary study design
Interventional

Secondary study design
Randomised controlled trial

Study setting(s)
Hospital

Study type(s)
Treatment

Participant information sheet
Not available in web format, please use the contact details below to request a patient information sheet

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 measure
To assess rate of re dislocation per technique of splinting

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/01/2005

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

Age group
Adult

Sex
Both

Target number of participants
75 patients per group

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
England

United Kingdom

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)

Sponsor details
The Department of Health, Richmond House, 79 Whitehall
London
United Kingdom
SW1A 2NL
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dhmail@doh.gsi.org.uk

Sponsor type
Government

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

Funder(s)

Funder type
Government

Funder Name
Good Hope Hospital 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