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A comparison of Arthroscopic subacromial decompression by soft tissue excision with arthroscopic excision with soft tissue and bone excision

Submission date 30/09/2004	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol
Registration date 30/09/2004	**Overall study status** Completed	[_] Statistical analysis plan [_] Results
Last Edited 01/10/2013	**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
N0191133189

Study information

Scientific Title

Study objectives
To compare the results of two surgical procedures for subacromial decompression of the shoulder, one removing soft tissue only and one removing bone and soft tissue.

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)
Not Specified

Participant information sheet

Health condition(s) or problem(s) studied
Surgery: Subacromial decompression

Interventions
Random allocation of patients into two groups (removing soft tissue only vs removing bone and soft tissue) and assessment of results post-operatively with questionnaires at follow up visits to assess patient satisfaction.

Intervention Type
Procedure/Surgery

Phase
Not Specified

Primary outcome measure
Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date
13/11/2003

Completion date
13/11/2006

Eligibility

Key inclusion criteria
Patients attending the Orthopaedic out-patient clinic for problems with shoulder impingement.

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
Not provided at time of registration

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
13/11/2003

Date of final enrolment
13/11/2006

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Queens Hospital
Burton upon Trent
United Kingdom
DE13 0RB

Sponsor information

Organisation
Department of Health

Sponsor details
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

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

Funder(s)

Funder type
Government

Funder Name
Burton Hospitals 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