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A true to life study of pain management early in rotator cuff tendonitis and adhesive capsulitis

Submission date 30/09/2005	**Recruitment status** No longer recruiting	[ ] Prospectively registered [ ] Protocol
Registration date 30/09/2005	**Overall study status** Completed	[ ] Statistical analysis plan [ ] Results
Last Edited 08/04/2014	**Condition category** Signs and Symptoms	[ ] 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
N0080151784

Study information

Scientific Title

Study objectives
To determine whether early intervention with corticosteroid injection is effective in reducing acute pain from rotator cuff tendonitis or adhesive capsulitis.

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)
Not specified

Study type(s)
Not Specified

Participant information sheet

Health condition(s) or problem(s) studied
Signs and Symptoms: Pain

Interventions
Randomised study

Intervention Type
Drug

Phase
Not Specified

Drug/device/biological/vaccine name(s)
Corticosteroids

Primary outcome measure
To measure the change in perceived pain and disability to inform treatment decisions and improve the evidence base for care of patients with this condition.

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/11/2004

Completion date
01/08/2006

Eligibility

Key inclusion criteria
150 patients

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
150

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/11/2004

Date of final enrolment
01/08/2006

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Consultant Rheumatologist
London
United Kingdom
E11 1NR

Sponsor information

Organisation
Department of Health

Sponsor details
Richmond House
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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
Whipps Cross University Hospital NHS Trust (UK)

Funder Name
NHS R&D Support Funding

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