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

Submission date	Recruitment status	Prospectively registered
30/09/2005	No longer recruiting	Protocol
Registration date	Overall study status	Statistical analysis plan
30/09/2005	Completed	Results
Last Edited	Condition category	Individual participant data
08/04/2014	Signs and Symptoms	<ul><li>Record updated in last year</li></ul>

### Plain English summary of protocol

Not provided at time of registration

### Contact information

### Type(s)

Scientific

#### Contact name

Dr Alan Hakim

#### Contact details

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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 79 Whitehall London United Kingdom SW1A 2NL +44 (0)20 7307 2622 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