

A Prospective Randomised Double Blind Trial of Subacromial Shoulder Injections for Impingement Syndrome

Submission date 30/09/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 30/09/2004	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 18/08/2015	Condition category Signs and Symptoms	<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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0247130745

Study information

Scientific Title

A Prospective Randomised Double Blind Trial of Subacromial Shoulder Injections for Impingement Syndrome

Study objectives

Is there any difference between local anaesthetic, steroid and saline injections in the treatment of subacromial impingement syndrome?

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

Signs and Symptoms: Pain

Interventions

Comparison of local anaesthetic, steroid and saline injections

Intervention Type

Drug

Phase

Not Applicable

Primary outcome measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

10/11/2003

Completion date

01/07/2004

Eligibility

Key inclusion criteria

New patient referrals to specialist shoulder clinic

Participant type(s)

Patient

Age group

All

Sex

Both

Target number of participants

Not provided at time of registration

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

10/11/2003

Date of final enrolment

01/07/2004

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Swindon & Marlborough NHS Trust
Swindon
United Kingdom
SN3 6BB

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

Swindon and Marlborough 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