

Does aspiration of simple traumatic elbow effusions benefit the patient?

Submission date 12/09/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
Last Edited 30/04/2015	Condition category Musculoskeletal Diseases	<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
N0106067610

Study information

Scientific Title

Does aspiration of simple traumatic elbow effusions benefit the patient?

Study objectives

Does aspiration of traumatic elbow effusions reduce pain and increase mobility and quality of life?

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)

Quality of life

Participant information sheet**Health condition(s) or problem(s) studied**

Musculoskeletal Diseases: Joint effusion

Interventions

1. Aspiration
2. No aspiration

Intervention Type

Procedure/Surgery

Primary outcome measure

1. Range of movement
2. Pain
3. Quality of life

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/03/2000

Completion date

30/06/2003

Eligibility

Key inclusion criteria

Adults attending Accident and Emergency Department with traumatic elbow effusions

Participant type(s)

Patient

Age group

Adult

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

01/03/2000

Date of final enrolment

30/06/2003

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Gloucestershire Royal Hospital

Gloucester

United Kingdom

GL1 3NN

Sponsor information

Organisation

Department of Health (UK)

Sponsor details

Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type

Government

Website

<http://www.doh.gov.uk>

Funder(s)

Funder type

Government

Funder Name

Gloucestershire R&D Consortium (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