

A randomised controlled trial of physiotherapy and osteopathy in the treatment of chronic adhesive capsulitis

Submission date 12/09/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 20/02/2020	Condition category Musculoskeletal Diseases	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<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

N0544093465

Study information

Scientific Title

-

Study objectives

Comparison of physiotherapy to osteopathy treatment for frozen shoulder

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)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Chronic adhesive capsulitis

Interventions

Patients with suspected chronic adhesive capsulitis, as indicated by painful and limited shoulder movement during the doctor's and physiotherapist's assessment will be offered participation in this study. This will involve random allocation to one of three groups: a control group receiving rest and advice only, a physiotherapy group receiving a combination of manual therapy and therapeutic exercise, and an osteopathy group receiving a combination of osteopathic manual therapy and exercise.

Intervention Type

Other

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

26/07/2000

Completion date

26/07/2003

Eligibility

Key inclusion criteria

60

Participant type(s)

Patient

Age group

Adult

Sex

Not Specified

Target number of participants

60

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

26/07/2000

Date of final enrolment

26/07/2003

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Box 194

Cambridge

United Kingdom
CB2 2QQ

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

Research organisation

Funder Name

Cambridge Consortium - Addenbrooke's (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