

A randomised controlled trial to investigate the effectiveness of acupuncture as a treatment for fibromyalgia

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 02/02/2017	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

Contact name

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0263104896

Study information

Scientific Title

A randomised controlled trial to investigate the effectiveness of acupuncture as a treatment for fibromyalgia

Study objectives

Is acupuncture better than placebo for treating fibromyalgia?

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

Health condition(s) or problem(s) studied

Musculoskeletal Diseases: Fibromyalgia

Interventions

A. Acupuncture

B. Placebo

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Pain threshold measurement by a dolorimeter number of painful trigger points. Scores on questionnaires validated for use in fibromyalgia. Visual analogue scale (VAS) of global health status.

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/09/2001

Completion date

01/08/2006

Eligibility

Key inclusion criteria

80 Patients from Pain Management.

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

80

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/09/2001

Date of final enrolment

01/08/2006

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Anaesthetics Department

London

United Kingdom

W1P 8AN

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

University College London Hospitals 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