

The effectiveness of magnetic bracelets as used in the consumer market for pain in osteoarthritis

Submission date 15/07/2002	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 15/07/2002	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 02/10/2007	Condition category Musculoskeletal Diseases	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data

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

Study information

Scientific Title

Study objectives

To determine the effectiveness of commercially available magnetic bracelets for pain control in osteoarthritis of the hip and knee.

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

Osteoarthritis

Interventions

1. Will receive bracelets with active magnets
2. Will receive the placebo bracelet, with less active magnets
3. Will receive a control bracelet, with no magnets

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

01/12/2001

Completion date

01/12/2003

Eligibility

Key inclusion criteria

Men and women aged 45 to 80 years at entry into the trial with osteoarthritis of the hip and or knee either radiologically proven or diagnosed by a consultant rheumatologist or consultant orthopaedic surgeon.

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

194

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/12/2001

Date of final enrolment

01/12/2003

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Department of Complementary Medicine
Exeter
United Kingdom
EX2 4NT

Sponsor information

Organisation

Arthritis Research Campaign (ARC) (UK)

Sponsor details

Copeman House
St Mary's Court
St Mary's Gate
Chesterfield
Derbyshire
United Kingdom
S41 7TD

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info@arc.org.uk

Sponsor type

Charity

Website

<http://www.arc.org.uk>

ROR

<https://ror.org/02jkpm469>

Funder(s)

Funder type

Charity

Funder Name

Arthritis Research Campaign (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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	Results	18/12/2004		Yes	No