

A randomised controlled trial of vertebroplasty for the treatment of osteoporotic vertebral crush fractures

Submission date 12/09/2003	Recruitment status No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 11/10/2016	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

N0213112414

Study information

Scientific Title

A randomised controlled trial of vertebroplasty for the treatment of osteoporotic vertebral crush fractures

Study objectives

Not provided at time of registration

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 contact details to request a participant information sheet

Health condition(s) or problem(s) studied

Osteoporotic crush fractures

Interventions

Randomised to vertebroplasty or best medical treatment.

Intervention Type

Procedure/Surgery

Primary outcome measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

28/11/2005

Completion date

30/11/2005

Eligibility

Key inclusion criteria

1. History of vertebral crush fractures proven on radiograph
2. Causes of crush fractures other than osteoporosis excluded
3. Persistent moderate/severe pain after 4 weeks conservative treatment
4. No more than four fractures

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

Not provided at time of registration

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

28/11/2005

Date of final enrolment

30/11/2005

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

St Richards Hospital

Chichester

United Kingdom

PO19 4SE

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

Hospital/treatment centre

Funder Name

Royal West Sussex 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