

The effects of an intervention to treat and prevent depression in older people following hip fracture

Submission date 01/09/2005	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 14/09/2005	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 20/12/2007	Condition category Mental and Behavioural Disorders	<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

N/A

Study information

Scientific Title

Study objectives

An intervention will reduce the incidence of depression and improve the symptoms of depression following hip fracture.

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

Depression in orthopaedic patients

Interventions

Cognitive behaviour therapy (CBT) for prevention and nurse intervention for treatment versus no intervention

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Rates of depression.

Secondary outcome measures

Functional outcome.

Overall study start date

01/01/2004

Completion date

01/06/2005

Eligibility

Key inclusion criteria

People over the age of 60 who have fractured their hip.

Participant type(s)

Patient

Age group

Senior

Sex

Both

Target number of participants

200

Key exclusion criteria

People who do not satisfy the inclusion criteria.

Date of first enrolment

01/01/2004

Date of final enrolment

01/06/2005

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Wythenshawe Hospital

Manchester

United Kingdom

M23 9LT

Sponsor information

Organisation

University of Manchester (UK)

Sponsor details

Wythenshawe Hospital
Manchester
England
United Kingdom
M23 9LT

Sponsor type

University/education

ROR

<https://ror.org/027m9bs27>

Funder(s)**Funder type**

Government

Funder Name

Health Foundation (UK) (ref: orthopaedic study)

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	01/11/2006		Yes	No