

# The effect of clodronate on the incidence of hip fracture and consequences for health and costs

<b>Submission date</b> 25/10/2000	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
<b>Registration date</b> 25/10/2000	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
<b>Last Edited</b> 12/09/2007	<b>Condition category</b> Musculoskeletal Diseases	<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

G9518113

## Study information

## **Scientific Title**

### **Study objectives**

To determine in an elderly female population who would best benefit from an intervention aimed at reducing the incidence of 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)**

Prevention

### **Participant information sheet**

### **Health condition(s) or problem(s) studied**

Orthopaedics, rheumatology

### **Interventions**

Clodronate/control

### **Intervention Type**

Other

### **Phase**

Not Specified

### **Primary outcome measure**

Fragility fractures, particularly hip fracture

### **Secondary outcome measures**

Not provided at time of registration

### **Overall study start date**

10/06/1996

**Completion date**

01/12/2002

## Eligibility

**Key inclusion criteria**

Women aged 75 years or older willing to participate

**Participant type(s)**

Patient

**Age group**

Senior

**Sex**

Female

**Target number of participants**

5700

**Key exclusion criteria**

Drugs and diseases likely to impair the efficacy of intervention or the interpretation of results

**Date of first enrolment**

10/06/1996

**Date of final enrolment**

01/12/2002

## Locations

**Countries of recruitment**

England

United Kingdom

**Study participating centre**

Department of Human Metabolism & Clinical Biochemistry

Sheffield

United Kingdom

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## Sponsor information

**Organisation**

Medical Research Council (MRC) (UK)

**Sponsor details**

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clinical.trial@headoffice.mrc.ac.uk

**Sponsor type**

Research council

**Website**

<http://www.mrc.ac.uk>

## **Funder(s)**

**Funder type**

Research council

**Funder Name**

Medical Research Council (UK)

**Alternative Name(s)**

Medical Research Council (United Kingdom), UK Medical Research Council, MRC

**Funding Body Type**

Government organisation

**Funding Body Subtype**

National government

**Location**

United Kingdom

## **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?
<a href="#">Results article</a>	Results	01/01/2007		Yes	No