Evaluation of early home rehabilitation program for geriatric patients with hip fractures

Submission date	Recruitment status	Prospectively registered
09/10/2002	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
09/10/2002	Completed	Results
Last Edited	Condition category	Individual participant data
01/07/2009	Injury, Occupational Diseases, Poisoning	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 422018

Study information

Scientific Title

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)

Not specified

Study type(s)

Not Specified

Participant information sheet

Health condition(s) or problem(s) studied

Gerentology, orthopaedics

Interventions

The patients will be randomized into two groups:

- 1. EHRP Group, the study group, where the patient is discharged (within 2-3 weeks post-surgery) when he/she could walk independently with a quadripod
- 2. TRP Group, the control group, where the patient is discharged (within 4-5 weeks post-surgery) when he/she could walk independently with a stick or without any other walking aids

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

01/04/2001

Completion date

30/04/2002

Eligibility

Key inclusion criteria

- 1. Geriatric patients less than 85 years of age with femoral-neck fractures
- 2. Patients who had Austin Moore Arthroplasty, Cemented Thompson replacement, internal fixation with hip screws or dynamic hip screw surgery
- 3. Patients who, pre-morbidly, could walk without support or with a stick
- 4. Patients with no active medical problems detected
- 5. Practice full weight bearing walking immediate post-operation, and
- 6. Patients living with family members

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

Not provided at time of registration.

Key exclusion criteria

Does not meet inclusion criteria

Date of first enrolment

01/04/2001

Date of final enrolment

30/04/2002

Locations

Countries of recruitment

China

Hong Kong

Study participating centre Physiotherapy Department

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Sponsor information

Organisation

Hong Kong Health Services Research Fund (Hong Kong)

Sponsor details

Health Welfare and Food Bureau Government Secretariat, HKSAR 20th floor Murray Building Garden Road

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Hong Kong

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+852 (0)2973 8288 hsrf@hwfb.gov.hk

Sponsor type

Government

Website

http://www.fhb.gov.hk/grants/english/funds/funds_hhsrf/funds_hhsrf_abt/funds_hhsrf_abt.html

ROR

https://ror.org/03qh32912

Funder(s)

Funder type

Not defined

Funder Name

Hong Kong Health Services Research Fund

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 summaryNot provided at time of registration