

Cardiac rehabilitation and prevention programme: its cost-effectiveness in Hong Kong

Submission date 10/10/2002	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 10/10/2002	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 02/07/2009	Condition category Circulatory System	<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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-
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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
511002

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)

Hospital

Study type(s)

Not Specified

Participant information sheet

Health condition(s) or problem(s) studied

Cardiovascular

Interventions

Patients were enrolled either into educational classes and exercise training 6 h/week for 8 weeks plus conventional medical management or conventional medical therapy alone. Costs of equipment and staff salary were estimated. A set of quality of life (QOL) questionnaires was used to collect data by interviewing.

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

01/04/2000

Completion date

30/04/2001

Eligibility

Key inclusion criteria

Those with acute myocardial infarction within 6 weeks or post-percutaneous transluminal coronary angioplasty (PTCA)

Participant type(s)

Patient

Age group

Other

Sex

Both

Target number of participants

Not provided at time of registration.

Key exclusion criteria

Does not meet inclusion criteria

Date of first enrolment

01/04/2000

Date of final enrolment

30/04/2001

Locations

Countries of recruitment

Hong Kong

Study participating centre

Department of Medicine

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Hong Kong

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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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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 (Hong Kong)

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	01/12/2004		Yes	No