

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

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

**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**  
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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**Sponsor type**

Government

**Website**

[http://www.fhb.gov.hk/grants/english/funds/funds\\_hhsrf/funds\\_hhsrf\\_abt/funds\\_hhsrf\\_abt.html](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?
<a href="#">Results article</a>	results	01/12/2004		Yes	No