Does intensive rehabilitation improve the functional outcome of traumatic brain injury?

Submission date	Recruitment status No longer recruiting	Prospectively registered	
10/10/2002		☐ Protocol	
Registration date	Overall study status	Statistical analysis plan	
10/10/2002	Completed	[X] Results	
Last Edited 02/07/2009	Condition category Injury, Occupational Diseases, Poisoning	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

531019

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)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Traumatic brain injury

Interventions

Head injured patients were randomised into the conventional group and the intensive group:

- 1. Subjects belonging to the conventional group underwent the usual programme of physiotherapy, occupational therapy and speech therapy 2 hours a day, 5.5 days a week
- 2. For those in the intensive group, therapy time was doubled to 4 hours per day without changing the content of the therapy

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

01/01/2002

Completion date

01/01/2005

Eligibility

Key inclusion criteria

- 1. Moderate (Glasgow Coma Score [GCS] 9 12) and severe (GCS 8/15) traumatic brain injury
- 2. Aged from 12 to 65 years old

Participant type(s)

Patient

Age group

Other

Sex

Both

Target number of participants

Not provided at time of registration

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

01/01/2002

Date of final enrolment

01/01/2005

Locations

Countries of recruitment

Hong Kong

Study participating centre Division of Neurosurgery

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

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

Government

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 summaryNot provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/06/2007		Yes	No