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Paired kidney analysis of tacrolimus and cyclosporine microemulsion-based therapy in Chinese cadaveric renal transplant recipients

Submission date	Recruitment status	[_] Prospectively registered
02/11/2005	No longer recruiting	[_] Protocol
Registration date	**Overall study status**	[_] Statistical analysis plan
08/12/2005	Completed	[X] Results
Last Edited	**Condition category**	[_] Individual participant data
10/09/2009	Urological and Genital Diseases	

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
N/A

Study information

Scientific Title

Study objectives
To study whether tacrolimus is superior to cyclosporine in preserving renal function

Ethics approval required
Old ethics approval format

Ethics approval(s)
Yes, 1998

Study design
Prospective 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
Cadaveric renal transplant

Interventions
Tacrolimus therapy or Neoral cyclosporine therapy

Intervention Type
Drug

Phase
Not Specified

Drug/device/biological/vaccine name(s)
Tacrolimus and cyclosporine

Primary outcome measure
Patient and graft survival, rejection rate

Secondary outcome measures
Course of renal function, cardiovascular risks, infection and malignancy

Overall study start date
01/06/1998

Completion date
31/12/2004

Eligibility

Key inclusion criteria
Chinese patients receiving first cadaveric renal transplants consecutively between 01/06/98 and 31/12/04 in Queen Elizabeth Hospital in Hong Kong

Participant type(s)
Patient

Age group
Adult

Sex
Both

Target number of participants
100

Key exclusion criteria
Consent not available

Date of first enrolment
01/06/1998

Date of final enrolment
31/12/2004

Locations

Countries of recruitment
Hong Kong

Study participating centre
Department of Medicine
Kowloon
Hong Kong
-

Sponsor information

Organisation
Queen Elizabeth Hospital (Hong Kong)

Sponsor details
30, Gascoigne Road
Kowloon
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-
simoncycheung@gmail.com

Sponsor type
Hospital/treatment centre

ROR
https://ror.org/05ee2qy47

Funder(s)

Funder type
Hospital/treatment centre

Funder Name
Queen Elizabeth Hospital (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	01/08/2006		Yes	No