

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

Submission date
02/11/2005

Recruitment status
No longer recruiting

☐ Prospectively registered

☐ Protocol

Registration date
08/12/2005

Overall study status
Completed

☐ Statistical analysis plan

☒ Results

Last Edited
10/09/2009

Condition category
Urological and Genital Diseases

☐ Individual participant data

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Dr Simon Cheung

Contact details

Department of Medicine
Queen Elizabeth Hospital
30 Gascoigne Road
Kowloon
Hong Kong

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simoncycheung@gmail.com

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

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Sponsor information

Organisation

Queen Elizabeth Hospital (Hong Kong)

Sponsor details

30, Gascoigne Road

Kowloon

Hong Kong

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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