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Cardiovascular effects of medical (ie use of calcimimetics) vs surgical parathyroidectomy in dialysis of patients. A prospective randomised trial

Submission date 29/09/2006	**Recruitment status** Stopped	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [_] Results [_] Individual participant data [_] Record updated in last year
Registration date 29/09/2006	**Overall study status** Stopped	
Last Edited 13/10/2011	**Condition category** 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
N0205164619

Study information

Scientific Title

Study objectives
1. In patients undergoing dialysis, do calcimimetics reduce calcium deposition in the arteries and thereby reduce risk of cardiovascular events?
2. Is medical parathyroidectomy superior to surgical parathyroidectomy in reducing cardiovascular risks and improving abnormalities of mineral metabolism?

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)
Not specified

Study type(s)
Not Specified

Participant information sheet

Health condition(s) or problem(s) studied
Urological and Genital Diseases: Renal

Interventions
Calcimimetics vs surgical parathyroidectomy.

Added June 2008: the trial was stopped due to poor recruitment.

Intervention Type
Drug

Phase
Not Specified

Drug/device/biological/vaccine name(s)
calcimimetics

Primary outcome measure
Progression of disease as determined by multislice CT coronary calcium scores, Dexa scans pulse wave velocity, and carotid intima media thickness.

Secondary outcome measures
Not provided at time of registration

Overall study start date
28/03/2005

Completion date
30/09/2007

Reason abandoned (if study stopped)
Poor recruitment

Eligibility

Key inclusion criteria
Not provided at time of registration

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
50

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
28/03/2005

Date of final enrolment
30/09/2007

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Renal Unit
London
United Kingdom
E1 1BB

Sponsor information

Organisation
Record Provided by the NHSTCT Register - 2006 Update - Department of Health

Sponsor details
The Department of Health, Richmond House, 79 Whitehall
London
United Kingdom
SW1A 2NL
+44 (0)20 7307 2622
dhmail@doh.gsi.org.uk

Sponsor type
Government

Website
http://www.dh.gov.uk/Home/fs/en

Funder(s)

Funder type
Government

Funder Name
Barts and The London NHS Trust (UK)

Funder Name
Queen Mary University of London - QMUL

Funder Name
NHS R&D Support Funding

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