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A randomised controlled trial into the influence of peri-operative skin preparation on the incidence of post-operative wound sepsis in patients undergoing renal and pancreas transplants

Submission date 30/09/2004	**Recruitment status** No longer recruiting	[ ] Prospectively registered [ ] Protocol
Registration date 30/09/2004	**Overall study status** Completed	[ ] Statistical analysis plan [ ] Results
Last Edited 14/11/2014	**Condition category** Surgery	[ ] Individual participant data [ ] Record updated in last year

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
N0453126680

Study information

Scientific Title

Study objectives
Can the peri-operative skin preparation solution used reduce the incidence of post-operative wound sepsis?

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)
Prevention

Participant information sheet

Health condition(s) or problem(s) studied
Surgery: Renal and pancreas transplant

Interventions
Peri-operative skin preparation vs standard practice

Intervention Type
Procedure/Surgery

Phase
Not Applicable

Primary outcome measure

Reduction in wound sepsis in the experimental group

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/06/2003

Completion date
31/05/2004

Eligibility

Key inclusion criteria
Normal transplant follow-up patients

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
Not provided at time of registration

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/06/2003

Date of final enrolment
31/05/2004

Locations

Countries of recruitment
England

United Kingdom

Study participating centre

Central Manchester & Manchester Children's University Hospitals
Manchester
United Kingdom
M13 9WL

Sponsor information

Organisation
Department of Health

Sponsor details
Richmond House
79 Whitehall
London
United Kingdom
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Sponsor type
Government

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

Funder(s)

Funder type
Government

Funder Name
Central Manchester and Manchester Children's University Hospitals NHS Trust (UK)

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