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Prospective randomised controlled study to evaluate the role of synbiotic cocktail 2000 before, during and after liver transplantation to increase resistance to infection

Submission date 12/09/2003	**Recruitment status** Stopped	[ ] Prospectively registered [ ] Protocol
Registration date 12/09/2003	**Overall study status** Stopped	[ ] Statistical analysis plan [ ] Results
Last Edited 24/05/2012	**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
N0256114034

Study information

Scientific Title

Study objectives
Whether giving pre and probiotics could decrease the incidence of infective complications in patients having liver transplantation.

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
Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied
Surgery: Liver transplant

Interventions
Prospective randomised controlled trial.

Intervention Type
Procedure/Surgery

Phase
Not Specified

Primary outcome measure

Service outcomes development.

Secondary outcome measures
Not provided at time of registration

Overall study start date
22/07/2002

Completion date
01/06/2004

Reason abandoned (if study stopped)
Lack of funding

Eligibility

Key inclusion criteria
120 Patients and 40 controls.

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
160

Key exclusion criteria
Does not match inclusion criteria

Date of first enrolment
22/07/2002

Date of final enrolment
01/06/2004

Locations

Countries of recruitment
England

United Kingdom

Study participating centre

University Department of Surgery
London
United Kingdom
NW3 2QG

Sponsor information

Organisation
Department of Health (UK)

Sponsor details
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

Website
http://www.doh.gov.uk

Funder(s)

Funder type
Hospital/treatment centre

Funder Name
The Royal Free Hampstead 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