The effect of probiotic therapy on nosocomial infections in the critically ill

Submission date	Recruitment status	Prospectively registered
30/09/2004	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
30/09/2004	Completed	☐ Results
Last Edited	Edited Condition category	Individual participant data
18/10/2017	Infections and Infestations	Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Dr D Knight

Contact details

Department of Anaesthesia and Intensive Care Queens Medical Centre University Hospital Nottingham United Kingdom NG7 2UH

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers N0192133923

Study information

Scientific Title

The effect of probiotic therapy on nosocomial infections in the critically ill

Study objectives

Does early enteral feeding with probiotics reduce the rate of nosocomial (hospital acquired) infections in the critically ill?

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

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

Health condition(s) or problem(s) studied

Infections and Infestations: Nosocomial infections

Interventions

- 1. Lactobacillus
- 2. Placebo

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

Probiotics

Primary outcome measure

Oropharyngeal nosocomial bacterial carriage rates, nosocomial infection rates, death rates, diarrhoea rates.

Secondary outcome measures

Not provided at time of registration

Overall study start date

07/10/2003

Completion date

31/12/2005

Eligibility

Key inclusion criteria

Not provided at time of registration

Participant type(s)

Patient

Age group

Not Specified

Sex

Both

Target number of participants

Not provided at time of registration

Key exclusion criteria

Does not meet inclusion criteria

Date of first enrolment

07/10/2003

Date of final enrolment

31/12/2005

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Department of Anaesthesia and Intensive Care

Nottingham United Kingdom NG7 2UH

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House 79 Whitehall London United Kingdom SW1A 2NL

Sponsor type

Government

Website

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

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Nottingham 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