

Randomised controlled trial of prophylactic antibiotics at the time of oocyte retrieval and their impact on incidence of bacterial contamination of the catheter used for embryo transfer, embryo implantation and pregnancy rates.

Submission date 30/09/2005	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 30/09/2005	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 01/07/2010	Condition category Pregnancy and Childbirth	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0013146096

Study information

Scientific Title

Study objectives

Impact of prophylactic antibiotics at the time of oocyte retrieval on incidence of bacterial contamination of the catheter used for embryo transfer, embryo implantation and pregnancy rates.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Added 25/06/10:

Received from South East London REC 1 (formerly Guy's Research Ethics Committee)

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

Embryo implantation

Interventions

Randomised controlled trial of patients into antibiotics or no treatment.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Incidence of bacterial contamination of mock catheter and embryo transfer catheter, embryo implantation and pregnancy rates.

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/03/2004

Completion date

01/05/2004

Eligibility

Key inclusion criteria

Patients undergoing embryo transfer without any indication of pelvic infection.

Participant type(s)

Patient

Age group

Adult

Sex

Female

Target number of participants

350

Key exclusion criteria

1. Patients allergic to penicillin
2. Patients previously in trial
3. Patients who required antibiotics at the time of egg collection

Date of first enrolment

01/03/2004

Date of final enrolment

01/05/2004

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Guy's & St Thomas' NHS Foundation Trust

London

United Kingdom

SE1 9RT

Sponsor information

Organisation

Department of Health

Sponsor details

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

Guy's and St. Thomas' NHS Foundation Trust (UK) - 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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/11/2006		Yes	No