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Assessment of patients' satisfaction with regard to transvaginal scanning position during ovarian stimulation protocol

Submission date	Recruitment status	[ ] Prospectively registered
29/09/2006	No longer recruiting	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
29/09/2006	Completed	[ ] Results
Last Edited	**Condition category**	[ ] Individual participant data
23/04/2015	Pregnancy and Childbirth	[ ] 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
N0128106659

Study information

Scientific Title
Assessment of patients' satisfaction with regard to transvaginal scanning position during ovarian stimulation protocol

Study objectives
To find out the best possible and comfortable position for the patient during transvaginal scanning.

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Study design
Randomised prospective crossover observational study

Primary study design
Interventional

Secondary study design
Randomised cross over trial

Study setting(s)
Not specified

Study type(s)
Treatment

Participant information sheet

Health condition(s) or problem(s) studied
Pregnancy and Childbirth: IVF treatment

Interventions
Interventions: assessment by questionnaire of two different positions during transvaginal scanning:
1. Supine position with pillows under bottom
2. Buttocks on the end of the bed with feet on a chair

Intervention Type
Other

Phase
Not Specified

Primary outcome measure

Comparison of patients' satisfaction regarding the two positions commonly used in transvaginal scanning.

Secondary outcome measures
Not provided at time of registration

Overall study start date
08/09/2001

Completion date
08/10/2004

Eligibility

Key inclusion criteria
1. 80 patients age 20-40, undergoing IVF treatment
2. Any patient for baseline and follicular tracking scan

Participant type(s)
Patient

Age group
Adult

Sex
Female

Target number of participants
80

Key exclusion criteria
Any patient with physical disability which might affect normal position.

Date of first enrolment
08/09/2001

Date of final enrolment
08/10/2004

Locations

Countries of recruitment
England

United Kingdom

Study participating centre

Liverpool Women's Hospital
Liverpool
United Kingdom
L8 7SS

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
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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
Liverpool Women's Hospital 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