

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

Submission date 29/09/2006	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 29/09/2006	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 23/04/2015	Condition category Pregnancy and Childbirth	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

Protocol serial number

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

Study type(s)

Treatment

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(s)

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

Key secondary outcome(s)

Not provided at time of registration

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

Healthy volunteers allowed

No

Age group

Adult

Sex

Female

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**

United Kingdom

England

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

Funder(s)**Funder type**

Government

Funder Name

Liverpool Women's Hospital NHS Trust (Uk)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration