# Efficacy of double intrauterine insemination in controlled ovarian hyperstimulation cycle

Submission date	Recruitment status	<ul><li>Prospectively registered</li></ul>
12/09/2003	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
12/09/2003	Completed	Results
Last Edited	Condition category	Individual participant data
19/05/2017	Pregnancy and Childbirth	Record updated in last year

### Plain English summary of protocol

Not provided at time of registration

### Contact information

### Type(s)

Scientific

#### Contact name

Dr Nikita Rawal

#### Contact details

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# Additional identifiers

EudraCT/CTIS number

**IRAS** number

ClinicalTrials.gov number

Secondary identifying numbers

N0072119579

# Study information

### Scientific Title

Efficacy of double intrauterine insemination in controlled ovarian hyperstimulation cycle

### Study objectives

Not provided at time of registration

### 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

### Health condition(s) or problem(s) studied

Double intrauterine insemination (IUI)

#### Interventions

Randomised controlled trial. Women in group A will have IUI 24 hours after human chorionic gonadotropin (HCG) injection. Group B will have IUI twice at 12 and 36 hours after HCG injection.

### Intervention Type

Other

### Phase

Not Applicable

### Primary outcome measure

Not provided at time of registration

### Secondary outcome measures

Not provided at time of registration

### Overall study start date

01/01/2003

### Completion date

31/12/2003

# Eligibility

### Key inclusion criteria

Couples with unexplained fertility of at least 2 years.

### Participant type(s)

**Patient** 

### Age group

Adult

### Sex

Female

### Target number of participants

Not provided at time of registration

### Key exclusion criteria

Not provided at time of registration

### Date of first enrolment

01/01/2003

### Date of final enrolment

31/12/2003

# Locations

### Countries of recruitment

England

**United Kingdom** 

## Study participating centre

**Countess of Chester NHS Foundation Trust** 

Chester United Kingdom CH2 1UL

# 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

Government

### **Funder Name**

Countess of Chester Hospital NHS Foundation 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