

# Efficacy of double intrauterine insemination in controlled ovarian hyperstimulation cycle

<b>Submission date</b> 12/09/2003	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
<b>Registration date</b> 12/09/2003	<b>Overall study status</b> Completed	<input type="checkbox"/> Protocol
<b>Last Edited</b> 19/05/2017	<b>Condition category</b> 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

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## Additional identifiers

**Protocol serial number**  
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

**Study type(s)**

Treatment

**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(s)**

Not provided at time of registration

**Key secondary outcome(s)**

Not provided at time of registration

**Completion date**

31/12/2003

**Eligibility**

**Key inclusion criteria**

Couples with unexplained fertility of at least 2 years.

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Adult

**Sex**

Female

**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**

United Kingdom

England

**Study participating centre**

Countess of Chester NHS Foundation Trust

Chester

United Kingdom

CH2 1UL

**Sponsor information****Organisation**

Department of Health (UK)

**Funder(s)****Funder type**

Government

**Funder Name**

Countess of Chester Hospital NHS Foundation Trust (UK)

# Results and Publications

Individual participant data (IPD) sharing plan

**IPD sharing plan summary**

Not provided at time of registration