

Scottish Unexplained Infertility Trial: a randomised trial of clomiphene versus intrauterine insemination for unexplained infertility

Submission date 16/03/2005	Recruitment status No longer recruiting	☐ Prospectively registered
Registration date 01/09/2005	Overall study status Completed	☐ Protocol
Last Edited 30/10/2012	Condition category Pregnancy and Childbirth	☐ Statistical analysis plan
		☒ Results
		☐ 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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Contact details

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Additional identifiers

Protocol serial number

CZH/4/17

Study information

Scientific Title

Acronym

SUIT

Study objectives

To assess the efficacy of clomiphene citrate versus intrauterine insemination (IUI) versus no treatment using the partner's semen sample in the treatment of unexplained infertility.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Primary study design

Interventional

Study design

Randomised controlled trial

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Infertility

Interventions

Clomifene citrate versus intrauterine insemination versus expectant management.

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

Clomifene citrate

Primary outcome(s)

To compare the three treatments in terms of efficacy, patient acceptability and costs.

Key secondary outcome(s)

Not provided at time of registration

Completion date

31/08/2006

Eligibility

Key inclusion criteria

570 couples with unexplained infertility. Those with infertility of less than 2 years or females aged above 38 years will be excluded.

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

01/09/2001

Date of final enrolment

31/08/2006

Locations**Countries of recruitment**

United Kingdom

Scotland

Study participating centre

Dept. of Obstetrics & Gynaecology

Aberdeen

United Kingdom

AB25 2ZD

Sponsor information**Organisation**

Individual Sponsor (UK)

Funder(s)

Funder type

Government

Funder Name

Chief Scientist Office of the Scottish Executive Health Department (UK) (ref: CZH/4/17)

Results and Publications

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	07/08/2008		Yes	No
Results article	results	01/02/2011		Yes	No