Comparison between two commercially available embryo culture media

| Submission date | Recruitment status | Prospectively registered |
|-------------------|---------------------------------|--|
| 03/02/2009 | No longer recruiting | Protocol |
| Registration date | Overall study status | Statistical analysis plan |
| 13/03/2009 | Completed | Results |
| Last Edited | Condition category | Individual participant data |
| 13/03/2009 | Urological and Genital Diseases | 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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers N/A

Study information

Scientific Title

A prospective randomised sibling-oocyte study of two sequential media for culturing cleavage stage embryos

Study objectives

Commercially available culture media are expected to be equally efficient in fertilisation and culture of early embryos.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Local ethics committee (Örebro läns landsting) (Sweden) gave approval on the 25th March 2003 (ref: 90/03)

Study design

Prospective randomised single centre single-blinded (patients only) sibling study

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Treatment

Participant information sheet

Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Infertility

Interventions

The study included two commercially available sequential media for fertilisation and culture of oocytes. One of the media system was standard in our clinic, while the other media system was standard in the private clinic in our town. It was not expected that there would be a difference.

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

- 1. Fertilisation rate, measured at 1 day
- 2. Polynuclei rate, measured at 2 days
- 3. Cleavage rate, measured at 2 days

Secondary outcome measures

- 1. Positive human chorionic gonadotrophin (hCG) rate, measured at 1 2 weeks.
- 2. Clinical pregnancy rate, measured at 12 weeks
- 3. Implantation rate, measured at 12 weeks
- 4. Delivery rate, measured at 9 months

Overall study start date

01/10/2007

Completion date

31/12/2007

Eligibility

Key inclusion criteria

All oocytes from patients (aged 24 - 40 years, female) undergoing in vitro fertilisation (IVF) treatment

Participant type(s)

Patient

Age group

Adult

Sex

Female

Target number of participants

110 patients/1200 oocytes

Key exclusion criteria

Does not meet inclusion criteria

Date of first enrolment

01/10/2007

Date of final enrolment

31/12/2007

Locations

Countries of recruitment

Sweden

Study participating centre Department of Women's and Children's Health Uppsala Sweden 751 85

Sponsor information

Organisation

Uppsala County Council (Uppsala Läns Landsting) (Sweden)

Sponsor details

Kvinno och Barndivisionen Akademiska sjukhuset Uppsala Sweden 751 85

Sponsor type

Government

Website

http://www.lul.se/

ROR

https://ror.org/01dv86r63

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Uppsala University Hospital (Sweden) - Centre of Reproduction

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 summaryNot provided at time of registration