The enigma of unexplained subfertility

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Prevention of multiple pregnancies in couples with unexplained or mild male subfertility: randomised controlled trial of in vitro fertilisation with single embryo transfer, or in vitro fertilisation in a modified natural cycle compared with intrauterine insemination with controlled ovarian hyperstimulation


ABSTRACT

Objectives: To compare the effectiveness of in vitro fertilisation with single embryo transfer or in vitro fertilisation in a modified natural cycle to intrauterine insemination with controlled ovarian hyperstimulation in terms of a healthy child.

Design: Multicentre open-label three arm parallel-group randomised controlled non-inferiority trial.

Setting: 17 centres in the Netherlands.

Participants: Couples seeking fertility treatment after at least 12 months of unprotected intercourse, with the female partner aged between 18 and 38 years, an unfavourable prognosis for natural conception and a diagnosis of unexplained or mild male subfertility.

Interventions: Three cycles of in vitro fertilisation with single embryo transfer (plus subsequent cryocycles), six cycles of in vitro fertilisation in a modified natural cycle and six cycles of intrauterine insemination with ovarian hyperstimulation within 12 months after randomisation.

Main outcome measures: The primary outcome was birth of a healthy child resulting from a singleton pregnancy conceived within 12 months after randomisation. Secondary outcomes were live birth, clinical pregnancy, ongoing pregnancy, multiple pregnancy, time to pregnancy, pregnancy complications, neonatal morbidity and mortality.

Results: 602 couples were randomly assigned between January 2009 and February 2012, with 201 allocated to in vitro fertilisation with single embryo transfer, 194 to in vitro fertilisation in a modified natural cycle, and 207 to intrauterine insemination with controlled ovarian hyperstimulation. Birth of a healthy child occurred in 104 couples (52%) in the in vitro fertilisation with single embryo transfer group, 83 (43%) in the in vitro fertilisation in a modified natural cycle group, and 97 (47%) in the intrauterine insemination with controlled ovarian hyperstimulation group. This corresponds to a risk, relative
In vitro fertilisation versus intrauterine insemination

to intrauterine insemination with ovarian hyperstimulation, of 1.10 for in vitro fertilisation with single embryo transfer (95% CI: 0.91 to 1.34) and of 0.91 for in vitro fertilisation in a modified natural cycle (95% CI: 0.73 to 1.14). These two 95% confidence intervals do not extend below the predefined threshold of 0.69 for inferiority. Multiple pregnancy rates per ongoing pregnancy were 6 % after in vitro fertilisation with single embryo transfer, 5% after in vitro fertilisation in a modified natural cycle and 7% after intrauterine insemination with ovarian hyperstimulation (one sided p-value=0.52 for in vitro fertilisation with single embryo transfer compared to intrauterine insemination with ovarian hyperstimulation and one sided p-value=0.33 for in vitro fertilisation in a modified natural cycle compared to intrauterine insemination with controlled ovarian hyperstimulation).

**Conclusions:** In vitro fertilisation with single embryo transfer and in vitro fertilisation in a modified natural cycle were non-inferior to intrauterine insemination with controlled ovarian hyperstimulation in terms of the birth of a healthy child and showed comparable, low multiple pregnancy rates.

**Trial registration:** The trial was registered (NTR 939, ISRCTN 52843371).
INTRODUCTION

Involuntary childlessness currently affects more than 70 million couples worldwide and has a large impact on the quality of life leading to lasting psychosocial effects (1,2). Most couples will seek fertility care and consult their general practitioner for guidance. Approximately half of these couples will be diagnosed as having unexplained or mild male subfertility (3). Most of them still have a good chance of conceiving, and will achieve pregnancy without treatment (4,5). Treatment is thus indicated only if the chances to conceive naturally are low and the success rate after treatment exceeds this probability (6,7).

Intrauterine insemination with controlled ovarian hyperstimulation is the first line fertility treatment in couples with an unfavourable prognosis of natural conception (8,9). However, concern exists about the increased rates of multiple pregnancy after intrauterine insemination with controlled ovarian hyperstimulation, as a result of the stimulation of multiple follicles (10). Intrauterine insemination with controlled ovarian hyperstimulation results in pregnancy rates of 13% per cycle at the expense of a multiple pregnancy rate that is estimated to be above 10% per ongoing pregnancy (6,11). Multiple pregnancies are associated with maternal morbidity such as pre-eclampsia, gestational diabetes and a 50% risk of premature delivery, resulting in considerable neonatal morbidity and mortality (12).

In vitro fertilisation with single embryo transfer is increasingly implemented, and might be equally effective, but safer as it prevents multiple pregnancies (13-15). Live birth rates with the transfer of one good quality embryo are 28% per cycle, while multiple pregnancy rates are as low as 1% per live birth (16). The downside of in vitro fertilisation is that it is an invasive and burdensome procedure with higher costs (17).

In vitro fertilisation in a modified natural cycle - a cycle in which monofollicular growth results in one oocyte at follicular aspiration, and in one embryo after fertilisation - might be another treatment option. In vitro fertilisation with unstimulated or mildly stimulated regimens are becoming more popular (18-21). Cumulative live birth rates over six cycles of more than 30% have been reported with almost no multiple pregnancies (22). In vitro fertilisation in a modified natural cycle has been advocated as less burdensome and less costly than “conventional” in vitro fertilisation (22,23).
Multiple pregnancies seem to be prevented by in vitro fertilisation with single embryo transfer and in vitro fertilisation in a modified natural cycle, but this will be acceptable only if these interventions are as effective as intrauterine insemination with controlled ovarian hyperstimulation in terms of live birth rates. We therefore designed a randomised trial to evaluate the effectiveness of these interventions in terms of healthy children born from singleton pregnancies.

METHODS

We did a multicentre, open-label, three arm parallel-group randomised controlled non-inferiority trial in 17 fertility clinics in the Netherlands between January 2009 and February 2013. The study protocol was approved by the Medical Ethical Committee of the Academic Medical Centre in Amsterdam (MEC 07-064) and the boards of directors of each participating centre (24). Full details of the trial protocol can be found on www.studies-obsgyn.nl/ines.

Participants
couples seeking fertility treatment after at least 12 months of unprotected intercourse were eligible. All participating couples provided written informed consent. All couples underwent basic fertility investigations, which included semen analysis, evaluation of ovulation and tubal patency testing (Chlamydia antibody test, hysterosalpingography or laparoscopy). Inclusion criteria were age of the female partner between 18 and 38 years, an unfavourable prognosis for natural conception and a diagnosis of unexplained or mild male subfertility. We classified couples as having unexplained subfertility when the fertility investigations showed at least one patent fallopian tube, an ovulatory menstrual cycle, and a normal semen analysis (pre-wash total motile sperm count above 10 million).(24) We considered couples who qualified for intrauterine insemination with donor sperm after at least six cycles of artificial intracervical insemination with donor sperm to have unexplained subfertility for the purpose of this study. Mild male subfertility was diagnosed when the semen analysis showed a pre-wash total motile sperm count between three and 10 million (according to the Dutch guidelines). We defined an unfavourable prognosis for natural conception as a probability of natural conception within
the next twelve 12 of less than 30%, as calculated with the validated synthesis model of Hunault (7). This model encompasses female age, duration of subfertility, whether infertility is primary or secondary, percentage of motile progressive sperm and referral status. It is readily available for all clinicians (http://www.freya.nl/web_bereken/bereken.php). Exclusion criteria were anovulation, double sided tubal disease, severe endometriosis, premature ovarian failure, and known endocrine disorders (such as Cushing syndrome or adrenal hyperplasia).

Randomisation and masking
Couples were randomly allocated in a 1:1:1 ratio to receive either three consecutive cycles of in vitro fertilisation with single embryo transfer plus subsequent cryocycles, six consecutive cycles of in vitro fertilisation in a modified natural cycle, or six consecutive cycles intrauterine insemination with controlled ovarian hyperstimulation within 12 months. Randomisation was performed with an online randomisation program, using biased coin minimisation, stratified for study centre. A web-based program generated a unique number with allocation code after entering patient’s initials and date of birth. Neither the recruiters nor the trial project group could access the randomisation sequence. Blinding was not possible owing to the nature of the interventions.

Interventions
In the in vitro fertilisation with single embryo transfer group, participating hospitals could adhere to local stimulation protocols, which were either long/short agonist or antagonist protocols. Controlled ovarian hyperstimulation started with 150 IU follicle-stimulating hormone. Treatment was continued until at least two follicles of at least 18 mm had developed. Ovulation triggering was induced by 10,000 IU human chorionic gonadotropin hormone (Pregnyl®, Merck Sharp & Dohme) and cumulus-oocyte complexes were recovered by transvaginal ultrasound-guided retrieval 36 hours thereafter. We adhered to an elective single embryo transfer policy: if one good quality embryo was available, we transferred one embryo (25). If more than one good quality embryo was available, suitable surplus embryos were cryopreserved. If no good quality embryos were available, two embryos would be transferred. For the morphological score the degree of fragmentation of the embryo and
the uniformity of the blastomeres were assessed daily (26). The embryo was
given a score of 1 (no fragmentation), 2 (<20% fragmentation), 3 (20-50% 
fragmentation) or 4 (> 50% fragmentation). We defined good quality embryos
as embryos having a cumulative embryo score of 24 or higher. Embryo transfer
followed on day three. All available frozen embryos were transferred after
thawing before a new treatment cycle was started.

During our trial the results of a pilot study, randomising women to three cycles
of intrauterine insemination with controlled ovarian hyperstimulation or one
cycle of in vitro fertilisation with single embryo transfer, was published. This
pilot study showed that the policy of transferring two embryos when no good
quality embryos are available is not effective in preventing multiple pregnancies
(27). We amended our study protocol and from February 2010, after allocation
of 48 women to the in vitro fertilisation with single embryo transfer group, a
strict single embryo transfer policy was implemented (that is, single embryo
transfer was performed irrespective of the quality of the embryo).

In the in vitro fertilisation in a modified natural cycle group, women were
monitored by transvaginal ultrasound from cycle days eight to 10 onward
and when the lead follicle had a mean diameter of at least 14 mm, they were
started on daily injections of 0.25 mg of the gonadotropin-releasing hormone
antagonist to prevent premature ovulation together with 150 IU follicle-
stimulating hormone to prevent collapse of the follicle and a concomitant fall
in oestradiol concentrations. Follicle-stimulating hormone was continued up to
the day of the ovulation triggering, and the gonadotropin-releasing hormone
antagonist was last given on the day of ovulation triggering. When a follicle with
a diameter of 17 to 18 mm was observed, ovulation triggering was achieved
by subcutaneous injection of 10 000 IU of human chorionic gonadotropin.
Oocyte retrieval was planned 34 hours thereafter and was performed without
anaesthesia or sedation. If an oocyte was obtained and fertilised, transfer of
the embryo was performed on day three. For luteal support, human chorionic
gonadotropin 1500 IU was given by subcutaneous injections on days five, eight
and 11 after oocyte retrieval. The next treatment cycle could start immediately
after the previous cycle.

In the intrauterine insemination with controlled ovarian hyperstimulation
group, women received controlled ovarian hyperstimulation according
to local protocol, with either 100 mg clomiphene citrate (cycle day three to
seven) or daily subcutaneous injections of 75 IU follicle stimulating hormone
(starting dose). Follicular growth was monitored by ultrasound and when at least one follicle of 17 or 18 mm was present, final oocyte maturation was induced by the administration of 5,000 IU human chorionic gonadotropin or 6,500 IE of recombinant human chorionic gonadotropin. Approximately 36 hours thereafter IUI was performed. Intrauterine insemination with controlled ovarian hyperstimulation cycles were cancelled when there were more than three follicles with a diameter of 16 mm, or more than five follicles with a diameter of 12 mm. Those couples were instructed to refrain from unprotected intercourse. The box summarizes the three interventions.

Box describing three interventions

<table>
<thead>
<tr>
<th>IUI-COH</th>
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<tbody>
<tr>
<td><strong>Hyperstimulation from cycle day 3 or 4 start with 100 mg clomiphene citrate or subcutaneous injections of 75 IU FSH</strong></td>
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<tr>
<td><strong>Monitoring of follicular growth by transvaginal ultrasound.</strong></td>
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<tr>
<td><strong>Induction of final oocyte maturation with 5,000 IU of hCG when ≥ 1 follicle with diameter of 17 or 18 mm</strong></td>
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</tr>
<tr>
<td><strong>IUI 36 hours thereafter.</strong></td>
<td></td>
</tr>
<tr>
<td><strong>Cancel criteria</strong></td>
<td></td>
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<tr>
<td>HCG administration and IUI will be withheld when ≥ 3 follicles with a diameter of 16 mm, or &gt; 5 follicles with a diameter of 12 mm.</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>MNC-IVF</th>
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</tr>
</thead>
<tbody>
<tr>
<td><strong>Ultrasound monitoring from cycle day 8-10 with monitoring serum concentrations of LH and E2</strong></td>
<td></td>
</tr>
<tr>
<td><strong>Start daily injections of 0.25 mg of a GnRH-antagonist with 150 IU FSH when a lead follicle with a mean diameter of ≥14 mm</strong></td>
<td></td>
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<tr>
<td><strong>Ovulation induction with 10,000 IU hCG when 1 follicle with a diameter of 17 to 18 mm or LH &gt; 30 with follicle&gt; 15mm</strong></td>
<td></td>
</tr>
<tr>
<td><strong>Oocyte retrieval 34 hours thereafter.</strong></td>
<td></td>
</tr>
<tr>
<td><strong>Embryo transfer day 2 or 3 day after oocyte retrieval. luteal support, HCG 1500 IU on day 5, 8 and 11</strong></td>
<td></td>
</tr>
<tr>
<td><strong>Cancel criteria</strong></td>
<td></td>
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<tr>
<td>LH-rise of &gt; 30.0 IU/L is noticed at a follicle size of &lt; 15 mm.</td>
<td></td>
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</tbody>
</table>

<table>
<thead>
<tr>
<th>IVF-SET</th>
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</thead>
<tbody>
<tr>
<td><strong>Down-regulation with a GnRH agonist in long protocol or fixed start antagonist protocol. Stimulation start dose 150 IU FSH.</strong></td>
<td></td>
</tr>
<tr>
<td><strong>Ultrasound monitoring according to local protocol</strong></td>
<td></td>
</tr>
<tr>
<td><strong>Ovulation induction with 10,000 IU hCG until ≥2 follicles &gt; 18 mm</strong></td>
<td></td>
</tr>
<tr>
<td><strong>Oocyte retrieval 36 hours thereafter.</strong></td>
<td></td>
</tr>
<tr>
<td><strong>Embryo transfer day 2, 3 or 4.</strong></td>
<td></td>
</tr>
<tr>
<td><strong>Cryopreservation of non-transferred good quality embryos will be cryo-preserved. (one embryo will be transferred per frozen-thaw cycle if it is of good quality)</strong></td>
<td></td>
</tr>
<tr>
<td><strong>Cancel criteria</strong></td>
<td></td>
</tr>
<tr>
<td>OHSS, non response</td>
<td></td>
</tr>
</tbody>
</table>

IVF-SET: in vitro fertilisation with single embryo transfer; IVF-MNC: in vitro fertilisation in a modified natural cycle; IUI-COH: intrauterine insemination with controlled ovarian hyperstimulation

FSH= follicle stimulating hormone, hCG = human chorionic gonadotropin, GnRH = Gonadotropin-releasing hormone, OHSS= ovarian hyperstimulation syndrome
**Follow-up**
Couples were followed up for 12 months after randomisation. We included all interventions that couples received within 12 months after randomisation. A pregnancy test was performed two weeks after embryo transfer or intrauterine insemination. Clinical and ongoing pregnancies were confirmed by ultrasonography. If a miscarriage occurred within 12 months after randomisation couples could continue their allocated treatment. Natural conceptions were also included in the analysis in the three groups.

**Outcomes**
The primary outcome was the birth of a healthy child, resulting from a singleton pregnancy conceived within 12 months after randomisation. A child was considered healthy when born at term, defined as a gestational age between 37 and 42 weeks, with a birth weight above the fifth percentile, according to the Dutch reference curves corrected for parity, sex and ethnicity, without congenital anomalies, and developing normally up to six weeks after birth (28). This information was collected from the parents, child health centres, or paediatricians.

Secondary outcomes were any live birth, multiple pregnancy (registered heartbeat of at least two foetuses at 12 weeks of gestation), clinical pregnancy (any registered embryonic heartbeat on ultrasound), ongoing pregnancy (registered heartbeat of a foetus at 12 weeks of gestation), neonatal mortality (death of the child within 28 days after birth), neonatal morbidity (preterm birth <37 weeks, birth weight <2500 grams), pregnancy complications, (pregnancy induced hypertension, (pre-) eclampsia, HELLP syndrome), and time to pregnancy.

All serious adverse events were reported to the trial coordinator. An independent data safety and monitoring committee (consisting of two gynaecologists and one statistician) monitored the progress of the study and the safety of the women and performed a blinded interim analysis to exclude large differences after 300 inclusions with ongoing pregnancy as a surrogate outcome.

**Study design, sample size, and statistical analysis**
The trial was designed as a non-inferiority trial. The null hypothesis assumed that the pregnancy rates between all interventions would be comparable. We expected 40% of the couples to have a live birth within 12 months after starting
with intrauterine insemination with controlled ovarian hyperstimulation (29). For the power calculation, using a 5% significance level, we needed 190 couples to achieve 80% power to exclude a difference of 12.5% or more to the detriment of in vitro fertilisation with single embryo transfer and in vitro fertilisation in a modified natural cycle. This difference corresponds to a relative risk of 0.69. To account for a 5% loss to follow-up we needed to include 200 couples in each group.

All randomised patients were included in all analyses. We estimated differences in the primary and secondary outcomes as relative risks (RR) with 95% confidence intervals. We constructed Kaplan-Meier survival curves in each treatment group for the time to ongoing pregnancy of a healthy child. We used SPSS (version 20.0) for all statistical analyses. We considered P-values <0.05 as indicating statistically significant differences.

Role of the funding source
The study was supported by a grant from ZonMW, the Dutch Organization for Health Research and Development (120620027), and a grant from Zorgverzekeraars Nederland, the Dutch association of health care insurers (09-003). The funders had no role in study design, collection, analysis or interpretation of the data. The corresponding author had full access to all data and final responsibility for the decision to submit for publication.

RESULTS

Between January 2009 and February 2012, 869 couples fulfilled the inclusion criteria and were invited to the trial, of whom 267 declined randomisation; 602 couples were included: 201 couples assigned to in vitro fertilisation with single embryo transfer, 194 to in vitro fertilisation in a modified natural cycle and 207 to intrauterine insemination with controlled ovarian hyperstimulation (figure 1).
**Figure 1. Flow chart**

![Flow chart]

**Table 1. Baseline characteristics of the couples**

<table>
<thead>
<tr>
<th></th>
<th>IVF-SET (n=201)</th>
<th>IVF-MNC (n=194)</th>
<th>IUI-COH (n=207)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mean (SD) of female partner, years,</td>
<td>33 (3.39)</td>
<td>33 (3.50)</td>
<td>34 (3.67)</td>
</tr>
<tr>
<td>Primary subfertility, N (%)</td>
<td>160 (80)</td>
<td>141 (73)</td>
<td>157 (76)</td>
</tr>
<tr>
<td>Median (IQR), duration of subfertility, years</td>
<td>2.13 (1.73-3.01)</td>
<td>2.14 (1.77-2.81)</td>
<td>2.30 (1.82-3.13)</td>
</tr>
<tr>
<td>White, N (%)</td>
<td>182 (91)</td>
<td>163 (85)</td>
<td>178 (90)</td>
</tr>
<tr>
<td>Smoking, N (%)</td>
<td>45 (23)</td>
<td>35 (18)</td>
<td>46 (22)</td>
</tr>
<tr>
<td>Median (IQR) body mass index, kg/m²</td>
<td>23 (21-26)</td>
<td>23 (21-25)</td>
<td>23 (21-26)</td>
</tr>
<tr>
<td>Median (IQR) total motile sperm count (x 10⁶)</td>
<td>51 (25-100)</td>
<td>53 (25-126)</td>
<td>59 (30-124)</td>
</tr>
<tr>
<td>Diagnosis subfertility N (%):</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Unexplained</td>
<td>183 (91)</td>
<td>173 (89)</td>
<td>189 (91)</td>
</tr>
<tr>
<td>Male factor</td>
<td>18 (9)</td>
<td>21 (11)</td>
<td>18 (9)</td>
</tr>
<tr>
<td>Hunault score; Mean (SD) *</td>
<td>20 (6.56)</td>
<td>21 (6.83)</td>
<td>19 (6.38)</td>
</tr>
</tbody>
</table>

* Hunault score was calculated according to the formula by Steeg et al 2007

Baseline characteristics in the three groups were comparable (table 1).

Tables 2 list all interventions that took place during the 12 months follow up. Table 2a shows the number of couples who received the allocated treatment per cycle, and table 2b shows the total number of cycles within 12 months.
Table 2 also shows additional treatments that couples received when they switched from the allocated treatment to another treatment or when they continued with another treatment after having completed their allocated treatment within 12 months after randomisation also shows additional treatments that couples received when they switched from the allocated treatment to another treatment or when they continued with another treatment after having completed their allocated treatment within 12 months of randomisation. In table 2 the total number of cycles within 12 months are presented.

In the in vitro fertilisation with single embryo transfer group 15% (30/201) of the couples discontinued treatment before receiving three fresh cycles of in vitro fertilisation with single embryo transfer or achieving pregnancy, 23% (45/194) in the in vitro fertilisation in a modified natural cycle group and 13% (26/207) in the intrauterine insemination with controlled ovarian hyperstimulation group. These couples discontinued treatment because they switched to another treatment, or chose to end treatment.

Table 2. Number of couples per intervention and 2b total number fo cycles performed within 12 months

<table>
<thead>
<tr>
<th>Number of couples per intervention</th>
<th>IVF-SET (N=201)</th>
<th>IVF-MNC (N=194)</th>
<th>IUI-COH (N=207)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number of couples per cycle</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1 cycle</td>
<td>177*</td>
<td>166</td>
<td>194</td>
</tr>
<tr>
<td>2 cycles</td>
<td>90*</td>
<td>135</td>
<td>173</td>
</tr>
<tr>
<td>3 cycles</td>
<td>34*</td>
<td>113</td>
<td>145</td>
</tr>
<tr>
<td>4 cycles</td>
<td>-</td>
<td>91</td>
<td>125</td>
</tr>
<tr>
<td>5 cycles</td>
<td>-</td>
<td>72</td>
<td>100</td>
</tr>
<tr>
<td>6 cycles</td>
<td>-</td>
<td>55</td>
<td>85</td>
</tr>
<tr>
<td>Additional treatment within 12 months</td>
<td>15</td>
<td>63</td>
<td>70</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Total number of cycles performed within 12 months</th>
<th>IVF-SET</th>
<th>IVF-MNC</th>
<th>IUI-COH</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number of cycles</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total number of fresh cycles</td>
<td>301</td>
<td>632</td>
<td>822</td>
</tr>
<tr>
<td>Total number of frozen cycles</td>
<td>158</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Total additional fresh IVF/ICSI cycles</td>
<td>5</td>
<td>38</td>
<td>68</td>
</tr>
<tr>
<td>Total additional frozen IVF/ICSI cycles</td>
<td>0</td>
<td>7</td>
<td>17</td>
</tr>
<tr>
<td>Total additional IUI cycles</td>
<td>45</td>
<td>61</td>
<td>31</td>
</tr>
</tbody>
</table>

* IVF-SET: in vitro fertilisation with single embryo transfer; IVF-MNC: in vitro fertilisation in a modified natural cycle; IUI-COH: intrauterine insemination with controlled ovarian hyperstimulation

* Data in the IVF-SET group is presented as the number of fresh cycles started. According to the protocol couples received a maximum of 3 fresh cycles of IVF-SET.
Pregnancy outcomes
Table 3 lists the outcomes of pregnancy. In all, 104 couples (52%) had healthy children after in vitro fertilisation with single embryo transfer, 83 (43%) after in vitro fertilisation in a modified natural cycle, and 97 (47%) after intrauterine insemination with controlled ovarian hyperstimulation. This corresponds to a relative risk, compared to intrauterine insemination with controlled ovarian hyperstimulation, of 1.10 (95%CI: 0.91 to 1.34) for in vitro fertilisation with single embryo transfer and of 0.91 (95%CI: 0.73 to 1.14) for in vitro fertilisation in a modified natural cycle. These two 95% confidence intervals do not extend below the predefined threshold of 0.69 for inferiority.
We found no evidence of a difference between groups in rates of live birth, ongoing pregnancy or clinical pregnancy. Multiple pregnancy rates were 6% after in vitro fertilisation with single embryo transfer, 5% after in vitro fertilisation in a modified natural cycle, and 7% after intrauterine insemination with controlled ovarian hyperstimulation (one sided p-value= 0.52 for in vitro fertilisation with single embryo transfer compared to intrauterine insemination with controlled ovarian hyperstimulation and one sided p-value =0.33 for in vitro fertilisation in a modified natural cycle compared to intrauterine insemination with controlled ovarian hyperstimulation. One triplet occurred in the intrauterine insemination with controlled ovarian hyperstimulation group; all other multiple pregnancies were twin pregnancies.
In the 48 couples who received in vitro fertilisation with single embryo transfer before the amendment of the protocol, 22 women (46%) had a healthy child and three women (6%) had an ongoing multiple pregnancy. In the 153 couples who received strict in vitro fertilisation with single embryo transfer after the amendment of the protocol, 82 (54%) women had a healthy singleton and four (3%) women had twins.

Neonatal outcomes
Neonatal death did not occur in this study. We found no differences between groups in the relative number of preterm babies or in those with low birth weight (<2500 grams) (table 3). Congenital anomalies were reported in 10 cases (table 3). According to the ICD-10 classification (international classification of diseases, 10th revision) one case of an aortic stenosis (Q 23) occurred in the in vitro fertilisation with single embryo transfer group. One case of an urachus fistula (Q 64), one case of Moebius syndrome (Q 87) and two cases
of Down syndrome (Q90) occurred in the in vitro fertilisation in a modified natural cycle group. In the intrauterine insemination with controlled ovarian hyperstimulation group, one case of patent/persistent foramen ovale (Q21), one case of congenital deafness (H90), one case of an umbilical hernia (K42), one case of Down syndrome (Q90) and one case of oesophageal atresia (Q39) occurred.

**Maternal outcomes**
Two women in the in vitro fertilisation with single embryo transfer group and one woman in the intrauterine insemination with controlled ovarian hyperstimulation group developed ovarian hyperstimulation syndrome and all were managed expectantly: two in an outpatient setting and one was admitted for one night. Pregnancy-induced hypertension, pre-eclampsia and HELLP syndrome occurred at similar frequencies in the three groups (table 3).

**Time to pregnancy**
Average time to pregnancy leading to a healthy child was 8.04 months for in vitro fertilisation with single embryo transfer, 8.32 months for in vitro fertilisation in a modified natural cycle, and 8.39 months for intrauterine insemination with controlled ovarian hyperstimulation (figure 2). The differences were not statistically significantly with for in vitro fertilisation with single embryo transfer (log-rank: \( p=0.38 \)) or for in vitro fertilisation in a modified natural cycle (log-rank: \( p=0.59 \)) compared with intrauterine insemination with controlled ovarian hyperstimulation.
<table>
<thead>
<tr>
<th>Pregnancy outcomes</th>
<th>IVF-SET (N=201)</th>
<th>IVF-MNC (N=194)</th>
<th>IUI-COH (N=207)</th>
<th>IVF-SET vs IUI-COH RR (95% CI)</th>
<th>IVF-MNC vs IUI-COH RR (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Healthy child</td>
<td>104 (52)</td>
<td>83 (43)</td>
<td>97 (47)</td>
<td>1.10 (0.91 to 1.34)</td>
<td>0.91 (0.73 to 1.14)</td>
</tr>
<tr>
<td>Live birth*</td>
<td>118 (59)</td>
<td>99 (51)</td>
<td>116 (56)</td>
<td>1.05 (0.89 to 1.24)</td>
<td>0.91 (0.76 to 1.09)</td>
</tr>
<tr>
<td>Ongoing pregnancy</td>
<td>121 (60)</td>
<td>102 (53)</td>
<td>119 (57)</td>
<td>1.05 (0.89 to 1.23)</td>
<td>0.91 (0.77 to 1.09)</td>
</tr>
<tr>
<td>Clinical pregnancy</td>
<td>144 (72)</td>
<td>121 (62)</td>
<td>135 (65)</td>
<td>1.10 (0.96 to 1.25)</td>
<td>0.96 (0.82 to 1.11)</td>
</tr>
<tr>
<td>Multiple pregnancy*</td>
<td>7 (6)</td>
<td>5 (5)</td>
<td>8 (7)</td>
<td>0.86 (0.32 to 2.30)</td>
<td>0.73 (0.25 to 2.16)</td>
</tr>
<tr>
<td>Neonatal outcomes***</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Neonatal mortality</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Birth weight &lt; 2500 grams</td>
<td>9 (8)</td>
<td>6 (6)</td>
<td>8 (7)</td>
<td>1.11 (0.44 to 2.77)</td>
<td>0.88 (0.32 to 2.45)</td>
</tr>
<tr>
<td>Preterm birth &lt; 37 weeks</td>
<td>8 (7)</td>
<td>10 (10)</td>
<td>13 (11)</td>
<td>0.60 (0.26 to 1.40)</td>
<td>0.90 (0.41 to 1.97)</td>
</tr>
<tr>
<td>Congenital anomalies</td>
<td>1 (1)</td>
<td>4 (4)</td>
<td>5 (4)</td>
<td>0.20 (0.02 to 1.66)</td>
<td>0.94 (0.26 to 3.40)</td>
</tr>
<tr>
<td>Maternal outcomes</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ovarian hyperstimulation</td>
<td>2 (1)</td>
<td>0</td>
<td>1 (0)</td>
<td>2.06 (0.19 to 2.54)</td>
<td>-</td>
</tr>
<tr>
<td>Pregnancy induced hypertension</td>
<td>1 (1)</td>
<td>4 (2)</td>
<td>4 (2)</td>
<td>0.26 (0.03 to 2.28)</td>
<td>1.07 (0.27 to 4.21)</td>
</tr>
<tr>
<td>Pre-eclampsia</td>
<td>2 (1)</td>
<td>6 (3)</td>
<td>7 (3)</td>
<td>0.29 (0.06 to 1.40)</td>
<td>0.91 (0.31 to 2.67)</td>
</tr>
<tr>
<td>HELLP syndrome</td>
<td>0</td>
<td>0</td>
<td>1 (0)</td>
<td>-</td>
<td>-</td>
</tr>
</tbody>
</table>

Values are numbers (percentages) unless stated otherwise
* One couple in the IVF-SET group with an ongoing pregnancy could not be contacted for follow-up of the pregnancy
** Percentage is a ratio of total ongoing pregnancies. Three monochorial twins: two in the IVF-MNC group and one in the IUI-COH group.
*** Percentage is a ratio of total live births.

Cumulative chances of ongoing pregnancy. IUI-COH=intrauterine insemination with controlled ovarian hyperstimulation; IVF-MNC=in vitro fertilisation in modified natural cycle; IVF-SET=in vitro fertilisation with single embryo transfer
Figure 3 shows time to ongoing pregnancy within 12 months after randomisation, classified as to whether the pregnancy was achieved with the allocated treatment, with additional treatment, or after natural conception. Twenty four (12%) couples had a live birth after natural conception in the in vitro fertilisation with single embryo transfer group, 30 (15%) in vitro fertilisation in a modified natural cycle group and 22 (11%) in the intrauterine insemination with controlled ovarian hyperstimulation group. Of the pregnancies achieved through allocated treatment in the in vitro fertilisation with single embryo transfer group, 68 (76%) were ongoing pregnancies after fresh embryo transfer and 21 (24%) after frozen embryo transfer.

DISCUSSION

In this multicentre, open-label, three arm parallel-group randomised controlled non-inferiority trial we showed that in vitro fertilisation with single embryo transfer and in vitro fertilisation in a modified natural cycle are non-inferior to intrauterine insemination with controlled ovarian hyperstimulation in terms of the birth of a healthy child in couples with unexplained or mild male subfertility and unfavourable fertility prospects. Rates of multiple pregnancy in all three treatment arms were comparable and amounted to less than 7%. Also, time to pregnancy was comparable in all arms.

Strengths and weaknesses of study
Couples were treated with three cycles of in vitro fertilisation with single embryo transfer, six cycles of in vitro fertilisation in a modified natural cycle or six cycles of intrauterine insemination with controlled ovarian hyperstimulation during 12 months to determine the chances of pregnancy following treatment over a realistic time period, reflecting daily clinical practice. Also, all additional interventions or pregnancies achieved through natural conception within the 12 months after randomisation were registered and included in the analysis. The trial was further strengthened by our primary outcome, the birth of a single healthy child. The European Society for Human Reproduction and Embryology (ESHRE) has recommended this outcome measure as the primary aim of reproductive medicine is to help couples with an unfulfilled wish for a child to have a healthy child (30). Our criteria for a healthy child
were quite strict, but if these were softened to include healthy late-pre term infants, more children born out of a twin pregnancy in our study would be considered healthy.

We initially chose elective single embryo transfer on the basis of data from a meta-analysis that showed that elective single embryo transfer in patients with a good prognosis resulted in a higher chance of delivering a term singleton live birth, compared with double embryo transfer. Birth rates overall were lower, but this could almost be compensated by additional cycles of frozen single embryo transfer (31). Our change in policy from elective single embryo transfer to a policy of strict single embryo transfer did not lead to lower pregnancy rates, but did lead to lower multiple pregnancy rates. The four dizygotic twin pregnancies after the policy change occurred after double embryo transfer at the request of the patients. This shows the difficulty to maintain a rigid policy of in vitro fertilisation with single embryo transfer and, in vitro fertilisation in a modified natural cycle.

The margin of equivalence, 12.5%, may be seen as relatively high, as one could argue that a 5% or 10% increase may already be important in clinical practice. We hypothesized that the pregnancy rates of in vitro fertilisation had to be considerably better than those of intrauterine insemination with controlled ovarian hyperstimulation since in vitro fertilisation is a more invasive and expensive treatment, and found 12.5% to be reasonable in this respect.

Generalisability
We included only couples with chances of conceiving below 30%, as it has earlier been established that intrauterine insemination with controlled ovarian hyperstimulation is not effective in couples with natural conception chances greater than 30% according to the model Hunault (6). The use of this prediction model obviates the need to minimise by female age, parity and duration of infertility, as all these factors are incorporated in the model.

We included only couples with women below 38 years, as this age limit had often been used in studies comparing single embryo transfer with double embryo transfer in good prognosis couples predating the design of our study (32-35). As a consequence, we did not want to expose couples where the woman is at the end of her reproductive period to single embryo transfer.
Interpretation in context of setting and intervention

Results from this trial partly confirm the findings of a meta-analysis (36). This Cochrane review, included four trials, and the results suggested that among treatment naïve couples with unexplained subfertility no significant difference existed in live birth rates for intrauterine insemination with controlled ovarian hyperstimulation versus in vitro fertilisation. However, two recent trials concluded pregnancy was achieved faster with immediate in vitro fertilisation (37,38). Apart from differences in population, the stimulation protocols were more aggressive than ours, and the multiple pregnancy rates were higher. We found no significant difference in time to pregnancy in the three arms of our study, and therefore no time is wasted by starting with intrauterine insemination with controlled ovarian hyperstimulation.

Many couples stop treatment or switch to another treatment, before an ongoing pregnancy is achieved. By allowing couples to receive other treatments if they did not achieve an ongoing pregnancy leading to a live birth after in vitro fertilisation with single embryo transfer, in vitro fertilisation in a modified natural cycle or intrauterine insemination with controlled ovarian hyperstimulation within a time horizon of one year, our study design reflects daily practice and leads to high external validity.

As we have excluded large differences in efficacy and safety between in vitro fertilisation with single embryo transfer and in vitro fertilisation in a modified natural cycle when compared to intrauterine insemination with controlled ovarian hyperstimulation, costs and patient preferences should be considered. The number of couples who discontinue treatment if a pregnancy does not occur can be considered a proxy for the burden of treatment. In this trial in vitro fertilisation in a modified natural cycle had a higher rate of couples discontinuing treatment. From that perspective in vitro fertilisation in a modified natural cycle might not be as patient-friendly as previously considered (22).

The number of multiple pregnancies in the intrauterine insemination with controlled ovarian hyperstimulation group was lower in this trial than previously reported (39-41). We aimed to achieve a maximum of three dominant follicles and used strict cancellation criteria. This resulted in low chances of multiple pregnancy while maintaining live birth rates, indicating the relative safety of intrauterine insemination with controlled ovarian hyperstimulation when strict criteria are maintained (11). Our rather cautious approach to stimulation also resulted in inducing ovulation and subsequent intra uterine insemination
with controlled ovarian hyperstimulation when only one mature follicle was present. Furthermore intrauterine insemination with controlled ovarian hyperstimulation was performed with either clomiphene citrate or follicle stimulating hormone; this may have influenced the rates of both pregnancy and multiple pregnancy. Whether the use of clomiphene citrate over follicle stimulating hormone should be preferred in intrauterine insemination with controlled ovarian hyperstimulation cycles in couples with unexplained subfertility is presently being studied in a randomised controlled trial (trial registration number: 4057).

We found comparable rates of multiple pregnancy between the three interventions. This would suggest that in vitro fertilisation with single embryo transfer or in vitro fertilisation in a modified natural cycle has no benefit in the reduction of multiple pregnancies over intrauterine insemination with controlled ovarian hyperstimulation. However, if a twin pregnancy must be avoided at all costs, intrauterine insemination with controlled ovarian hyperstimulation should not be performed, as a small risk of a multiple pregnancy still exists. In vitro fertilisation with the transfer of a single embryo, irrespective of embryo quality, should be the treatment of choice, whether after in vitro fertilisation in a modified natural cycle or in vitro fertilisation in a modified natural cycle with conventional ovarian hyperstimulation. To the detriment of in vitro fertilisation in a modified natural cycle, no surplus embryos are available for cryopreservation.

Recently, the question was posed whether couples with unexplained subfertility and an unfavourable prognosis on natural conception should be treated at all. The updated 2013 guideline from the National Institute for Health and Care Excellence abandoned intrauterine insemination with controlled ovarian hyperstimulation for these couples and recommended extended expectant management for all couples with unexplained subfertility instead, on the basis of a lack of data indicating efficacy of intrauterine insemination for unexplained subfertility (42). However, trials comparing intrauterine insemination with controlled ovarian hyperstimulation versus extended expectant management in couples with an unfavourable prognosis on natural conception have not yet been performed. We believe that such a trial is necessary before such guidelines are implemented.
Conclusions
We have shown that in vitro fertilisation with single embryo transfer and in vitro fertilisation in a modified natural cycle were non-inferior to intrauterine insemination with controlled ovarian hyperstimulation in terms of the birth of a healthy child with comparable multiple pregnancy rates. The absence of a marked difference in pregnancy outcomes suggests that the more invasive in vitro fertilisation with single embryo transfer and in vitro fertilisation in a modified natural cycle may not be desirable alternatives for intrauterine insemination with controlled ovarian hyperstimulation. In view of these results, there seems no reason to abandon intrauterine insemination with controlled ovarian hyperstimulation as a first-line treatment of couples with unexplained or mild male subfertility and an unfavourable prognosis for natural conception.
REFERENCES


