

IVF for unexplained subfertility; whom should we treat?

R. van Eekelen^{1,2,*}, N. van Geloven³, M. van Wely¹, S. Bhattacharya⁴,
F. van der Veen¹, M.J. Eijkemans², and D.J. McLernon⁵

¹Centre for Reproductive Medicine, Academic Medical Centre, Meibergdreef 9, 1105 AZ, Amsterdam, The Netherlands ²Department of Biostatistics and Research Support, Julius Centre, University Medical Centre Utrecht, Heidelberglaan 100, 3584 CX, Utrecht, The Netherlands ³Medical Statistics, Department of Biomedical Data Sciences, Leiden University Medical Centre, Einthovenweg 20, 2333 ZC, Leiden, The Netherlands ⁴Cardiff University School of Medicine, Heath Park Way, Cardiff, CF14 4XN, UK ⁵Medical Statistics Team, Institute of Applied Health Sciences, University of Aberdeen, Foresterhill, AB24 2ZD Aberdeen, UK

*Correspondence address. Centre for Reproductive Medicine, Academic Medical Centre, Meibergdreef 9, 1105 AZ, Amsterdam, The Netherlands and Department of Biostatistics and Research Support, Julius Centre, University Medical Centre Utrecht, Heidelberglaan 100, 3584 CX, Utrecht, The Netherlands; E-mail: r.vaneekelen@amc.uva.nl

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STUDY QUESTION: Which couples with unexplained subfertility can expect increased chances of ongoing pregnancy with IVF compared to expectant management?

SUMMARY ANSWER: For couples in which the woman is under 40 years of age, IVF is associated with higher chances of conception than expectant management.

WHAT IS KNOWN ALREADY: The clinical indications for IVF have expanded over time from bilateral tubal blockage to include unexplained subfertility in which there is no identifiable barrier to conception. Yet, there is little evidence from randomized controlled trials that IVF is effective in these couples.

STUDY DESIGN, SIZE, DURATION: We compared outcomes in British couples with unexplained subfertility undergoing IVF ($n = 40\,921$) from registry data to couples with the same type of subfertility on expectant management. Those couples on expectant management (defined as no intervention aside from the advice to have intercourse) comprised a prospective nation-wide Dutch cohort ($n = 4875$) and a retrospective regional cohort from Aberdeen, Scotland ($n = 975$). We excluded couples who had tried for <1 year to conceive and also those with anovulation, uni- or bilateral tubal occlusion, mild or severe endometriosis or male subfertility i.e. impaired semen quality according to World Health Organization criteria.

PARTICIPANTS/MATERIALS, SETTING, METHODS: We matched couples who received IVF and couples on expectant management based on their characteristics to control for confounding. We fitted a Cox proportional hazards model including patient characteristics, IVF treatment and their interactions to estimate the individualized chance of conception over 1 year—either following IVF or expectant management for all combinations of patient characteristics. The endpoint was conception leading to ongoing pregnancy, defined as a foetus reaching a gestational age of at least 12 weeks.

MAIN RESULTS AND THE ROLE OF CHANCE: The adjusted 1-year chance of conception was 47.9% (95% CI: 45.0–50.9) after IVF and 26.1% (95% CI: 24.2–28.0) after expectant management. The absolute difference in the average adjusted 1-year chances of conception was 21.8% (95%CI: 18.3–25.3) in favour of IVF. The effectiveness of IVF was influenced by female age, duration of subfertility and previous pregnancy. IVF was effective in women under 40 years, but the 1-year chance of an IVF conception declined sharply in women over 34 years. In contrast, in woman over 40 years of age, IVF was less effective, with an absolute difference in chance compared to expectant management of 10% or lower. Regardless of female age, IVF was also less effective in couples with a short period of secondary subfertility (1 year) who had chances of natural conception of 30% or above.

LIMITATIONS, REASONS FOR CAUTION: The 1-year chances of conception were based on three cohorts with different sampling mechanisms. Despite adjustment for the three most important prognostic patient characteristics, namely female age, duration of subfertility and primary or secondary subfertility, our estimates might not be free from residual confounding.

WIDER IMPLICATIONS OF THE FINDINGS: IVF should be used selectively based on judgements on gain compared to continuing expectant management for a given couple. Our results can be used by clinicians to counsel couples with unexplained subfertility, to inform their expectations and facilitate evidence-based, shared decision making.

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Introduction

Subfertility is defined as not conceiving within 1 year of regular unprotected intercourse and this affects approximately one in nine heterosexual couples (Datta *et al.*, 2016). Following standard investigations, no cause can be identified in one-third of these couples who are said to have unexplained subfertility. IVF with or without ICSI, is a commonly used treatment for couples with prolonged unresolved subfertility and over 470 000 treatment cycles were recorded in Europe in 2013 (Calhaz-Jorge *et al.*, 2017). IVF is a burden to couples in terms of mental and physical stress, is associated with high expectations and considerable investment in terms of emotions, finances and time (Rooney and Domar, 2016). The number of IVF cycles conducted increases annually, posing an increasing burden on health services in countries where IVF is publicly funded (HFEA, 2004; Andersen *et al.*, 2007; NVOG, 2010; NICE, 2013; Kamphuis *et al.*, 2014; Calhaz-Jorge *et al.*, 2017; HFEA, 2018). This increase is generally considered to be the consequence of the increasingly liberal utilization of IVF for a variety of indications, including unexplained subfertility (HFEA, 2004; Kamphuis *et al.*, 2014; HFEA, 2015). Yet, there is little robust evidence supporting the effectiveness of IVF in couples with unexplained subfertility compared to a wait-and-see approach i.e. expectant management (Pandian *et al.*, 2015; Tjon-Kon-Fat *et al.*, 2016).

There is a single trial evaluating the effectiveness of IVF versus expectant management for couples with unexplained subfertility in terms of live birth which reported the chance of live birth following IVF (11 out of 24 couples) to be 12 times that of expectant management (1 out of 27 couples) (Hughes *et al.*, 2004). Although the results seem to support IVF, there is considerable uncertainty around this result based on very small numbers of participants and it is inappropriate for clinical practice across the globe to be based on this quality of evidence (Tjon-Kon-Fat *et al.*, 2016).

Observational studies have separately quantified the predicted chances of conception after IVF and after a period of expectant management (Leushuis *et al.*, 2009; McLernon *et al.*, 2016; van Eekelen *et al.*, 2017a). There are two problems that hamper the comparability of these predictions, which currently limit their clinical utility. First, the prognoses were derived from separate studies with dissimilar patient characteristics. For instance, women with unexplained subfertility who received IVF are generally older than women who pursued expectant management. Second, the prognosis after IVF is expressed per embryo transfer or per complete IVF cycle while the prognosis associated with expectant management is expressed in terms of calendar time, commonly over 1 year (Daya, 2005).

We can address these problems by adjusting for differences between couples who were treated with IVF and couples who pursued expectant

management and expressing predicted chances over a uniform time horizon. To this end, we opted for a pragmatic approach by analysing data from three observational cohorts: the UK national IVF registry and two groups of couples (from the Netherlands and Scotland, respectively) who embarked on a variable period of expectant management.

Our aim was threefold. First, to use individual patient data from these three cohorts to compare the average absolute unadjusted 1-year chance of conception after IVF or expectant management. Second, to compare the adjusted 1-year chance of conception after IVF or expectant management and third, to estimate the effectiveness of IVF in individual patients based on their clinical characteristics.

Materials and Methods

The population comprised couples with unexplained subfertility seen in fertility clinics. The exposure was all IVF cycles and subsequent embryo transfers performed within 1 year after the start of ovarian stimulation. The comparator in the unexposed group was expectant management for 1 year after completion of the fertility workup. The outcome of interest was conception leading to ongoing pregnancy.

IVF cohort

Data on couples treated with IVF between 1999 and 2011 were obtained from the Human Fertilisation and Embryology Authority (HFEA) registry which collects data from all licensed clinics in the UK (McLernon *et al.*, 2016). From 2009 onwards, the number of women included was limited because explicit consent was required for the use of their data for research purposes (McLernon *et al.*, 2016).

Expectant management cohorts

We combined data from two separate cohorts comprising couples with unexplained subfertility who underwent expectant management. The first was a prospective cohort assembled across 38 hospitals in The Netherlands between January 2002 and February 2004. Couples were followed for natural conception from the completion of the fertility workup onwards. The detailed protocol for this has been described elsewhere (van der Steeg *et al.*, 2007). The second was a retrospective population-based cohort from the Grampian region of Scotland comprising subfertile couples who registered at Aberdeen Fertility Clinic. Using a unique, pseudonomized identifier, we linked patient records, including demographic and diagnostic information, from the fertility clinic to treatment records from Aberdeen Assisted Reproduction Unit Database and to pregnancy outcomes from the

Aberdeen Maternity and Neonatal Databank ([van Eekelen et al., 2018](#)). This process was carried out according to the Standard Operating Procedures of the Data Management Team, University of Aberdeen. We selected couples living in the Aberdeen City District whose births occurred at Aberdeen Fertility Clinic. Pregnancy outcomes from natural conceptions were identified by linkage with the Aberdeen Maternity and Neonatal Databank, which captures all birth outcomes in this region ([Ayorinde et al., 2016](#)).

Inclusion and exclusion criteria

Couples who had been trying for a pregnancy for <1 year, those with anovulation, uni- or bilateral tubal occlusion, mild or severe endometriosis and male subfertility i.e. impaired semen quality according to World Health Organization criteria were excluded from the UK IVF and Scottish cohorts ([WHO, 1999](#); [WHO, 2010](#)). For the Dutch cohort, the same exclusion criteria were applied, except that mild endometriosis was considered as a part of unexplained subfertility and male subfertility was defined as a total motile count below 1 million ([van Eekelen et al., 2017a](#)).

Treatment protocols

Decisions regarding treatment were based on local and national protocols. In short, the UK IVF registry comprises every IVF cycle, with guidelines changing over time ([NICE, 2013](#)). Treatment decisions for the Dutch cohort were left to the discretion of physicians in agreement with their patients ([NVOG, 2004](#); [van der Steeg et al., 2007](#)) and in the Scottish cohort by the local protocol and national guideline ([NICE, 2013](#)).

Expectant management was defined as no intervention aside from the advice to have intercourse.

Definitions for outcome and follow up

Our outcome of interest was conception leading to an ongoing pregnancy, defined as a foetus reaching a gestational age of at <12 weeks visualised by ultrasound. The date of conception was defined as the first day of the last menstruation period prior to conception. We analysed data up to a maximum of 1 year of follow up.

Follow up for couples on expectant management started at completion of the fertility workup and ended, for those who did not conceive, at 1 year after the workup, on the date of last contact or the date of starting ovarian stimulation for IUI or IVF treatment (whichever came first) i.e. we censored their time-to-pregnancy. We assumed that couples who continued with expectant management were no different, in terms of their clinical characteristics and resulting prognosis, to those who were censored (non-informative censoring).

Couples who received IVF were followed from the start of ovarian stimulation in the first cycle up until their last embryo transfer. Since the IVF registry contained all UK IVF cycles from 1999 to 2011, all ongoing IVF pregnancies within 1 year of initiating the first cycle (i.e. all fresh and frozen cycles) were recorded and we thus had complete 1-year follow up during which couples received 1.5 embryo transfers on average. This assumes that couples who discontinued treatment had zero chance of conception after IVF afterwards, for instance for reasons related to an insufficient number of oocytes collected during follicle aspiration, a low fertilization rate or financial reasons ([Daya, 2005](#)).

To align with our assumption of pursuing one full year of expectant management, we also considered the hypothetical scenario in which couples continued their IVF attempts for a full year of follow up during which they underwent three to four embryo transfers on average. In the supplementary analysis following this scenario, we censored time-to-pregnancy in couples receiving IVF after their last unsuccessful IVF transfer, defined as the first day of menstruation before the last embryo transfer. We thus also assumed non-informative censoring in IVF i.e. that couples who continued IVF were similar to couples who dropped out of IVF.

Missing data

To be able to compare couples who received IVF and couples who underwent expectant management, we had to make assumptions around the dates of ovarian stimulation and first day of menstruation in couples who had IVF. As couples start their IVF treatment with ovarian stimulation, we elected to follow couples from that date until conception (the first day of last menstruation before the final embryo transfer) to align with the general definition of time to natural conception. Since dates of initiation of ovarian stimulation were not available in the UK IVF database and are not applicable to frozen/thawed cycles, we assumed a period of 15 days before the date of embryo transfer ([Alport et al., 2011](#)).

In the Dutch cohort, the date of workup completion could be derived and this date was used as the start of follow up ([van Eekelen et al., 2017a](#)). For the Scottish cohort, this date was not available and was estimated at 6 weeks after the date of registration, which was the average time between registration and completion of the fertility workup in the Dutch cohort.

The prognostic patient characteristics that were recorded in all cohorts were female age, duration of subfertility and (female) primary or secondary subfertility. In the UK IVF cohort, data for primary or secondary subfertility from 2008 onwards ($n = 7532$, 18%) were not systematically recorded and were considered as missing. Because of these missing values, we applied multiple imputation including all relevant prognostic characteristics and a covariate for the cumulative hazard of pregnancy to account for the aspect of time in the data, creating 10 imputation sets ([White and Royston, 2009](#)). In the Dutch cohort, fewer than 1% of data used for the present study were missing and were accounted for in a previous study by multiple imputation, creating 10 imputation sets ([van Eekelen et al., 2017a](#)). In the Scottish cohort, fewer than 1% of data were missing and we applied multiple imputation identical to the approach in the UK IVF cohort. Ten imputation sets were thus created separately for the three cohorts, then combined to derive 10 combined datasets and we pooled their results using Rubin's Rules ([Rubin, 2004](#)).

Matching procedure

To ensure that there was minimal confounding due to the three prognostic patient characteristics (female age, duration of subfertility and previous pregnancy), we applied matching ([Austin, 2014](#)). In this matching procedure, we paired couples on expectant management to couples that received IVF that had the same (rounded) female age, duration of subfertility and primary or secondary subfertility status. We found all possible pairs with replacement, which allows each patient to be used as a match more than once. This yields higher quality matches

than matching without replacement due to data on all matches being used (Abadie and Imbens, 2006). Then, we weighted couples such that the expectant management group was the reference or 'target population'. Thus, in the resulting complete 'matched' dataset, the average patient characteristics and sample size of couples on expectant management were now identical to couples who received IVF. Using this matched data, we estimate what would happen if couples on expectant management would instead start IVF (referred to as the average treatment effect in controls, or ATC) (Austin, 2014).

Statistical analysis

Average effect of IVF

We calculated the unadjusted 1-year chance of conception after IVF as the observed fraction of couples who conceived within 1 year of IVF on the original, unmatched dataset. We estimated the unadjusted 1-year chance of conception after expectant management with the Kaplan–Meier method on the original, unmatched dataset. We calculated the average unadjusted effect as the absolute difference of these two chances. To estimate the adjusted chances and the adjusted average effect, we repeated both these analyses on the matched dataset.

Individualized effectiveness of IVF

We defined the individualized effectiveness of IVF as the absolute difference between the estimated 1-year chance of conception after IVF and the 1-year chance when pursuing expectant management for a couple based on female age, duration of subfertility and primary/secondary subfertility status. To estimate these individual chances, we fitted a Cox proportional hazards model on the original, unmatched dataset using treatment (IVF or expectant management), the patient characteristics and the interaction between treatment and patient characteristics as covariates. This was done following three steps.

We first determined how female age and duration of subfertility could best be entered into our statistical model; we evaluated both linear and non-linear associations with the log hazard of conception using linear terms or restricted cubic splines, then tested which fitted better using Wald tests and Akaike's Information Criterion (AIC) (Akaike, 1974; Harrell et al., 1996).

Once a suitable form for female age and duration of subfertility was determined, we included IVF treatment, female age, duration of subfertility, primary or secondary subfertility and all interaction terms with IVF treatment in the model to assess if the effect of IVF depended on these characteristics. We then tested all interaction terms simultaneously with an overall Wald test. If this test was significant, we performed backwards selection on the full model using Wald tests per separate interaction and AIC to determine which interaction was informative and removed those that were not (Akaike, 1974). We checked the proportional hazards assumption for all covariates in the model using scaled Schoenfeld residuals (Grambsch and Therneau, 1994) and accounted for the non-proportional hazard for IVF treatment versus expectant management by stratifying on treatment group.

After the final model fit, we visualized the association between patient characteristics which varied the effect of IVF by estimating the 1-year chances of conception for couples with different characteristics.

In addition, we estimated chances for all combinations of patient characteristics, tabulating the estimated chances, their corresponding

95% CIs, absolute differences, relative differences and the number needed to treat (NNT).

Supplementary analyses

In the first supplementary analysis, in order to estimate the outcome if couples would continue to have IVF over 1 full year, we used the Kaplan–Meier method both for couples receiving IVF and for couples pursuing expectant management on the original and matched datasets.

In the second supplementary analysis, we again estimated individualized chances after both IVF and expectant management but now expressed over a period of 6 months. We tabulated these 6-month chances as well as their corresponding 95% CIs, absolute differences, relative differences and the NNT.

The study was approved by the North of Scotland Research Ethics Committee (17/NS/0122). Data linkage and all statistical analyses were performed in the Data Safe Haven of the University of Aberdeen using R version 3.4.3 (R Core Team (2017). R: A language and environment for statistical computing. R Foundation for Statistical Computing, Vienna, Austria. <http://www.R-project.org/>) and RStudio using the *survival* package for the Kaplan–Meier method, *mice* for multiple imputation of missing data, *rms* for functions for splines and fitting Cox models and *Matching* to conduct the matching by patient characteristics.

Results

Data from a total of 46 771 couples were available for analysis (Fig. 1). Out of 40 921 couples in the UK IVF cohort who received 61 019 embryo transfers in total, 16 281 conceived (39.8% of couples, 26.7% per embryo transfer) within 1 year of starting IVF. In total, 32 396 (79%) couples received IVF and 8525 (21%) received ICSI. There were 4891 multiple gestations after IVF (12% of couples, 30% of conceptions). Out of 4875 couples in the Dutch cohort pursuing expectant management, 903 (18.5%) couples conceived naturally within 1 year after completion of the fertility workup. There were 11 multiple gestations (0.2% of couples, 1.2% of conceptions). Out of 975 couples in the Scottish cohort pursuing expectant management, 229 (23.5%) couples conceived naturally within 1 year after completion of the fertility workup. There were no multiple gestations.

The median duration of follow up for couples receiving IVF was one embryo transfer (25th–75th percentile: 0–7 months) as 29% of couples conceived after their first embryo transfer and 21% discontinued IVF treatment after their first unsuccessful embryo transfer. The median follow up for couples pursuing expectant management was 7 months (25th–75th percentile: 3–12 months).

Patient characteristics

The baseline characteristics of couples, stratified by cohort, are presented in Table I. In comparison with women who were managed expectantly, those who received IVF were older (mean 35.1 years in the UK IVF, 32.5 years in the Dutch and 33.2 years in the Scottish cohorts), had been trying to conceive for longer (median 4.0 years in UK IVF, 1.6 years in the Dutch and 2.1 years in the Scottish cohorts) but were just as likely to have primary subfertility (60% in the UK IVF, 66% in the Dutch and 59% in the Scottish cohorts).

The distributions of female age and duration of subfertility for couples who received IVF and couples who pursued expectant management are shown in Fig. 2A and B.

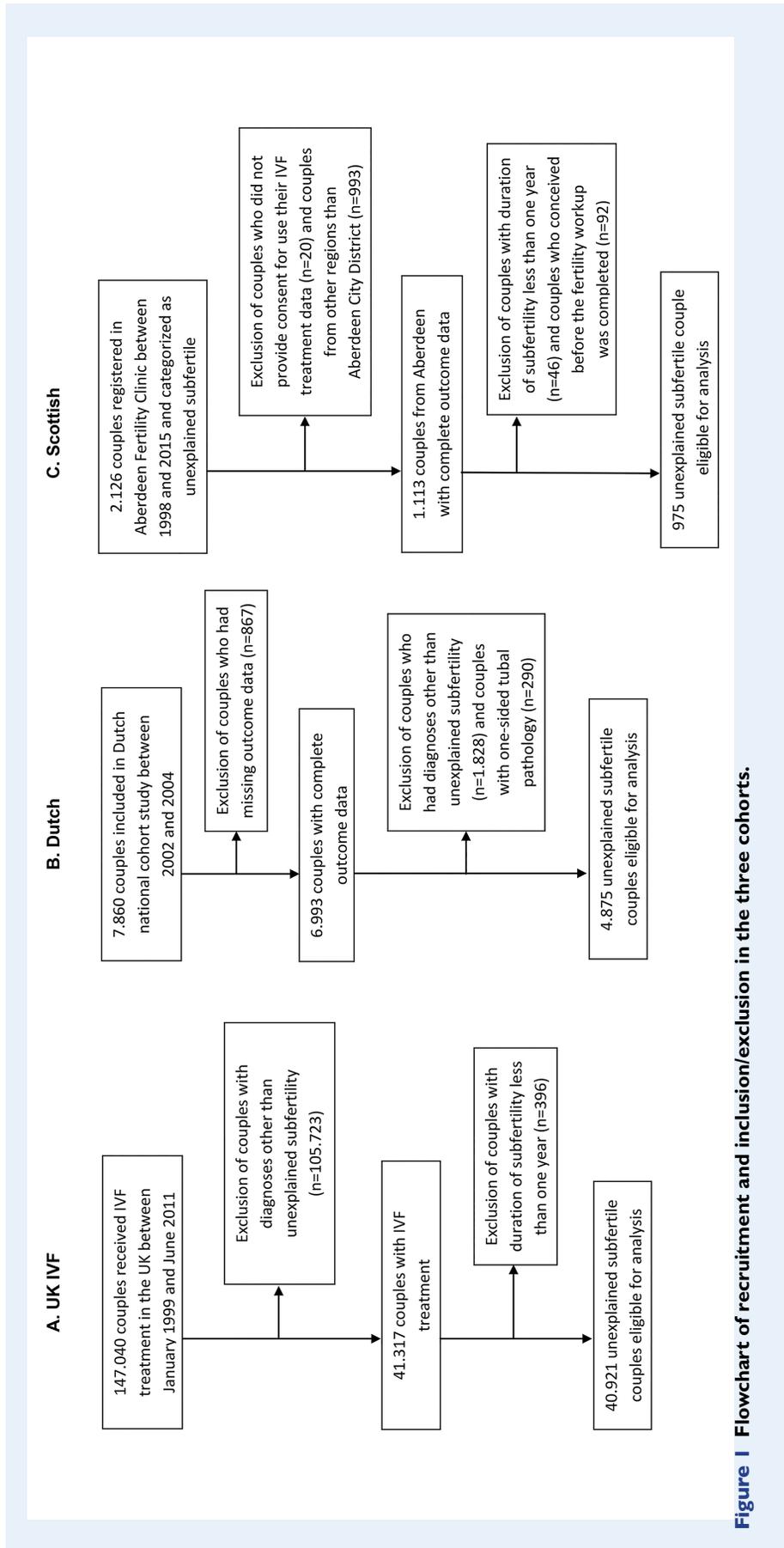


Figure 1 Flowchart of recruitment and inclusion/exclusion in the three cohorts.

Table 1 Baseline characteristics at the start of follow up for the three cohorts included in the analysis.

	UK IVF (n = 40 921)	Dutch (n = 4875)	Scottish (n = 975)
Female age in years (mean, 5th–95th percentile)	35.1 (28–42)	32.5 (24.9–39.4)	33.2 (26.1–41.1)
Duration of subfertility in years (median, 5th–95th percentile)	4.0 (1–13)	1.6 (1–4.9)	2.1 (1.1–5.1)
Primary subfertility (n, %)	24 572 (60%)	3231 (66%)	571 (59%)

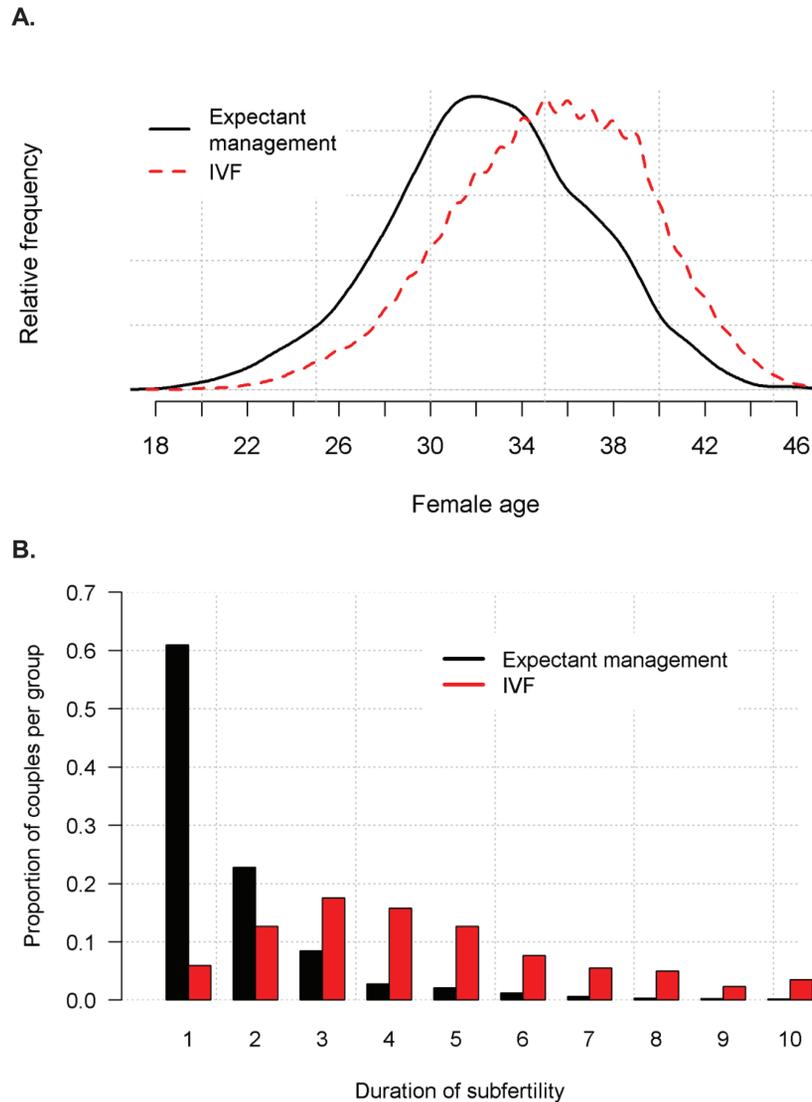


Figure 2 Overlap of patient characteristics for couples who received IVF and couples who underwent expectant management. (A) Distribution of female age per treatment group, depicted by relative frequency (density). (B) Distribution of duration of subfertility per treatment group, depicted as the proportion of couples per group who had a certain (rounded) duration.

Unadjusted average chance of conception

The unadjusted 1-year chance of conception after starting IVF was 39.8% (95% CI: 39.3–40.3) and after expectant management was 26.1% (95% CI: 24.7–27.5). The average absolute difference in the

unadjusted 1-year chance of conception was 13.6% (95% CI: 11.6–15.7) in favour of IVF. The 1-year chances following expectant management in the Dutch and Scottish cohorts were similar (26.9% and 23.8%, respectively).

Table II Estimated effects of patient characteristics on conception leading to ongoing pregnancy.

	HR for conception after IVF (95% CI)	HR for conception after expectant management (95% CI)
Female age, years (34 versus 27)*	0.99 (0.94–1.04)	0.70 (0.60–0.82)
Female age, years (40 versus 35)*	0.43 (0.41–0.46)	0.64 (0.49–0.84)
Duration of subfertility, years (6 versus 2)*	0.86 (0.80–0.92)	0.39 (0.30–0.50)
Primary versus secondary subfertility	0.98 (0.94–1.02)	0.71 (0.63–0.81)

*Contrasts between values for female age and duration of subfertility were chosen to depict their non-linear estimated effects.

Results are from the model including interaction (via stratification) with treatment.

Adjusted average chance of conception

A total of 5818 out of 5850 (99%) couples pursuing expectant management were matched with 31 867 out of 40 921 (78%) counterparts who received IVF and had the same characteristics. The adjusted 1-year chance of conception was 47.9% (95% CI: 45.0–50.9) after starting IVF and 26.1% (95% CI: 24.2–28.0) after expectant management. The average absolute difference in the adjusted 1-year chance of conception was 21.8% (95% CI: 18.3–25.3) in favour of IVF.

Individualized effectiveness of IVF

Both female age and duration of subfertility were non-linearly associated with conception (Wald tests for non-linearity both $P < 0.001$, splines with five and three knots, respectively).

There were statistically significant interactions between all three patient characteristics and IVF treatment (overall $P < 0.001$, individual interactions all $P < 0.001$).

The estimated effects of couple characteristics on conception in terms of hazard ratios (HRs) are presented in Table II. In general,

as female age increased, the chance of conception decreased both after expectant management and after IVF, but the detrimental effect of female age above 34 years on the chance of conception was stronger in the latter (HR of 40 versus 35 years: 0.43 after IVF and 0.64 after expectant management). As duration of subfertility increased, the chance of conception decreased in both groups, but this effect was stronger for those on expectant management (HR of 6 versus 2 years: 0.86 after IVF and 0.39 after expectant management). Couples with primary subfertility on expectant management had a lower chance of conception compared to couples with secondary subfertility (HR of primary versus secondary: 0.71) but there was no noticeable difference in the IVF group (HR: 0.98).

The predicted 1-year chance of conception in couples with primary subfertility of 2 years duration and female age ranging between 26 and 42 years are shown in Fig. 3. The effectiveness of IVF decreased in women over 34 years.

The predicted 1-year chances of conception in couples with primary subfertility where female age is 35 years and the duration of subfertility ranges from 1 to 8 years are visualized in Fig. 4. The effectiveness of IVF increased as the duration of subfertility increased.

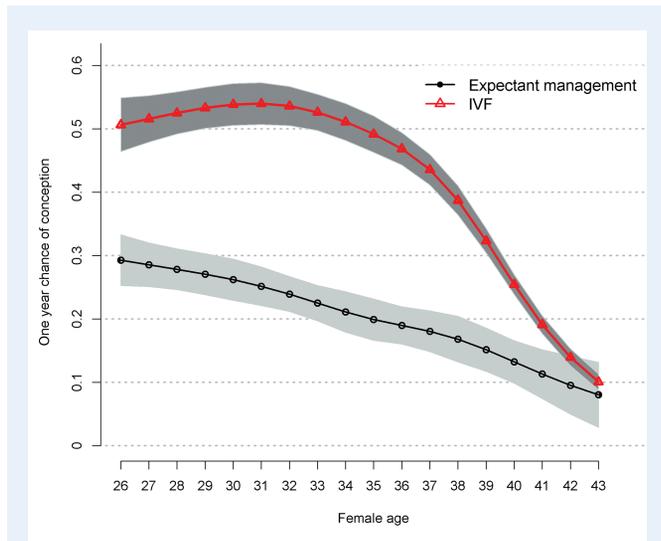


Figure 3 Association between female age and the 1-year chance of conception after receiving IVF or pursuing expectant management for a primary subfertile couple who have been trying to conceive for 2 years. Grey bands are 95% confidence limits.

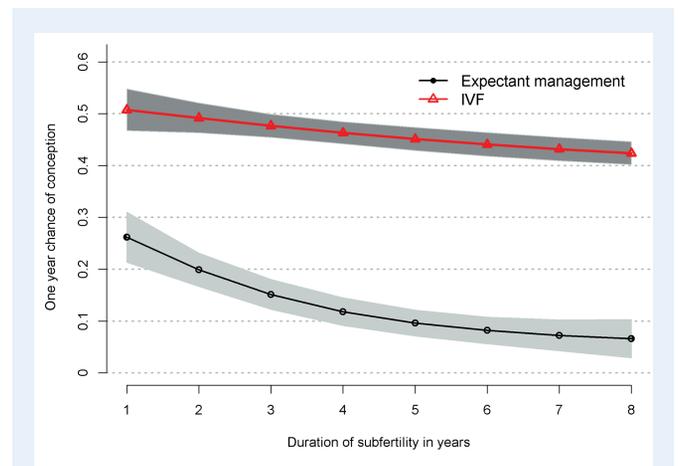


Figure 4 Association between duration of subfertility and the 1-year chance of conception receiving IVF or pursuing expectant management for a primary subfertile couple of which the woman is 35 years old. Grey bands are 95% confidence limits.

Table III Association between primary or secondary subfertility and the 1-year chance of conception after receiving IVF or pursuing expectant management for a couple of which the woman is 35 years old who have been trying to conceive for 2 years.

	1-year chance of conception after IVF (95% CI)	1-year chance of conception after expectant management (95% CI)
Primary subfertile couple	49.2 (46.3–52.1)	19.9 (16.7–23.1)
Secondary subfertile couple	50.0 (47.0–53.0)	26.7 (22.2–31.2)

The predicted 1-year chances of conception for couples with 2 year duration where female age is 35 years stratified for primary and secondary subfertility are presented in Table III. IVF was more effective for couples with primary subfertility than for couples with secondary subfertility.

In Supplementary Tables, we present full tables containing the predicted 1-year chance of conception after IVF and after starting expectant management for all combinations of patient characteristics. Also provided are the absolute differences between these chances, the relative differences and the NNT to achieve one additional conception. Predictions are presented separately for primary (Supplementary Table SI) and secondary (Supplementary Table SII) subfertility for ranges of female age from 26 to 42 years and duration of subfertility from 1 to 8 years. For instance, a typical couple undergoing IVF, where the woman is 35 years old with 4 years duration of primary subfertility, has an estimated 1-year chance of conception of 46% (95% CI: 44–48) after IVF compared to 12% (95% CI: 9–14) after expectant management, with an absolute difference of 34% and a NNT of 2.9.

On the other hand, a typical couple pursuing expectant management, where the woman is 33 years old with 2 years of primary subfertility, has an estimated 1-year chance of conception of 53% (95% CI: 50–55) after IVF compared to 23% (95% CI: 20–25) after expectant management, with an absolute difference of 30% and a NNT of 3.3.

In couples where the woman is under 40 years, IVF was effective compared to expectant management. In contrast, in couples where the woman is over 40 years, IVF was less effective as the absolute difference between chances was ~10% or lower. In couples with 1-year duration of secondary subfertility, regardless of the age of the woman, IVF was also less effective since their chances of natural conception remained relatively high at 30% or above.

Supplementary analyses

In the supplementary analysis where we estimated outcomes in couples who continued with IVF for a full year, the unadjusted 1-year chance of conception after IVF was estimated at 51.6% (95% CI: 50.9–52.2). The average absolute difference in the unadjusted 1-year chance of conception became 25.4% (95% CI: 23.1–27.7) in favour of IVF.

The adjusted 1-year chance of conception after receiving IVF for one full year was estimated at 59.7% (95% CI: 55.3–64.0). The average absolute difference in the adjusted 1-year chance of conception became 33.6% (95% CI: 28.8–38.3) in favour of IVF.

In Supplementary Tables SIII and SIV, we present the same individualized predictions as in Supplementary Tables SI and SII but now expressed over 6 months instead of 1 year.

Discussion

In couples with unexplained subfertility, we found that IVF increased the average 1-year chance of conception compared to expectant management. Factors affecting the effectiveness of IVF were female age, duration of subfertility and primary/secondary subfertility.

Although couples who received IVF had, on average, a higher female age and a higher duration of subfertility compared to couples who continued expectant management, the large sample size of treated and untreated couples resulted in sufficient overlap of case-mix to enable us to accurately estimate all the separate interactions between patient characteristics and treatment. A second strength was our ability to control for confounding in the average adjusted chance by matching on female age, duration of subfertility and primary versus secondary subfertility.

We were able to predict individualized chances of conception following either IVF or expectant management on the same time axis representing 1 year of 'real' calendar time. This is intuitive, allows for a straightforward comparison, allows for most couples to complete at least one full IVF cycle and is easier to communicate to patients compared to chances per embryo transfer or per IVF cycle. A longer follow up might increase the rates after both IVF and expectant management but may be more difficult for decision making, as the longer the follow up period becomes, the less likely couples are to continue IVF.

Aside from calculating the observed fraction of couples who conceived within 1 year in the matched data (~48%), we also estimated the adjusted chance of conception when receiving IVF for one full year i.e. when continuing IVF (~60%). The latter might be an optimistic estimate, as not all couples can continue with additional IVF cycles, for instance because of an insufficient number of oocytes or financial reasons.

Limitations of this study include the availability of only three important patient characteristics in all data sources, the missing date of completion of the fertility workup in the Scottish data and the possibility of residual confounding due to the observational nature of the data. We had to make an assumption on the time between registration and completion of the fertility workup in the Scottish cohort. In the Dutch cohort, this was on average 6 weeks (van Eekelen et al., 2018). In a previously conducted validation study, we found similar chances of ongoing pregnancy in the Scottish and Dutch cohort when assuming 6 weeks between registration and completion of the fertility workup; hence, this assumption was deemed reasonable (van Eekelen et al., 2018). The dropout rate after the first embryo transfer of 21% is >12% reported in a recent Dutch validation study, but the difference can be explained by the geographical variation

in reimbursement for the UK IVF cohort compared to full reimbursement up to three cycles at the time of the Dutch study (Leijdekkers *et al.*, 2018).

In addition, the three different data sources used different sampling mechanisms, which could potentially compromise the comparability of study populations. Couples pursuing expectant management were recruited at completion of the fertility workup (Dutch cohort) or identified retrospectively (Scottish cohort). In contrast, couples who received IVF were registered in the UK IVF database with no prior data other than diagnosis. Therefore, we were unable to assess or adjust for any selection bias that might occur between completion of the fertility workup and the start of treatment, as only couples that did not conceive naturally during that period will have ended up in the UK IVF registry, a selection which might not be fully captured by the duration of subfertility (van Eekelen *et al.*, 2017b).

As the UK IVF data were only available up to 2011 and treatment success rates were found to increase over time, our estimates for the 1-year chance after IVF might be conservative for today's practice. However, IVF rates in the UK in 2016 were found to plateau in 2013 to 25%–26% per cycle (HFEA, 2016, 2018). A recent external validation of the outcome prediction in subfertility model developed on UK IVF data up to 2008 showed good performance in Dutch data collected up to 2014, meaning that our data might reasonably reflect today's practice and pregnancy outcomes (McLernon *et al.*, 2016; Leijdekkers *et al.*, 2018). The decade has witnessed changes in embryo transfer protocols in the UK from predominantly double embryo transfer (DET) to increasing numbers of elective single embryo transfer (eSET) resulting in a decline in multiple pregnancy rates from 27% in 2008 to 16% in 2014 (Harbottle *et al.*, 2015; HFEA, 2015). Nevertheless, the impact of this change in IVF policy on our estimated chances of conception might be minor as the cumulative chances of IVF success are comparable following DET and eSET combined with subsequent transfers of frozen/thawed embryos (Lukassen *et al.*, 2005; McLernon *et al.*, 2010; Harbottle *et al.*, 2015).

The primary outcome was ongoing pregnancy because the increased logistical efforts and associated costs involved in following couples to delivery were not possible in the Dutch cohort. Ongoing pregnancy is generally considered an appropriate proxy for live birth in clinical research; ~95% of ongoing pregnancies lead to live birth (Clarke *et al.*, 2010; Braakhekke *et al.*, 2014).

A large randomized controlled trial (RCT) would be the ideal study design to assess the effectiveness of IVF compared to expectant management. Conducting such a trial now would be challenging as IVF has become an established treatment for unexplained subfertility and many couples are unconfident about the value of expectant management, overestimate IVF success and push for early active treatment (van den Boogaard *et al.*, 2011; Kersten *et al.*, 2015). In addition, many clinicians fail to take into account couples' chances of natural conception in their consultations and believe that it would be unethical to withhold early access to IVF (Kersten *et al.*, 2015). This has created a genuine lack of equipoise without which no trial can be conducted. We therefore felt that the best and most pragmatic option was to compare observational data from cohorts on expectant management and IVF (van Eekelen *et al.*, 2017b).

A key benefit of the present study is the provision of the adjusted average effectiveness of IVF compared to expectant management and,

in addition, individualized estimates, which are easy to interpret and allow for direct comparisons.

Our results may be used by clinicians to counsel couples with unexplained subfertility to inform their expectations and to avoid unnecessary treatment for some while allowing timely access to IVF for others. They can also be used to allow funders and commissioners to make decisions on access to publicly funded IVF.

Our results need to be validated in other datasets or, ideally, in RCTs involving couples with characteristics in whom the effectiveness of IVF is unclear and some equipoise remains. In addition, data on long-term follow up after the first live birth is necessary to counsel couples who wish to have multiple children.

Conclusion

The effectiveness of IVF over expectant management in unexplained subfertility depends on the characteristics of the couple and thus, IVF should be used selectively based on judgements on gain for a given couple. Our results can be used by clinicians to counsel couples with unexplained subfertility, to inform their expectations and facilitate evidence-based, shared decision making.

Supplementary data

Supplementary data are available at *Human Reproduction* online.

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Authors' roles

NvG, DJM, SB, FvdV, MvW and MJE conceived the study. MDJ oversaw the storage of all data in the Safe Haven. RvE performed the data linkage. RvE, NvG, MJE and DJM designed the statistical analysis plan. RvE and DJM analysed the data. RvE, NvG, FvdV and SB drafted the manuscript. All authors contributed critical revision to the paper and approved the final manuscript.

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Conflict of interest

SB reports acting as editor-in-chief of *HROpen*. Other authors have no conflicts.

References

- Abadie A, Imbens GW. Large sample properties of matching estimators for average treatment effects. *Econometrica* 2006;**74**:235–267.
- Akaike H. A new look at the statistical model identification. *IEEE Trans Automat Contr* 1974;**19**:716–723.
- Alport B, Case A, Lim H, Baerwald A. Does the ovarian stimulation phase length predict in vitro fertilization outcomes? *Int J Fertil Steril* 2011;**5**:134–141.
- Andersen AN, Goossens V, Gianaroli L, Felberbaum R, de Mouzon J, Nygren KG. Assisted reproductive technology in Europe, 2003. Results generated from European registers by ESHRE. *Hum Reprod* 2007;**22**:1513–1525.
- Austin PC. The use of propensity score methods with survival or time-to-event outcomes: reporting measures of effect similar to those used in randomized experiments. *Stat Med* 2014;**33**:1242–1258.
- Ayorinde AA, Wilde K, Lemon J, Campbell D, Bhattacharya S. Data resource profile: the Aberdeen maternity and neonatal databank (AMND). *Int J Epidemiol* 2016;**45**:389–394.
- Braakhekke M, Kamphuis EI, Dancet EA, Mol F, van der Veen F, Mol BW. Ongoing pregnancy qualifies best as the primary outcome measure of choice in trials in reproductive medicine: an opinion paper. *Fertil Steril* 2014;**101**:1203–1204.
- Calhaz-Jorge C, De Geyter C, Kupka MS, de Mouzon J, Erb K, Mocanu E, Motrenko T, Scaravelli G, Wyns C, Goossens V. Assisted reproductive technology in Europe, 2013: results generated from European registers by ESHRE. *Hum Reprod* 2017;**32**:1957–1973.
- Clarke JF, van Rumste MM, Farquhar CM, Johnson NP, Mol BW, Herbison P. Measuring outcomes in fertility trials: can we rely on clinical pregnancy rates? *Fertil Steril* 2010;**94**:1647–1651.
- Datta J, Palmer MJ, Tanton C, Gibson LJ, Jones KG, Macdowall W, Glasier A, Sonnenberg P, Field N, Mercer CH et al. Prevalence of infertility and help seeking among 15 000 women and men. *Hum Reprod* 2016;**31**:2108–2118.
- Daya S. Life table (survival) analysis to generate cumulative pregnancy rates in assisted reproduction: are we overestimating our success rates? *Hum Reprod* 2005;**20**:1135–1143.
- Grambsch PM, Therneau TM. Proportional hazards tests and diagnostics based on weighted residuals. *Biometrika* 1994;**81**:515–526.
- Harbottle S, Hughes C, Cutting R, Roberts S, Brison D. Elective single embryo transfer: an update to UK best practice guidelines. *Hum Fertil (Camb)* 2015;**18**:165–183.
- Harrell FE Jr, Lee KL, Mark DB. Multivariable prognostic models: issues in developing models, evaluating assumptions and adequacy, and measuring and reducing errors. *Stat Med* 1996;**15**:361–387.
- HFEA, Human Fertilisation and Embryology Authority. *Annual report and accounts 2004/05*. UK: Williams Lea Group, 2004.
- HFEA, Human Fertilisation and Embryology Authority. *Annual report and accounts 2015/16*. Williams Lea Group, UK: 2015.
- HFEA, Human Fertilisation and Embryology Authority. *Annual Report and Accounts 2016/17*. Williams Lea Group, UK: 2016.
- HFEA, Human Fertilisation and Embryology Authority. *Fertility Treatment 2014–2016: Trends and Figures*, 60, 2018.
- Hughes EG, Beecroft ML, Wilkie V, Burville L, Claman P, Tummon I, Greenblatt E, Fluker M, Thorpe K. A multicentre randomized controlled trial of expectant management versus IVF in women with fallopian tube patency. *Hum Reprod* 2004;**19**:1105–1109.
- Kamphuis EI, Bhattacharya S, van der Veen F, Mol BW, Templeton A. Are we overusing IVF? *BMJ* 2014;**348**:g252.
- Kersten FA, Hermens RP, Braat DD, Hoek A, Mol BW, Goddijn M, Nelen WL. Overtreatment in couples with unexplained infertility. *Hum Reprod* 2015;**30**:71–80.
- Leijdekkers JA, Eijkemans MJC, van Tilborg TC, Oudshoorn SC, McLernon DJ, Bhattacharya S, Mol BWJ, Broekmans FJM, Torrance HL. Predicting the cumulative chance of live birth over multiple complete cycles of in vitro fertilization: an external validation study. *Hum Reprod* 2018;**33**:1684–1695.
- Leushuis E, van der Steeg JW, Steures P, Bossuyt PM, Eijkemans MJ, van der Veen F, Mol BW, Hompes PG. Prediction models in reproductive medicine: a critical appraisal. *Hum Reprod Update* 2009;**15**:537–552.
- Lukassen HG, Braat DD, Wetzels AM, Zielhuis GA, Adang EM, Scheenjes E, Kremer JA. Two cycles with single embryo transfer versus one cycle with double embryo transfer: a randomized controlled trial. *Hum Reprod* 2005;**20**:702–708.
- McLernon DJ, Harrild K, Bergh C, Davies MJ, de Neubourg D, Dumoulin JC, Gerris J, Kremer JA, Martikainen H, Mol BW et al. Clinical effectiveness of elective single versus double embryo transfer: meta-analysis of individual patient data from randomised trials. *BMJ* 2010;**341**:c6945.
- McLernon DJ, Steyerberg EW, Te Velde ER, Lee AJ, Bhattacharya S. Predicting the chances of a live birth after one or more complete cycles of in vitro fertilisation: population based study of linked cycle data from 113 873 women. *BMJ* 2016;**355**:i5735.
- NICE, National Institute for Health and Care Excellence. Guideline on: Fertility problems: assessment and treatment (2013). <https://www.nice.org.uk/guidance/cg156> (5 February 2017, date last accessed).
- NVOG, Dutch Society for Obstetrics and Gynaecology. In: *Guideline on basic fertility workup*, 2004.
- NVOG, Dutch Society for Obstetrics and Gynaecology. Guideline on: subfertility (2010). <http://bit.ly/1UhuYMV> (5 February 2017, date last accessed).
- Pandian Z, Gibreel A and Bhattacharya S. In vitro fertilisation for unexplained subfertility. *Cochrane Database Syst Rev* 2015;**2**:Cd003357.
- R Core Team (2017). R: a language and environment for statistical computing. *R Foundation for Statistical Computing, Vienna, Austria*. <http://www.R-project.org/>.
- Rooney KL, Domar AD. The impact of stress on fertility treatment. *Curr Opin Obstet Gynecol* 2016;**28**:198–201.
- Rubin D. *Multiple Imputation for Nonresponse in Surveys*. New York: John Wiley and Sons, 2004.
- Tjon-Kon-Fat RI, Bendsdorp AJ, Scholten I, Repping S, van Wely M, Mol BW, van der Veen F. IUI and IVF for unexplained subfertility: where did we go wrong? *Hum Reprod* 2016;**31**:2665–2667.
- van den Boogaard NM, van den Boogaard E, Bokslag A, van Zwieten MC, Hompes PG, Bhattacharya S, Nelen W, van der Veen F, Mol BW. Patients' and professionals' barriers and facilitators of tailored expectant management in subfertile couples with a good prognosis of a natural conception. *Hum Reprod* 2011;**26**:2122–2128.
- van Eekelen R, Scholten I, Tjon-Kon-Fat RI, van der Steeg JW, Steures P, Hompes P, van Wely M, van der Veen F, Mol BW, Eijkemans MJ et al. Natural conception: repeated predictions over time. *Hum Reprod* 2017a;**32**:346–353.

- van Eekelen R, van Geloven N, van Wely M, McLernon DJ, Eijkemans MJ, Repping S, Steyerberg EW, Mol BW, Bhattacharya S, van der Veen F. Constructing the crystal ball: how to get reliable prognostic information for the management of subfertile couples. *Hum Reprod* 2017b;**32**:2153–2158.
- van Eekelen R, McLernon DJ, van Wely M, Eijkemans MJ, Bhattacharya S, van der Veen F, van Geloven N. External validation of a dynamic prediction model for repeated predictions of natural conception over time. *Hum Reprod* 2018;**33**:2268–2275.
- van der Steeg JW, Steures P, Eijkemans MJ, Habbema JD, Hompes PG, Broekmans FJ, van Dessel HJ, Bossuyt PM, van der Veen F, Mol BW. : Pregnancy is predictable: a large-scale prospective external validation of the prediction of spontaneous pregnancy in subfertile couples. *Hum Reprod* 2007;**22**:536–542.
- White IR, Royston P. Imputing missing covariate values for the cox model. *Stat Med* 2009;**28**:1982–1998.
- WHO, World Health Organisation. *Laboratory Manual for the Examination of Human Semen and Sperm-Cervical Mucus Interaction*, 4th edn. Cambridge: Cambridge University Press, 1999.
- WHO World Health Organisation. *Laboratory Manual for the Examination and Processing of Human Semen*, 5th edn. Geneva: World Health Organization, 2010.