Title
Adjustment to unmet parenthood goals: Systematic review of long-term adjustment after failed fertility treatment

Running title
Adjustment to unmet parenthood goals

Sofia Gameiro
Cardiff Fertility Studies Research Group, School of Psychology, Cardiff University

Amy Finnigan
School of Medicine, Cardiff University

Address for correspondence:
Sofia Gameiro, Cardiff Fertility Studies Research Group, School of Psychology, Cardiff University, Tower Building, Park Place, Cardiff, Wales CF10 3AT, UK. Email: GameiroS@cardiff.ac.uk
Table of contents

Introduction

Methods

- Systematic search
- Inclusion and exclusion criteria
- Data extraction
- Data synthesis and analysis
- Risk of bias assessment

Results

- Description of studies
- Quantitative meta-synthesis: Psychosocial adjustment after failed fertility treatment
- Qualitative meta-synthesis: Patients' perceived experiences of the period after failed treatment

Discussion

- Strengths and limitations

Conclusions
Abstract

**Background:** Fertility treatment is not guaranteed to succeed and around 30% of patients do not achieve parenthood. Failed treatment represents the loss of parenthood and often triggers intense and prolonged grief reactions. There is an increasing awareness of the need to support patients in the aftermath of failed treatment, however there are no effective interventions for the effect. This need for support is expected to increase as the number of people delaying parenthood increases and therefore so does the number for whom assisted reproduction will not offer a solution.

**Objective and rationale:** This mixed-methods review aims at investigating if patients who undergo failed fertility treatment experience significant psychosocial adjustment difficulties that warrant the provision of psychosocial support, and at developing a comprehensive model of adjustment to unmet parenthood goals that can be used to assist the design of theory led psychosocial interventions.

**Search methods:** Five databases were systematically searched between 1978 and December 2015. Search terms included fertility treatment AND psychosocial adjustment AND post-treatment. Quantitative studies had to include group mean comparisons on psychosocial adjustment (mental-health, wellbeing) between patients who did failed treatment and a control group. Qualitative studies had to focus on experiences of psychosocial adjustment after failed treatment. Screening, data extraction and critical appraisal were done independently by the authors using pre-defined protocols. Two meta-analyses were performed on mental-health and wellbeing with a random effect model. Primary outcome was Hedge’s g. Publication bias was checked with visual inspection of funnel plots, Egger’s test and the trim-and-fill method. A 3-stage thematic analysis of results reported in primary qualitative papers was implemented. First-order descriptive and second-order interpretative themes were extracted.

**Outcomes:** Nine quantitative (9052 individuals, 8 countries) and 9 qualitative (267 individuals, 6 countries) studies were included. Six (67%) of the quantitative studies reported on mental-health and 7 (78%) on wellbeing. The meta-analyses showed that the failed group had worse mental-health \( (g = -0.450, P=0.002, 95\% CI [-0.734 -0.267]; I^2=85\%, P<0.001) \) and wellbeing \( (g = -0.319, P<0.001, 95\% CI [-0.439 -0.198], I^2=45\%, P=0.001) \) than controls. The qualitative review resulted in 28 first-order themes that were grouped into 6 second-order themes: individual and relational adjustment, social adjustment, acceptance, pursuit of new life goals, meaning making, and fertility care perceptions and needs. The data showed that individual, relational and social adjustment tended to increase with time since treatment, and that individuals’ care perceptions and needs also changed. The data also suggested that individuals who engage in the psychological tasks of accepting and making meaning of their situation and pursuing new life goals adjust better and have fewer support needs. These
predictions were articulated in the Three Tasks Model of Adjustment to Unmet Parenthood Goals.

**Wider implications:** Results provide compelling evidence for the provision of psychosocial care directed at helping individuals relinquishing their parenthood goals. The model developed offers comprehensive guidance on the therapeutic mechanisms that psychosocial care should target to promote adjustment. Future research should test the model with prospective cohorts studies or by developing and testing interventions based on its predictions.

**Key-words:** parenthood; infertility; childlessness; psychosocial adjustment; mental-health; wellbeing; psychological care; mixed-methods; systematic review; meta-analysis.
Introduction

Around 10% of individuals experience infertility and half of these undergo fertility treatments such as intra-uterine insemination (IUI) or In-Vitro Fertilization (IVF) to conceive (Boivin et al. 2007), (Datta et al. 2016). For most people fertility treatment represents their only chance to achieve their parenthood goals, that is, to have the number of children they desire. However, treatment is not guaranteed to succeed and around 30% of patients do not achieve parenthood or end up having less children than they would like to (Pinborg et al. 2009, Troude et al. 2016). For these people the remaining options are to forgo their parenthood goals or apply for adoption. Therefore, for thousands of people failed fertility treatment represents the loss of biological parenthood, or parenthood in general, and usually triggers intense and prolonged grief reactions (Daniluk 2001, Volgsten et al. 2010). There is an increasing awareness of the need to support individuals in the aftermath of failed fertility treatment and of the preventive role fertility clinics can play in this context (Gameiro et al. 2013). However, the recently published guidelines of the European Society for Human Reproduction and Embryology (ESHRE) could not list a single evidence-based intervention with this aim, suggesting that patients' support needs are not being addressed (Gameiro et al. 2015). This support need is expected to increase as the number of people delaying parenthood increases and therefore so does the number for whom assisted reproduction will not offer a solution (Leridon 2004). The present mixed-methods review aims to investigate if patients who undergo failed fertility treatment experience significant psychosocial adjustment difficulties that warrant the provision of psychosocial care and at developing a comprehensive model of adjustment to unmet parenthood goals that can be used to guide the design of psychosocial interventions.

Despite the call for psychosocial care in the aftermath of treatment (Gameiro et al. 2015), it is still unclear if individuals who do failed treatment adjust significantly worse than those for whom treatment is successful. Identity theories (Thoits 1992) would suggest this to be the case because failed treatment represents the loss of parenthood. For some it may only represent the loss of biological parenthood, for others the complete loss of parenthood, and for others yet the impossibility to have more children. Despite the significant differences in these experiences, in this article and we will only refer generically to the loss of parenthood or parenthood goals.

Research has shown that undesired childlessness is associated with poorer psychosocial adjustment (e.g., Graham 2015) and those using fertility treatment can be considered especially at risk for maladjustment because of their high commitment to become parents. Indeed, 84% of patients undergo all treatment cycles recommended by their physician (Gameiro et al. 2013) despite treatment being emotionally and physically demanding and disruptive of personal and professional routines. During the last decade several studies
investigated adjustment after failed treatment but so far the evidence is mixed. Some studies show impairments in mental-health and wellbeing for patients who experienced failed treatment (e.g., Kuivasaari-Pirinen et al. 2014, Yli-Kuha et al. 2010) but others do not (Hammarberg et al. 2001, Wischmann et al. 2012). This inconsistency may be partially explained by the conceptual and methodological heterogeneity observed.

First, different comparison groups were used. Researchers either compared patients who ended successful versus failed treatment (treatment outcome), or patients with children versus without children at the end of treatment (parenthood status). While the first approach is based on the assumption that it is the non-realization of the individuals’ parenthood goals that hinders adjustment (less children than desired), the second is based on the assumption that it is their parenthood status (no children at all). These two factors (unmet parenthood goals, parenthood status) are difficult to disentangle and there is evidence to suggest that they interact, whereby the inability to have a(nother) desired child results in higher suffering for childless than non-childless individuals (Gameiro et al. 2016). Second, sometimes the failed treatment group included individuals who already had children before treatment or who adopted or conceived spontaneously after treatment and this may have attenuated group differences in adjustment. Third, the follow-up assessment period varied from immediately after treatment to up to 20 years after, which may have created mixed results because loss reactions are expected to attenuate with time (Bonanno 2004). A meta-analysis taking these potential moderators into account can clarify on the severity of adjustment difficulties experienced after failed fertility treatment.

Another important issue is to understand how to support people in the aftermath of failed fertility treatment. So far researchers focused on understanding short-term reactions to failed treatment cycles (e.g., Berghuis and Stanton 2002, Terry and Hynes 1998). The study of long-term (i.e., more than 1 year after) adjustment to unmet parenthood goals is still an emergent research topic within the field that has mostly been investigated with qualitative research methods aiming to describe patients’ experiences and needs during the post-treatment period. To date only a few studies focused on identifying risk factors or exploring the mechanisms of adjustment after failed treatment (e.g., Verhaak et al. 2007), despite the existence of multiple psychological theories that can be used for the effect. In the following paragraphs we briefly describe the theoretical paradigms that have been used so far and related evidence.

Stress and coping theory states that adjustment depends on the fit between the stressor experienced and the strategies used to address it (Lazarus and Folkman 1984). Stanton and colleagues (1994) hypothesize that when stress results from a significant loss the use of emotional approach coping strategies, such as efforts to acknowledge, understand and express emotions, should promote adjustment, while emotional avoidance strategies have
been found to hinder it (Berghuis and Stanton 2002). The only two studies that investigated coping after failed treatment showed that (active and passive) avoidant emotional coping was associated with higher distress up until 5 years after treatment (Daniluk and Tench 2007, Peterson et al. 2009). In addition, meaning based coping was associated with lower personal and relational distress for women, but higher social distress for men (Peterson et al. 2009).

Other risk factors for long-term maladjustment were lack of alternative non-parenting roles and social support. The latter may hinder the use of emotional approach coping. Cross sectional studies that investigated adjustment to definitive childlessness had similar findings (Kraaij et al. 2009, Lechner et al. 2007). In addition, these studies showed that cognitive coping is also important for adjustment. While catastrophizing (i.e., emphasizing the negative aspects of childlessness) was associated with worse adjustment, positive reappraisal coping (i.e., attaching a positive meaning to childlessness in terms of personal growth) was associated with better adjustment.

Self-regulation theories hypothesize that when all treatment options are exhausted and parenthood goals are highly blocked, individuals who disengage from parenthood and engage in other meaningful goals adjust better than individuals who remain engaged (Heckhausen et al. 2001). Findings from studies focusing on undesired childlessness support this view from both a behavioural and cognitive perspective (e.g., Heckhausen et al. 2001, Kraaij et al. 2009). However, research shows that people take time to disengage from parenthood: 44% of individuals still desire to have (more) children 3 to 5 years after ending treatment (Verhaak et al. 2007), 25% 10 years after (Wischmann et al. 2012), and 6% 11 to 17 years after (Gameiro et al. 2016). As hypothesized by identity theories (Thoits 1992), childlessness seems to be especially distressing when parenthood is an essential goal in life or when people have few alternative identities or goals (McQuillan et al. 2003, Moura-Ramos et al. 2012).

According to the dual process model of grief (Stroebe and Schut 1999), adjustment to loss depends on peoples' ability to process the loss and, in addition, its implications, for instance the impact it may have on the partnership. Around 42% of individuals experience symptoms of complicated grief after ending fertility treatment (Lechner et al. 2007). For some individuals these symptoms can extend until menopause and be re-experienced when other parenthood milestones (e.g., grandparenthood) are not achieved (Daniluk 2001, Wirtberg et al. 2007). Cognitive Behavioural models of grief (Boelen et al. 2006) explain the recurrence of grief symptoms by the difficulty individuals have to integrate the loss into their autobiographical knowledge. These models also claim that grief reactions are augmented by negative beliefs about the loss, for instance the idea that not having a(nother) child will threaten the partnership. Finally, as was observed in the Daniluk et al. (2007) study, avoidance seems to
be associated with grief symptoms (Boelen et al. 2006), and it is known that individuals experiencing undesired childlessness tend to avoid the ‘fertile world’ (Volgsten et al. 2010).

Overall these theories highlight the benefits of integrating the loss of parenthood goals and focusing on other goals, and the central role that cognition has in this process. Other theories of adjustment to stressful life events also highlight that individuals strive to find meaning in crises and to regain self-worth and a sense of control over their lives (e.g., Taylor 1983). However, none of these theories fully captures the specificities of the parenthood goal that influence how individuals adjust to its loss. For instance, the loss is never definitive until women achieve menopause, it is invisible, in the sense that there are no exterior signs of the loss (e.g., no body), it centres around a goal that was not yet achieved and it is often a loss that is shared with one (partner) or more (family) people. We argue that added value will be gained from developing a model that describes the specific psychological mechanisms that underlie adjustment to unmet parenthood goals and identifies unique risk factors for maladjustment in this context. Such model can set the basis for increasing awareness about the negative psychological impact of not realizing one’s parenthood goals and to guide the provision of psychosocial care in this context.

In sum, it is still unclear if individuals who experience failed fertility treatment need psychosocial care and what such support should entail. The goals of the present mixed-methods review were threefold. First, a meta-analysis was performed to investigate if individuals who experience failed fertility treatment present worse mental-health and wellbeing than individuals for whom treatment is successful. We distinguished between mental-health, which captures the absence of psychopathological symptoms, and wellbeing, considered to be a more holistic and subjective evaluation of one’s life satisfaction or happiness (Diener 2000). Second, moderation analysis were performed to investigate if group differences in adjustment vary with the type of comparisons performed, the percentage of children in the unsuccessful group, the follow-up period considered and the quality of the studies. Third, a review of the existing qualitative research was conducted based on thematic analysis (Thomas and Harden 2008) to extract information about the patients’ perceived experiences of this period. Based on the qualitative data, we made inferences about the underlying mechanisms of adjustment to unmet parenthood goals and the risk factors for maladjustment, which we present in the form of a comprehensive model.

Methods

Systematic search
The electronic databases of Medline, Embase, PubMed, PsychInfo and Web of Science were searched between 1978 (the year of the first IVF treatment performed) and December
2015 (inclusively) based on search terms for fertility treatment, defined using terminology from the International Committee for Monitoring Assisted Reproductive Technology and the WHO-revised glossary of Assisted Reproductive Technologies (e.g., artificial insemination, assisted reproduction, in vitro fertilization, IVF; Zegers-Hochschild et al. 2009), AND psychosocial adjustment (e.g., adjustment, adaptation, wellbeing, mental-health, quality of life) AND post-treatment (e.g., post, following, after). The detailed search strategy is presented in Table S1 of the supplementary material. The search was adapted for each database. Medical Subject Headings (MeSH) terms were used in PubMed. No restriction was made on the type (journal, conference paper or dissertation) or language of publication. All citations were transferred to EndNote (Thomson Reuters, San Francisco, CA, USA). The results were crosschecked with five articles that had been identified previously as being eligible for the review to ensure the search was suitable for its purpose. The reference sections of all identified articles were examined to identify other relevant manuscripts.

**Inclusion and exclusion criteria**

This systematic review included quantitative and qualitative studies. Studies were included if they reported on the psychosocial adjustment of a group of individuals (women, men or couples) who experienced failed fertility treatment and if assessments were conducted at least one year after the last treatment cycle. This is the period recommended to ensure patients have definitely finished treatment (Gameiro et al. 2013). Further, literature on stressful life events shows that people take on average 2 years to adjust (Bonanno 2004). Therefore, a minimum period of 1 year allowed us to focus on long-term reactions to loss and not on the immediate short-term distress. To be included quantitative studies had to report on the mental-health or wellbeing of individuals who experienced failed fertility treatment (i.e., failed group) and a control group. Qualitative studies had to report on a group of individuals’ stated experiences of psychosocial adjustment after failed fertility treatment.

AF screened the titles and abstracts and AF and SG independently screened the full articles. Any disagreements were resolved with discussion. Duplicate or secondary publications on the same sample reporting on the same outcome were excluded from the meta-analysis to avoid multiple publication bias. In these cases the paper with the most appropriate research questions for the review was prioritized. If this criterion did not apply, the publication reporting data for the largest sample was selected. Excluded studies were classified according to reason for exclusion (see Figure 1).
**Data extraction**

**Quantitative studies**

AF extracted data using a standardised protocol. To characterise the set of studies, we extracted data (where available) regarding the authors and country, type of fertility treatment (e.g. IUI, IVF), type and size of failed and control groups, whether the failed treatment group included individuals with children (adoption or spontaneous conception), years since the last treatment cycle, study design (cross-sectional or longitudinal) and outcomes reported (mental-health, wellbeing). AF also extracted the necessary quantitative data to include in the meta-analytical syntheses of group comparisons. SG reviewed data extraction and disagreement was resolved via discussion until consensus was reached.

**Qualitative studies**

SG extracted data using a standardised protocol. To characterise the set of studies, data was extracted (where available) regarding the authors, country and goal of the study, study sample and percentage with children, study design, years since treatment, and data collection and analytical method. The data extracted for analysis was the content of the results section of each study.

**Data synthesis and analysis**

**Quantitative studies**

To investigate group mean differences in mental-health and wellbeing between patients who experienced failed fertility treatment and an adequate control group two meta-analyses were carried out. Only one effect size was returned per study to create an independent set of effect sizes for each analysis. For mental-health we considered measures of anxiety, depression, mental-health, psychopathology or stress because all report on the participants' level of psychopathologic symptoms. We did not include studies reporting on the frequency of specific mental-health diagnosis (e.g., mood disorder) because frequency data cannot be pooled together with group means. For wellbeing we considered measures of general wellbeing, life satisfaction, quality-of-life, general health or vitality, because they capture participants' subjective evaluation of their positive functioning. For studies that reported on multiple measures for the same outcome, specific measures were prioritized. For mental-health, depression was prioritized because it is the measure that best captures the emotional response to the outcome of treatment (Verhaak et al. 2007), followed by mental-health, anxiety, distress and psychopathology. For wellbeing, satisfaction with life was prioritized because it is one of the most commonly used measures in the parenthood literature, followed by quality of life, wellbeing, general health and vitality. If the study reported data for couples, extraction of data took in consideration if the non-independence between male and female
data was addressed (Kenny et al. 2006). If it was not taken in consideration either the male or female data were used to ensure independence, women were prioritized because they are more negatively impacted by failed treatment (e.g., Verhaak et al. 2005).

A random effects model was adopted due to the heterogeneity of our studies (Borenstein et al. 2009). The $I^2$ index was used to assess the proportion of effect size variability that was due to non-chance factors (Higgins et al. 2003). $I^2$ values of 0%, 25%, 50% and 75% indicate no, low, moderate and high heterogeneity, respectively. The probability level of $P<0.05$ was used to indicate statistical significance. The primary outcome measure used in the meta-analyses was the standardized mean difference between the unsuccessful and control groups. Hedge’s $g$ correction was used to adjust for potential bias due to the small sample sizes found in our studies (Borenstein et al. 2009). Effect sizes were aligned across studies so that a negative standardized mean difference indicated poorer adjustment for the failed treatment group than the control group. Effect sizes of .20, .50 and .80 should be interpreted as small, medium and large, respectively (Cohen 1992).

To investigate the moderation effect of type of control group (based on treatment outcome, parenthood status), existence of children in the failed treatment group (no, yes) and years since last treatment cycle subgroup ($X^2$ test) and meta-regression (Z-test, unrestricted maximum likelihood) analyses were done. We used Comprehensive Meta-Analysis version 2.2.064 (Biostat, Englewood, USA).

**Qualitative studies**

Methods of qualitative reviews are emergent and still subject of debate. In this review we opted to follow the synthesis method described by Thomas and Harden (2008). It was designed to identify which interventions need to be developed to address a particular health issue by understanding it from the point of view of the people targeted. We conducted a 3-stage thematic analysis (Dixon-Woods et al. 2005). The first two stages were descriptive. First, SG extracted all results sections verbatim and SG and AM conducted an independent ‘line by line’ coding of the text to develop descriptors of post-treatment experiences. Two meetings were conducted to contrast and review the descriptors generated. Disagreement was resolved by discussion until the coders agreed if a descriptor should be changed, added or removed from the final list. Second, after a definitive list of descriptors was agreed on, SG developed first order descriptive themes to group the descriptors. The themes needed to capture something important about the data in relation to the research question, representing some level of patterned response across the studies’ data (Braun and Clarke 2008). We only considered themes that appeared more than twice across all results and were present in at least two different papers. AF reviewed this process and disagreements were solved by discussion until both authors agreed on the grouping of descriptors and the labels of the second order themes. The last stage was interpretative. SG generated second order
analytical themes that were sufficiently abstract to include or explain all first order themes.

This process aimed to create interpretative constructs and testable hypothesis (Thomas and Harden 2008) and constituted the crucial step to develop an explanatory model of adjustment after failed treatment. Finally, SG wrote a narrative summary and developed a graphical depiction of the data. AF reviewed SG’s analysis and a similar discussion meeting was conducted to reach agreement on the themes and on their graphical and textual depiction. The percentages of agreement in the coding of the first and second order themes were calculated.

**Risk of bias assessment**

**Quantitative studies**

AF and SG designed a critical appraisal form to assess study quality in terms of population representativeness of the study sample (3 points), measurement validity (1 point) and reliability (1 point), and evaluation of research hypotheses (3 points). These criteria are described in detail in Table S3 of the supplemental material. Each study quality score could range from 0 to 8. Papers were classified into low [≤2], moderate [3-5] or high [≥6] quality studies. AF and SG independently assessed study quality using the form and disagreement was resolved via discussion until consensus was achieved. Subgroup analyses were performed to investigate if the pooled estimates varied as a function of study-quality.

Publication bias was checked with visual inspection of funnel plots (Hedge’s g against its standard error) and Egger’s test (Egger et al. 1997). Trim and fill was used to adjust the standardized mean difference between the failed and control groups for the presence of publication bias (Duval and Tweedie 2000).

**Qualitative studies**

Two possible sources of bias were considered: the quality of the primary research and the researchers. The quality of primary research was assessed using the Critical Appraisal Skills Programme Qualitative Research Checklist (10 criteria, Critical Appraisal Skills Programme (CASP) 2013), [http://media.wix.com/ugd/dded87_29c5b002d99342f788c6ac670e49f274.pdf](http://media.wix.com/ugd/dded87_29c5b002d99342f788c6ac670e49f274.pdf) (see Table S7 of supplementary material). Based on the number of met criteria, each study was classified as being of low [≤3], moderate [4-7] or high [≥8] quality.

As in any qualitative analysis, the researchers’ previous understandings may have shaped their interpretation of the data. SG has been researching the psychosocial issues of infertility for around 10 years and has in-depth knowledge of the field literature, including of the post-treatment period. AF is a medical student with no knowledge on this topic prior to the start of the study. None of the authors has regular contact with individuals from the study population. In order to increase the trustworthiness of the findings and reduce possibility of researcher bias, AF and SG independently assessed study quality and disagreement was resolved via
discussion until consensus was reached. The themes had to be closely linked to the data collected (Braun and Clarke 2008). As a final step, the findings were presented to a group of five infertility researchers and to a group of six patients and revised according to the feedback received.

Results

Description of studies

Figure 1 presents the study decision-flow chart whereby 18 studies were included in the mixed-methods systematic review. The systematic search returned 6397 non-duplicated records. Ninety-eight per cent of these studies (n = 6265) were excluded based on their title and abstract, which reduced the number of eligible studies to 132. From these 15 were retained after inspection of the full text and three further studies were identified through the inspection of references lists. The three most common reasons for exclusion were that assessment was done during the first 12 months after treatment, the paper did not report on primary research or the paper did not focus (nor reported) on psychosocial adjustment (see Table S2 of supplementary material for reasons for exclusion). During this process we contacted seven authors to clarify about missing or inconsistent data. Only two replied with the required information.

Table 1 presents the characteristics of the nine included quantitative studies (designated in the reference list by asterisks). The studies sampled 2278 individuals with failed fertility treatment from seven countries. All participants had undergone IVF but some also underwent other treatments including ICSI and gamete intrafallopian transfer (GIFT). Two types of control group were used, based on the outcome of treatment (successful, 33%) or on parenthood status (with children, either adopted or conceived spontaneously or with treatment, 67%) at time of assessment. The time since last treatment varied from one (12 months) to 20 years. Two studies (22%) had a longitudinal design. Only three studies (33%) had a response rate of 80% or higher. Six (67%) studies reported on mental-health and six on wellbeing.

Quality ratings for the studies are also shown in Table 1. Study quality ratings indicated no low, seven moderate (78%) and two high-quality studies (22%) (See Tables S3 to S6 in
supplementary material for details on quality assessment). The percentage of agreement between SG and AF on quality ratings was 93%.

Table 2 presents the characteristics of the nine included qualitative studies (designated in the reference list by asterisks). The studies sampled 267 individuals who experienced failed treatment from six countries. In six studies (67%) the sample included people with children. Eight studies (88%) were cross-sectional and one (12%) was longitudinal with three follow-up assessments occurring more than one year after treatment (Daniluk 2001). Data was collected via interviews and the most common analytical technique was phenomenological analysis (n = 4, 44%), but thematic, content and discourse analysis and grounded theory were also used.

Quality ratings for the qualitative studies are also shown in Table 2. Study quality ratings indicated no low, two moderate (22%) and seven high-quality studies (78%) (See Table S7 in supplementary material for details on quality assessment). The percentage of agreement between SG and AF on quality ratings was 83%.

Quantitative meta-synthesis: Psychosocial adjustment after failed fertility treatment

Figure 2 shows the pooled standardized mean differences in mental-health and wellbeing between the failed and control groups for the random effects model. The meta-analysis showed that the failed treatment group had significantly worse mental-health than the control group (g = -0.496, P=0.001, 95%CI [-0.791 to -0.200]), with significant heterogeneity between studies (I²=86%, P<0.001).

The Egger’s test indicated the presence of publication bias (intercept= -3.192, t = 4.85, P = 0.004), which was confirmed through visual inspection of the funnel plot. The trim and fill method also confirmed the presence of publication bias. This method identified one missing study, estimating an adjusted point estimate of -0.458 that was still significant (95% CI -0.726 to -0.189).
Table 3 presents the standardized mean differences for mental-health and wellbeing between the failed and control groups for the pre-defined moderators. Subgroup analysis for the type of group comparisons done showed that mean group differences were bigger when the control group is based on the outcome of treatment than when it is based on parenthood status. The remaining subgroup and meta-regression analyses were statistically non-significant.

### Wellbeing

Six studies reported data on group mean differences for at least one wellbeing measure that could be included in the meta-analysis. Life satisfaction scores were considered for three studies, quality of life for two (Weaver et al. 1997, Wischmann et al. 2012) and wellbeing (Johansson et al. 2010) for one. The meta-analysis showed that the failed treatment group had significantly worse wellbeing than the control group (g = -0.415, P<0.001, 95%CI [-0.627 -0.203]), with non-significant heterogeneity between studies (I²=53%, P=0.059). Visual inspection of the forest plot (see Figure 1) suggests that the study by Leiblum and colleagues (1998) can be considered an outlier (its effect size was more than twice the pooled standard deviation distant from the pooled effect size). Therefore we repeated the meta-analysis excluding this study. The meta-analysis still showed that the failed treatment group had significantly worse wellbeing than the control (g = -0.324, P<0.001, 95%CI [-0.454 -0.193]), with no heterogeneity between studies (I²=0%, P=0.552). Subsequent analyses were run without including this study.

The Egger’s test indicated no presence of publication bias for wellbeing (intercept = -0.523, t= 0.428, P = 0.349). The trim and fill, however, identified one missing study, estimating an adjusted point estimate of -0.298 that was still significant (95% CI -0.422 -0.175).

Subgroup and meta-regression analyses could only be performed for the moderators existence of children in the failed treatment group and years since treatment and none were statistically significant.

### Qualitative meta-synthesis: Patients’ perceived experiences of the period after failed treatment

To illustrate the analysis process, please consider the following lines of text presented in the study by Boden (2007): “On a positive note, many couples expressed their belief that the experience of treatment had enhanced their relationship with their partner, bringing them, as participants described, ‘closer together’”. 
To summarize these data the following descriptor was developed: ‘treatment strengthens the relationship’. After, this and other similar descriptors, for instance ‘stronger partnership as result of infertility and treatment experience’ and ‘treatment strengthens the marital relationship and leads to marital satisfaction’, were grouped into the first order theme ‘treatment strengthens partnership’. Finally, this and other first order themes, for instance, ‘intense grief’, ‘anger directed at everything and everyone’, ‘sexual difficulties’ and ‘men stoic and supportive of their partner’ were classified into the second order theme ‘Individual and relational adjustment’. Table S8 of the supplemental material presents, for each study included, the list of descriptors of post-treatment experiences and their classification into first and second order themes. Inter-coder agreement for 1st and 2nd order themes was 83.3% and 92.2%, respectively.

A total of 28 first order themes and six second order themes were developed and are presented in Table 4. Table S9 presents all first order themes developed, the number of times they were cited in the results sections of the studies included, the number of studies citing it, and its association with the second order themes. In total 33 first order themes were developed, however, as can be seen in table S9, five were only cited once and were therefore not considered.

Table 4 around here

A comprehensive description of the themes is provided below. The themes Individual and relational adjustment, Social adjustment and Fertility care perceptions and needs capture changes across time in experiences and adjustment. The themes Acceptance, Meaning making and Pursuit of new life-goal capture the psychological tasks patients engaged with that were associated with positive adjustment in the aftermath of failed treatment.

Individual and relational adjustment. Participants described the period immediately after ending treatment as of intense grief (pain, sadness and emptiness) and anger felt towards ‘everything and everyone’. Grief endured in time and was lived in isolation. The grief experience was different for women and men. For women it was very intense and characterized by a sense of emptiness, meaningless, loss of control and self-blame. Women tended to be open about their grief with their family and friends. Men felt pain, tried to overcome grief by denying it and tended to be private in their suffering. Men were surprised with the intense grief reactions of their (female) partners and felt it was their duty to be stoic and supportive. However, they felt they lacked the knowledge to do this. In addition, most participants reported that the treatment experience strengthened their partnership. However, the sexual desire of many couples was affected during treatment and it did not recover once treatment was over. Despite the initial difficulties and grief, as time passed participants
reported a sense of survival and personal and spiritual growth that was associated with having undergone the treatment and the childlessness experience.

*Social adjustment.* Most participants experienced social isolation. Isolation was the result of not meeting parenthood social norms and therefore being unable to share parenthood experiences with peers. Men also felt isolated from family members, with whom they felt they communicated less due to not having children. The absence of children was a loss experienced by the whole family and raised concerns about not leaving a legacy to continue the family bloodline and about loneliness in later life. Many women talked about how social constructions and pressure for parenthood made it harder for them to accept their situation. For instance, women’s identities relied so strongly on parenthood that not having the desired children made them doubt their femininity or feel abnormal. Some women felt the need to show others they did everything they could to become parents to avoid social scrutiny. Others talked about how it was difficult to escape society’s expectations. Overall participants felt that not meeting their parenthood goals hindered their relationship with their social network. Despite the overall isolation, participants also described choosing to share their struggles with a selected few close friends and work colleagues who provided them with good emotional and instrumental support. Other people with infertility also provided them with useful information. Finally, for women social isolation also resulted from active efforts to avoid situations where children were present in order to escape suffering.

*Fertility care perceptions and needs.* Participants said that when all treatment options were exhausted they felt abandoned by their clinic. They perceived a lack of support from the clinic in both deciding if and when ending treatment and once this decision was made. They expressed a need for psychosocial care or counselling to help them cope with the intense grief they experienced, and for sexual counselling to help them overcome the sexual difficulties that developed during treatment. Participants acknowledged the limitations of reproductive medicine and this facilitated acceptance of their situation. Reaching acceptance was associated with developing a positive outlook on ending treatment, for instance as allowing participants to let go of their medical experience, as actively managing their child-desire and as allowing others to also have an opportunity to achieve parenthood.

*Meaning making.* In the process of coming to terms with their unmet parenthood goals, participants engaged in an intense re-evaluation effort whereby they tried to make sense of their past efforts to achieve parenthood and they re-evaluated their life values and priorities in search for a new identity and future in the absence of the desired children. Traditional notions of marriage (e.g., marriages do not survive without children), femininity (e.g., women are born to be mothers) and family (e.g., children are part of family life) hindered this meaning making process. Therefore many participants questioned such beliefs and reflected on the meaning of marriage and family without children. This critical approach to social
constructions of parenthood facilitated acceptance. Overall participants felt that meaning making was associated with higher acceptance of one’s unmet parenthood goals.

**Acceptance.** Immediately after finishing treatment participants maintained a strong hope to have (more) biological children and felt unwilling or unable to accept their situation. Acceptance was a process that unfolded over time because of the need to ‘move on’ with life. Many patients differentiated rational from emotional acceptance and stated that although rationally they accepted their situation, this was not synonymous with or followed by emotional acceptance. Indeed, the hope to have the desired children could persist for many years after treatment and for some women it only ended when they reached menopause, as menstruation meant that a pregnancy was still possible. Acceptance was easier when patients were convinced they did their best and explored all available treatment options, when their efforts were acknowledged by their family and friends and met with empathy instead of pressure for them to become parents. Acceptance was also easier when participants felt they received good medical advice during treatment, when their prognosis was clear and poor and when the reasons for treatment failure were identified. Acceptance was associated with a change in the participants’ view of the world. This translated in recognizing their limits to control reality, in the development of a more realistic view of the world or in increasingly valuing the ‘good things’ in life. Acceptance was also associated with better adjustment. It translated in a sense of freedom from the infertility experience and hope towards the future. This new state was described as of restored strength and agency, and equilibrium with both the self and in the partnership. In contrast, the inability to accept their situation was associated with maladjustment, in particular with continuous grief, feeling ‘stuck in life’, low self-esteem and different avoidance strategies.

**Pursuit of new life-goals.** When ending treatment participants felt incapable of envisioning themselves as anything else than parents. For most participants the need to move on with their lives resulted from a desire to overcome the pain of grief and to reject self-pity and continuous mourning. Coherently, the willingness to ‘move on’ was lower when there was a stronger desire for children. Overall participants perceived that the needs to accept their situation and ‘move on’ were interrelated. Participants’ typical strategy was to invest in new goals and activities that distracted them from their pain, compensated for their childlessness status and boosted their self-esteem. Despite desiring to ‘move on’, many participants did not know how to do this and expressed that the process required an active effort to look into the future, which implied a confrontation with later life stages and their lack of legacy. Participants talked about many different ways of finding new fulfilling life goals and activities and these included caring for other people (e.g., family members) and pets, supporting friends with infertility and advocating for the rights of other infertile people, travelling and focusing on their career and, when full acceptance was achieved, some individuals even
refocused on caring for other children (e.g., siblings). Overall participants felt that the pursuit of new life-goals was associated with higher acceptance of their unmet parenthood goals.

**Discussion**

Findings from this mixed-methods systematic review show that people who do not manage to conceive with fertility treatment experience worse mental-health (i.e., more psychopathologic symptoms) and wellbeing (lower positive functioning and life satisfaction) than those who conceive. Positive adjustment in this context seems to be associated with a progressive acceptance of the loss; making meaning of past efforts to conceive and the present situation; and pursuing new life goals or activities. Individuals who engage with these three psychological tasks tend to report a sense of personal growth and to develop a positive outlook on ending treatment and renewed views of the world. The results reported suggest that the risk for maladjustment varies according to individual, social and treatment related factors. Overall the data indicate that the post treatment period is worthy of clinical attention and reinforce advice for health professionals to support individuals during this period.

Long-term adjustment after failed fertility treatment is an emergent research topic within reproductive psychology. Only 18 studies could be identified for inclusion with an even split between quantitative and qualitative research. An examination of the papers excluded (see Figure 1) indicates that researchers tended to focus on the first 12 months following treatment (only capturing short term reactions), did not report on time since treatment, focused on topics either than adjustment, or, if they focused on adjustment, did not use a control group. From the included studies only two reported on correlates or predictors of adjustment (Gameiro et al. 2016, Verhaak et al. 2007) making it impossible to quantitatively investigate risk factors. In sum, research on this topic is still at a descriptive state that needs to be moved towards an explanatory theory-driven approach. This mixed-methods review is timely in that it provides the evidence base for the development of testable hypotheses organized in a comprehensive theoretical model.

Results from the meta-analysis show that individuals who do not manage to conceive with fertility treatment report significantly worse mental-health (medium effect sizes) and wellbeing (small to medium effect sizes) than those who do. Moderation analyses focusing on the type of control group and percentage of children in the failed treatment group suggest that the poorer adjustment of the failed group is better explained by their inability to meet their parenthood goals than their childless status. Indeed, group differences in mental-health were larger when based on the treatment outcome than on parenthood status (-0.604 versus -0.108) and the existence of children in the failed treatment group did not prove to be a significant moderator for mental-health nor wellbeing. These data suggests that what is associated with worse adjustment is having less children than desired or, in other words, not
meeting ones parenthood goals. Recent research shows that both the population of people with undesired childlessness and fewer children than desired are increasing as a result of parenthood delay (Schmidt et al. 2012).

Contrary to our hypothesis, group differences in mental-health and wellbeing did not subside within the 20 years time period surveyed. However, this result needs to be carefully considered because it may be explained by a methodological limitation in the operationalization of time since treatment. Indeed, seven out of the nine studies included provided a range for time since last treatment (instead of a specific time or average) and this could span several years. In these cases we had to use the mean and this is likely to have obscured associations.

Findings from the qualitative review indicate that positive adjustment is dependent on individuals accepting and making meaning of their unmet parenthood goals and pursuing new life-goals. Individuals perceived these as three separate but inter-dependent psychological tasks. Acceptance was portrayed as a process (and not an outcome). It appears to involve the emotional processing of one’s unmet parenthood goals. According to psychological theories, this implies the activation and restructuring of maladaptive fear structures, exposure and habituation to the situation and regulation of the negative affect it elicits (Park 2010). Therefore, acceptance may be fostered by emotional approach coping to the loss (Boelen et al. 2006, Stanton et al. 1994) so that, in accordance with grief models, its integration in autobiographical knowledge is achieved (Boelen et al. 2006). However, unmet parenthood goals can be an ambiguous loss because the chances of achieving parenthood are rarely completely null. This may preclude the integration of the loss. Indeed, our qualitative data showed that acceptance was easier when chances to conceive were lower (poor prognosis, clear advice to stop treatment from fertility team, reasons for failure identified), or in other words, when the loss was perceived as definitive.

Meaning making seems to reflect an adaptive cognitive processing of the situation. The literature about adaptation to stressful life events or trauma (e.g.; Park 2010, Taylor 1983) considers meaning making as a critical task for adjustment and states that individuals will always engage in it in order to restore a sense of control and predictability over their lives (Kelley and Michela 1980). Our results seem to be consistent with this literature in that patients perceived meaning making and acceptance to be related and in that acceptance was associated with new views of the world and redefinition of priorities in life (Park 2010).

Grief theorists also claim that such cognitive processing needs to address dysfunctional beliefs associated with the loss (Boelen et al. 2006). For instance, individuals talked about guilt related with ending treatment and concerns about their role in their family. Previous research also highlighted concerns about the impact of not achieving parenthood on the partnership and one’s ability to endure the related grief (Boivin et al. 2005). In this context,
self-blaming and catastrophizing were shown to be negatively associated with adjustment (Kraaij et al. 2009). On the contrary, positive reappraisal coping may assist meaning making and was associated with positive adjustment (Kraaij et al. 2009, Peterson et al. 2009).

Finally, results suggest that the pursuit of new life goals is associated with higher acceptance of the unmet parenthood goals and, in the long-term, better adjustment. Individuals claimed that pursuing new life goals required an active effort from their part. According to the dual process model of grief, this can be regarded as an effort to process the implications of the loss (Stroebe and Schut 1999) by finding new purpose(s) for living and regaining a sense of agency. A recently published meta-analysis showed that, in the context of a parenthood blockage, reengagement in new life goals is associated with better wellbeing (i.e., higher positive mood; Mesquita da Silva et al. 2016). Our results support these data, but also show that engaging with new life goals is harder for individuals who highly value parenthood or perceive a lack of other meaningful goals to pursue (Thoits 1992). Our data also show that individuals start by pursuing other goals just as a distraction from the suffering of not achieving their main goal of parenthood, but that they end up perceiving benefits from this. These results are consistent with literature showing that pursuing new goals is adaptive even when individuals have not completely disengaged from parenthood (Wrosch et al. 2003).

Another important finding was how the unmet parenthood goals affected individuals’ social relationships and how their social context influenced their adjustment process. This is not surprising as it is well known that the socio-cultural context has a strong influence in shaping the experience of infertility and childlessness of both women and men (Greil et al. 2011, Martins et al. 2016). Social isolation was a common theme in the data, either resulting from less empathic reactions from the social network or from self-avoidance of that social network, when it implied contact with children. Social support is a consistent protective factor in the context of infertility and its treatment (Martins et al. 2016, Rockliff et al. 2014). Our data was no different in this regard, but also showed that the social context could influence adjustment via other routes than support, more specifically via social norms (i.e., social pressure for parenthood) and prevalent representations of parenthood, marriage and femininity. Overall, the pervasiveness of social and cultural first order themes suggests that psychosocial care needs to include a thorough assessment of the social context and provide individuals with strategies to cope with it.

Strengths and limitations

The strengths of this mixed-methods review were the systematic review of almost 40 years of research on adjustment after failed fertility treatment, which yielded 18 studies from ten countries sampling adjustment experiences of 2545 patients. Data were independently
extracted and quality evaluations and analyses were made according to a-priori defined and
drigorous protocols in order to minimize the risk of bias. Another strength was the inclusion of
quantitative and qualitative research to answer different research questions, allowing for a
more comprehensive picture of adjustment to emerge. The primary research was of
moderate to high quality. In quantitative studies quality was due to the fact that most studies
had a representative sample of the population, used valid and reliable outcome measures
and had enough power to detect significant differences. In qualitative studies the quality was
due to the adequacy of the design, recruitment strategy and data collection method used.
However, the quantitative studies did not focus on investigating risk factors for
maladjustment and used heterogeneous samples of individuals in terms of time since
treatment and parenthood status. It was also not totally clear if those individuals who did not
reach parenthood via treatment were still pursuing it, for instance by doing more treatment at
different clinics or by choosing to adopt. These data, together with previous research (e.g.,
Gameiro et al. 2014, Wischmann et al. 2012), suggest that the larger the proportion of the
sample still pursuing parenthood the worse adjustment will be. The main problem of the
qualitative data was that, due to the nature of qualitative research, the data extracted from
the papers were interpretations that other researchers made of patients’ interpretations of
their experiences. Another problem was that the goals of the qualitative studies varied
slightly, for instance one study focused only on the perceived gains of doing fertility treatment
(Lee et al. 2009). This might have affected the prevalence of the emergent themes.
Despite these limitations, overall the results from the quantitative and qualitative review
were consistent and its integration provided a coherent picture of adjustment after failed
treatment. The only inconsistency detected was that qualitative data provided a more positive
perspective on adjustment over time than the quantitative data. However, we explained that
the quantitative data might not be reliable. Furthermore, it should be noted that the
quantitative data tapped into hedonic wellbeing, which is the extent to which individuals are
‘feeling well’, and the qualitative data also captured eudaimonic wellbeing, which is the extent
to which individuals are ‘doing well’ in terms of purpose, meaning and fulfilment in life (Ryan
and Deci 2001). By analysing both types of data we achieved a more nuanced view of
wellbeing, which can explain the discrepancy observed.

Conclusions
Based on the data reported we developed a comprehensive model to explain adjustment
to unmet parenthood goals, labelled the Three Tasks Model of Adjustment to Unmet
Parenthood Goals and presented in Figure 3. The model predicts that, in the face of unmet
parenthood goals, individuals able to accept their situation, make meaning of their
experience and pursue new life goals will adjust better than those who do not engage with
these psychological tasks. These three main predictions are consistent with knowledge from
other theories previously used to explain adjustment after failed treatment and general
theories of adjustment to stressful life events. The Three Tasks Model of Adjustment to
Unmet Parenthood Goals also predicts that specific variables may influence adjustment,
either directly (e.g., women are more likely to present worse adjustment), by impacting on the
individual's ability to perform the three psychosocial tasks (e.g., one is less likely to pursue
new goals if they do not find alternative meaningful goals), or by moderating the relationship
between the three psychological tasks and adjustment (e.g., pursuit of new life goals is more
likely to lead to better adjustment when one is supported by their network than when is not
supported). The variables identified were, at the individual level, gender, importance of
parenthood and availability of alternative life goals; at the social level, social support, social
representations of parenthood and parenthood social norms; and at the treatment level,
prognosis, perceived quality of medical advice, reasons for failure identified and treatment
options exhausted.

In this review we considered two adjustment outcomes: mental health and wellbeing. The
meta-analysis showed that these are sensitive measures in this context and should therefore
continue to be used in future research. The qualitative data suggests that patients assess
their adjustment not only in terms of how they feel (hedonic wellbeing, e.g. intense grief,
anger) but also in terms of how they think they are realizing their human potential
(eudaimonic wellbeing, e.g. resilience and personal and spiritual growth; Ryff 1995). Future
research could consider using outcomes that tap into eudaimonic wellbeing. This seems
particularly important in this context because the pursuit and realisation of intrinsic goals
seems to be more strongly related with eudaimonic than hedonic wellbeing (Ryan and Deci
2001). Examples of eudaimonic outcomes would be self-acceptance (positive evaluations on
oneself and one’s past life, personal growth (sense of continued growth and development as
a person) or purpose in life (the belief that one’s life is purposeful and meaningful (Ryff and
Keyes 1995).

In conclusion, this mixed-methods review offers compelling evidence for the need to
provide psychosocial care directed at helping individuals relinquishing their parenthood
goals. Patients themselves recognize the need for psychosocial care during this period and
express frustration at feeling abandoned by their clinic. The Three Tasks Model of
Adjustment to Unmet Parenthood Goals offers comprehensive guidance on the therapeutic
mechanisms that seem to underlie positive adjustment. The model was developed based on
the experiences of patients who experienced failed fertility treatment, but there is nothing to
suggest that it cannot be applied to everyone who is confronted with the loss of their
parenthood goals, regardless of the pathway that led to it. Future research should now test
the model in these different contexts. This can be done by conducting prospective cohort
studies to test the model predictions. Another way to test the model is by developing and testing psychosocial interventions that are based on its three main hypothesis, in other words, interventions designed to engage individuals with acceptance, meaning making and pursuit of new goals. An important question in this context is how such specialized care can reach individuals who are already outside the reproductive healthcare system. The ESHRE guideline argues that this is the responsibility of fertility clinics (Gameiro et al. 2015). We concur that specialized expertise is needed to develop and test effective interventions. A sensible approach would be to develop self-help interventions that individuals could access via multiple outlets (the most obvious one being the internet) instead of having to go back to their fertility clinics. Another crucial approach is to promote knowledge transfer about the topic so that primary care physicians and mental-health professionals are aware of the deleterious impact of unmet parenthood goals on wellbeing and of how it can be addressed.

**Author's roles**

S.G. did data extraction, critical appraisal, data analysis and interpretation and writing of the report. A.F. did data extraction, critical appraisal, data analysis and interpretation. All authors approved the final version for submission.

**Acknowledgements**

We would like to thank our colleagues Professors Jacky Boivin and Christianne Verhaak and Dr Emily Koert for their careful reviews of draft versions of the present paper.

**Funding**

This work was supported by the UK Economic and Social Research Council [grant number 508860].

**Conflict of interest**

There are no conflicts of interest.

References


Thomas, J. and A. Harden. Methods for the thematic synthesis of qualitative research in systematic reviews. *BMC Medical Research Methodology*. 2008: 45;

Throsby, K. No one will ever call me Mummy: Making sense of the end of IVF treatment. 2001. London School of Economics - Gender Institute, London.


Table 1. Characteristics of the nine included quantitative studies

<table>
<thead>
<tr>
<th>Authors, year Country</th>
<th>Fertility treatment</th>
<th>Failed treatment group</th>
<th>Control group</th>
<th>Existence of children in failed treatment group (YES, NO)(^a)</th>
<th>Years since last treatment cycle</th>
<th>Design</th>
<th>Response /attrition rate</th>
<th>Quality</th>
<th>Mental-Health Wellbeing(^b)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gameiro et al 2014 The Netherlands</td>
<td>Fertility drugs/ IUI/ AI/ IVF/ ICSI/ gamete donation</td>
<td>1493 women</td>
<td>5655 women with children(^b)</td>
<td>NO</td>
<td>range 11-17</td>
<td>Cross-sectional</td>
<td>60.4%</td>
<td>High</td>
<td>✓</td>
</tr>
<tr>
<td>Hammarberg et al 2001 Australia</td>
<td>IVF/ ET/ GIFT</td>
<td>65 women</td>
<td>51 women successful treatment</td>
<td>YES</td>
<td>range 2.5-3.5</td>
<td>Cross-sectional</td>
<td>55%</td>
<td>Moderate</td>
<td>✗</td>
</tr>
<tr>
<td>Johansson et al 2010 Sweden</td>
<td>IVF/ ET/ ICSI</td>
<td>149 women + 121 men</td>
<td>118 women + 93 men with children</td>
<td>YES</td>
<td>range 4-5.5</td>
<td>Cross-sectional</td>
<td>67.5%</td>
<td>Moderate</td>
<td>✓</td>
</tr>
<tr>
<td>Kuivasari-Pirinen et al 2014 Finland</td>
<td>IVF/ ICSI</td>
<td>209 women</td>
<td>296 women successful treatment</td>
<td>YES</td>
<td>range 1.3-11.7</td>
<td>Longitudinal</td>
<td>100% / 54.7%</td>
<td>Moderate</td>
<td>✗</td>
</tr>
<tr>
<td>Leiblum et al 1998 USA</td>
<td>IVF</td>
<td>18 women</td>
<td>41 women successful treatment</td>
<td>NO</td>
<td>range 2-13</td>
<td>Cross-sectional</td>
<td>31%</td>
<td>Moderate</td>
<td>✓</td>
</tr>
<tr>
<td>Sydsjö et al 2015 Sweden</td>
<td>IVF</td>
<td>62 men</td>
<td>230 men with children</td>
<td>NO</td>
<td>20</td>
<td>Cross-sectional</td>
<td>66.5%</td>
<td>Moderate</td>
<td>✓</td>
</tr>
<tr>
<td>Verhaak et al 2007 The Netherlands</td>
<td>IVF/ ICSI</td>
<td>39 women</td>
<td>68 women successful treatment</td>
<td>NO</td>
<td>range 3-5</td>
<td>Longitudinal</td>
<td>84% / 78%</td>
<td>High</td>
<td>✓</td>
</tr>
<tr>
<td>Weaver et al 1997 UK</td>
<td>IVF/ GIFT</td>
<td>21 couples</td>
<td>20 couples successful treatment</td>
<td>YES</td>
<td>range 1.25-2.25</td>
<td>Cross-sectional</td>
<td>95%</td>
<td>Moderate</td>
<td>✓</td>
</tr>
<tr>
<td>Wischmann et al 2012 Germany</td>
<td>IUI /IVF/ ICSI</td>
<td>40 couples</td>
<td>91 couples successful treatment</td>
<td>NO</td>
<td>10 (since first consultation)</td>
<td>Cross-sectional</td>
<td>35.7%</td>
<td>Moderate</td>
<td>✗</td>
</tr>
</tbody>
</table>

Legend. SC = spontaneous conception; IVF = in-vitro fertilisation; ICSI = intra-cytoplasmic sperm injection; FET = frozen thawed embryo transfer cycle; IUI = intrauterine insemination; GIFT = gamete intrafallopian transfer; AI = artificial insemination; ET = embryo transfer; NR = not reported; \(^a\) children could be adopted or conceived spontaneously or with previous fertility treatment; \(^b\)✓ = reported; ✗ = not reported. Note: 180 (2.5%) patients who participated in the Gameiro et al 2014 study also participated in the Verhaak et al 2007. Due to the extremely low sample overlap we still included the two studies.
Table 2. Characteristics of the nine included qualitative studies

<table>
<thead>
<tr>
<th>Author, year</th>
<th>Country</th>
<th>Goal</th>
<th>Failed treatment group</th>
<th>% with children</th>
<th>Design</th>
<th>Years since treatment</th>
<th>Data collection method</th>
<th>Analytical method</th>
<th>Quality</th>
</tr>
</thead>
<tbody>
<tr>
<td>Boden 2007</td>
<td>UK</td>
<td>Extend the understanding of how women and their partners experience infertility and come to terms with their childlessness</td>
<td>18 couples + 15 women</td>
<td>NR</td>
<td>Cross-sectional</td>
<td>5</td>
<td>Interviews</td>
<td>Thematic analysis</td>
<td>High</td>
</tr>
<tr>
<td>Daniluk 2001</td>
<td>Canada</td>
<td>Understand how couples make sense of their infertility and reconstruct their lives when faced with the permanence of their biological childlessness</td>
<td>37 couples</td>
<td>46</td>
<td>Longitudinal</td>
<td>1, 1.83 &amp; 2.66</td>
<td>Semi-structured interviews</td>
<td>Phenomenological analysis</td>
<td>High</td>
</tr>
<tr>
<td>Johansson &amp; Berg 2004</td>
<td>Sweden</td>
<td>Describe women’s experiences of ongoing childlessness 2 years after the end of IVF treatment</td>
<td>8 women</td>
<td>13</td>
<td>Cross-sectional</td>
<td>2</td>
<td>Unstructured interviews</td>
<td>Phenomenological analysis</td>
<td>Moderate</td>
</tr>
<tr>
<td>Lee et al 2009</td>
<td>China</td>
<td>Provide an in-depth description of the gains perceived by Chinese men and women and how they re-constructed their lives after unsuccessful IVF treatment</td>
<td>4 couples (4 women + 4 men) + 6 women</td>
<td>36</td>
<td>Cross-sectional</td>
<td>range 0.5 – 3</td>
<td>Semi-structured interviews (joint and individual)</td>
<td>Grounded Theory</td>
<td>High</td>
</tr>
<tr>
<td>McCarthy 2008</td>
<td>USA</td>
<td>Understand the lived experience of infertility for women in the aftermath of unsuccessful medical treatment</td>
<td>22 women</td>
<td>32</td>
<td>Cross-sectional</td>
<td>3.9, SD 3.2</td>
<td>Interviews</td>
<td>Hermeneutic-phenomenology</td>
<td>High</td>
</tr>
<tr>
<td>Su &amp; Chen 2006</td>
<td>Taiwan</td>
<td>Explore the lived experiences of infertile women who terminated treatment after IVF failure</td>
<td>24 women</td>
<td>NR</td>
<td>Cross-sectional</td>
<td>1.33, SD 0.33 range 1-1.92</td>
<td>Telephone interviews</td>
<td>Phenomenological analysis</td>
<td>High</td>
</tr>
<tr>
<td>Throsby 2001</td>
<td>UK</td>
<td>Identify the factors which inform the decision to end treatment; consider the ways in which the unsuccessful engagement with IVF marked both the experience of infertility and the participants’ own perceptions of that technology; explore the ways in which those feelings might change over time and the influence they might have on subsequent choices</td>
<td>13 couples + 15 women</td>
<td>54</td>
<td>Cross-sectional</td>
<td>2</td>
<td>Interviews</td>
<td>Discourse analysis</td>
<td>High</td>
</tr>
<tr>
<td>Volgsten et al 2010</td>
<td>Sweden</td>
<td>Explore the experience of undergoing unsuccessful in vitro fertilization (IVF) treatment and of remaining childless 3 years after IVF in both women and men</td>
<td>7 couples + 3 women + 2 men</td>
<td>21</td>
<td>Cross-sectional</td>
<td>3.17</td>
<td>Semi-structured interviews</td>
<td>Content analysis</td>
<td>High</td>
</tr>
<tr>
<td>Wirtberg et al 2007</td>
<td>Sweden</td>
<td>Obtain an increased knowledge and deeper insight into the long-lasting effects of, and coping with, involuntary childlessness for a group of women who had sought help for infertility and had completed infertility treatment over 20 years before</td>
<td>14 women</td>
<td>0</td>
<td>Cross-sectional</td>
<td>20</td>
<td>Semi-structured interviews</td>
<td>NR</td>
<td>Moderate</td>
</tr>
</tbody>
</table>

Legend. IVF = In Vitro Fertilization; NR = not reported.
Table 3. Subgroup and meta-regression analyses of mean differences between the failed treatment and control groups for mental-health and wellbeing.

<table>
<thead>
<tr>
<th>Subgroup analysis</th>
<th>Mental-health</th>
<th>Wellbeing</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Effect size</td>
<td>95% CI</td>
</tr>
<tr>
<td></td>
<td>k</td>
<td>LL</td>
</tr>
<tr>
<td>Comparison groups</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Based on treatment outcome (failed, successful)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Based on parenthood status (no children, children)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Existence of children in failed treatment group</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Based on treatment outcome (failed, successful)</td>
<td>4</td>
<td>-0.648</td>
</tr>
<tr>
<td>Based on parenthood status (no children, children)</td>
<td>2</td>
<td>-0.108</td>
</tr>
<tr>
<td>Quality</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Moderate</td>
<td>4</td>
<td>-0.608</td>
</tr>
<tr>
<td>High</td>
<td>2</td>
<td>-0.323</td>
</tr>
<tr>
<td>Meta-regression</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Years since treatment</td>
<td>6</td>
<td>0.025</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Legend. *P<0.05, **P<0.01, ***P<0.001, ART = assisted reproductive techniques, k=number of studies, CI = confidence intervals, LL = lower limit UL = upper limit, NA = not applicable as there were not enough studies in the group, bold indicates P<0.05
Table 4. First and second order themes developed to describe patients’ experiences after failed fertility treatment. Second order themes are presented in bold with a grey background. For each of the second order themes the associated first order themes are listed below.

<table>
<thead>
<tr>
<th>Individual and relational adjustment</th>
<th>Social adjustment</th>
<th>Fertility care perceptions and needs</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Immediately after ending treatment</strong></td>
<td>• Intense grief (especially for women)</td>
<td>• Feeling abandoned\lack of support from clinic</td>
</tr>
<tr>
<td></td>
<td>• Anger directed at ‘everything &amp; everyone’</td>
<td>• Need for psychosocial &amp; sexual care/counselling</td>
</tr>
<tr>
<td></td>
<td>• Treatment strengthens partnership</td>
<td>• Acknowledgement of reproductive medicine limitations</td>
</tr>
<tr>
<td></td>
<td>• Men stoic &amp; supportive of their partner</td>
<td><strong>As time since treatment passed</strong></td>
</tr>
<tr>
<td></td>
<td>• Sexual difficulties</td>
<td>• Positive outlook on ending treatment</td>
</tr>
<tr>
<td><strong>As time since treatment passed</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>• Resilience &amp; personal &amp; spiritual growth</td>
<td>• Social isolation</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Parenthood loss hinders partner and social relationships</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Social representations of parenthood hinder adjustment</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Support from selected family members &amp; friends</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Social avoidance</td>
</tr>
<tr>
<td></td>
<td>• Social isolation</td>
<td>• Immediate after ending treatment</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Parenthood loss hinders partner and social relationships</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Social representations of parenthood hinder adjustment</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Support from selected family members &amp; friends</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Social avoidance</td>
</tr>
<tr>
<td><strong>Meaning making</strong></td>
<td><strong>Acceptance</strong></td>
<td><strong>Pursuit of new life-goals</strong></td>
</tr>
<tr>
<td>• Need to find meaning in life (past efforts, new identity &amp; future)</td>
<td>• Immediately after ending treatment</td>
<td>• New fulfilling goals/activities include…</td>
</tr>
<tr>
<td></td>
<td>• Questioning of traditional notions of marriage, family… in light of parenthood loss</td>
<td>• Inability/unwillingness to accept childlessness</td>
</tr>
<tr>
<td></td>
<td>• Re-evaluation of life values &amp; priorities</td>
<td><strong>As time since treatment passed</strong></td>
</tr>
<tr>
<td></td>
<td>• Acceptance and meaning making are interdependent</td>
<td>• Acceptance easier when…</td>
</tr>
<tr>
<td></td>
<td></td>
<td>[all options explored, efforts acknowledged, no pressure for parenthood, good medical advice, poor prognosis, reasons for failure identified]</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Acceptance associated with new views of the world</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Non-acceptance associated with worse adjustment</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Acceptance associated with better adjustment</td>
</tr>
<tr>
<td></td>
<td>• Need to find meaning in life (past efforts, new identity &amp; future)</td>
<td>• ’Moving on’ requires active effort</td>
</tr>
<tr>
<td></td>
<td>• Questioning of traditional notions of marriage, family… in light of parenthood loss</td>
<td>• Need to ‘move on’ with life</td>
</tr>
<tr>
<td></td>
<td>• Re-evaluation of life values &amp; priorities</td>
<td>• Acceptance and pursuit of new life-goals are interdependent.</td>
</tr>
</tbody>
</table>
Records identified through database search (n=8496)

Records after duplicates removed (n=6397)

Records screened (n=6397)

Records excluded based on title and abstract (n=6265)

Full-text articles assessed for eligibility (n=132)

Full-text articles selected for inclusion (n=15)

Full-text articles included (n=18)

Full-text articles found through reference sections (n=3)

Reasons for exclusion:
- Post-treatment assessment before 12 months (n=33, 30%)
- Review, commentary, etc. (n=26, 23%)
- Topic not post-treatment adjustment (n=26, 23%)
- No unsuccessful group or no comparison with control group on the study outcomes (n=12, 10%)
- Data on adjustment outcomes insufficient or unclear (n=9, 8%)
- Post-treatment assessment period not specified (n=4, 3%)
- Duplicate publication (n=2, 2%)
- Publications not accessible (n=1, 1%)

Quantitative Studies (n=9)

Qualitative Studies (n=9)

Figure 1. Decision flow chart for identified studies. * For detailed exclusion strategy see Table S2 of the supplementary material.
### Mental-health

<table>
<thead>
<tr>
<th>Study names</th>
<th>Hedges's g</th>
<th>Standard error</th>
<th>Lower limit</th>
<th>Upper limit</th>
<th>Z-Value</th>
<th>p-Value</th>
<th>-2.00</th>
<th>-1.00</th>
<th>0.00</th>
<th>1.00</th>
<th>2.00</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gameiro et al 2014</td>
<td>-0.093</td>
<td>0.031</td>
<td>-0.155</td>
<td>-0.032</td>
<td>-2.968</td>
<td>0.003</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Johanssen et al 2010</td>
<td>-0.742</td>
<td>0.196</td>
<td>-1.167</td>
<td>-0.377</td>
<td>-3.905</td>
<td>0.000</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Leiblum et al 1998</td>
<td>-0.753</td>
<td>0.287</td>
<td>-1.317</td>
<td>-0.190</td>
<td>-2.620</td>
<td>0.009</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Syydso et al 2011</td>
<td>-0.415</td>
<td>0.144</td>
<td>-0.696</td>
<td>-0.133</td>
<td>-2.884</td>
<td>0.004</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Verhaak et al 2007</td>
<td>-0.596</td>
<td>0.123</td>
<td>-0.838</td>
<td>-0.355</td>
<td>-4.833</td>
<td>0.000</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Weaver et al 1997</td>
<td>-0.589</td>
<td>0.299</td>
<td>-1.175</td>
<td>-0.004</td>
<td>-1.972</td>
<td>0.049</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

### Wellbeing

<table>
<thead>
<tr>
<th>Study name</th>
<th>Hedges's g</th>
<th>Standard error</th>
<th>Lower limit</th>
<th>Upper limit</th>
<th>Z-Value</th>
<th>p-Value</th>
<th>-2.00</th>
<th>-1.00</th>
<th>0.00</th>
<th>1.00</th>
<th>2.00</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hammarberg et al 2001</td>
<td>-0.502</td>
<td>0.189</td>
<td>-0.872</td>
<td>-0.132</td>
<td>-2.658</td>
<td>0.008</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Johansson et al 2010</td>
<td>-0.355</td>
<td>0.183</td>
<td>-0.715</td>
<td>-0.004</td>
<td>-1.939</td>
<td>0.052</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Kuivasaari-Pirinen et</td>
<td>-0.300</td>
<td>0.091</td>
<td>-0.478</td>
<td>-0.122</td>
<td>-3.305</td>
<td>0.001</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Leiblum et al 1998</td>
<td>-1.168</td>
<td>0.299</td>
<td>-1.754</td>
<td>-0.582</td>
<td>-3.907</td>
<td>0.000</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Weaver et al 1997</td>
<td>-0.494</td>
<td>0.222</td>
<td>-0.919</td>
<td>-0.048</td>
<td>-2.177</td>
<td>0.029</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Wischmann et al 2012</td>
<td>-0.080</td>
<td>0.197</td>
<td>-0.466</td>
<td>0.307</td>
<td>-0.404</td>
<td>0.686</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Figure 2. Standardized mean differences (Hedge’s g, Standard Error (SE), 95% Confidence Intervals (CIs), Z value and P value) between failed treatment and control groups for mental-health and wellbeing. Negative Hedge’s values indicate worse adjustment for the failed treatment group than the control group.
Figure 3. A schematic representation of the Three Tasks Model of Adjustment to Unmet Parenthood Goals