

# Best Practice & Research Clinical Obstetrics & Gynaecology

## Economic evaluation of Medically Assisted Reproduction: an educational overview of methods and applications for healthcare professionals

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<b>Abstract:</b>	<p>Economic evaluations of the value-for-money of Medically Assisted Reproduction (MAR) interventions are increasingly important due to growing pressure on healthcare budgets. While such evaluations are commonplace in the published literature, the number/methodological complexity of different evaluations available, and the challenges specific to MAR interventions, can complicate the interpretation of such analyses for fertility treatments. This article aims to serve as an educational resource and provide context on the design/interpretation of economic analyses for MAR interventions. Several areas are relevant for first-line providers and decision makers: scope of analysis, comparator used, perspective/time horizon considered, outcomes used to measure success, and how results from cost-effectiveness studies can be summarized and used in clinical practice. We aim to help clinicians better understand the strengths/weaknesses of economic analyses, to enable the best use of the evidence in practice, so resources available for MAR interventions can provide maximum value to patients and society.</p>

## Highlights

- This educational resource provides context on economic analyses for MAR treatment
- Several factors can obstruct clear interpretation of economic analyses for MAR
- These include the stakeholder/decision-maker, scope of analysis and comparator
- Also the perspective/time horizon and outcome to measure success need consideration
- Greater appreciation of these is essential to improve current methodologies

1 **Economic evaluation of Medically Assisted Reproduction: an educational overview of**  
2 **methods and applications for healthcare professionals**

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40 **Abstract**

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3 41 Economic evaluations of the value-for-money of Medically Assisted Reproduction (MAR)  
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5 42 interventions are increasingly important due to growing pressure on healthcare budgets.  
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7 43 While such evaluations are commonplace in the published literature, the  
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9 44 number/methodological complexity of different evaluations available, and the challenges  
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11 45 specific to MAR interventions, can complicate the interpretation of such analyses for fertility  
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13 46 treatments. This article aims to serve as an educational resource and provide context on the  
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15 47 design/interpretation of economic analyses for MAR interventions. Several areas are relevant  
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17 48 for first-line providers and decision makers: scope of analysis, comparator used,  
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19 49 perspective/time horizon considered, outcomes used to measure success, and how results  
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21 50 from cost-effectiveness studies can be summarized and used in clinical practice. We aim to  
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23 51 help clinicians better understand the strengths/weaknesses of economic analyses, to enable  
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25 52 the best use of the evidence in practice, so resources available for MAR interventions can  
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27 53 provide maximum value to patients and society.  
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32 54 **Keywords:** Economic evaluation, assisted reproduction, cost-effectiveness analysis, fertility  
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## 56 **Introduction**

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3 57 Increasingly, publicly and privately funded fertility clinics are under threat due to constrained  
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5 58 resources, increasing demand, costs of increasing technical improvements and reprioritisation  
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7 59 in healthcare budgets. Under these conditions, there has never been a more important time  
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9 60 for using economic evaluations to optimize outcomes for patients and society with limited  
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11 61 resources available. The current environment underpins the need for clinicians to better  
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13 62 understand how to apply a cost-effectiveness rationale in clinical practice and resist the  
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15 63 temptation to focus only on costs without considering the effectiveness and the overall value-  
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17 64 for-money of Medically Assisted Reproduction (MAR) interventions.

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21 65 Clinicians, as first-line providers and often as decision makers, are in a unique position to  
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23 66 improve MAR delivery. Each choice that they make can have cost and outcome implications.  
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25 67 As such, to help clinicians translate economic evaluations into practice, the aim of this paper  
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27 68 is to provide recommendations for designing economic evaluations and to help clinicians better  
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29 69 understand their application. This paper focuses on several key questions on economic  
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31 70 evaluation that MAR practitioners and policy makers are likely to encounter in daily practice  
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33 71 and about which a better understanding is essential.

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37 72 The key themes of this paper were agreed at a panel meeting of experts, including all co-  
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39 73 authors, who have experience in conducting and/or contributing to economic evaluations of  
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41 74 MAR, which was convened in Frankfurt in November 2019.

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### 47 **Does the analysis produce broad or narrow cost-effectiveness estimates?**

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50 77 An economic evaluation always compares two or more interventions in terms of their costs and  
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52 78 outcomes. The goal is to provide information on how efficient an investment of resources in  
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54 79 one intervention is compared with its alternatives. There can be several possible conclusions:  
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56 80 the costs and outcomes can be equivalent, the intervention can be either more or less costly  
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58 81 than its comparator, and more or less effective in generating a desired outcome. From an  
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82 economic perspective, costlier/less-effective interventions should never be implemented,  
83 whereas less costly/more-effective interventions should always be selected. More difficult  
84 questions, where judgements of cost-effectiveness need to be made, occur when an  
85 intervention is costlier but also more effective, or when it is less costly but also less effective.

86 Depending on the scope of interventions being compared, a different type of economic  
87 evaluation will have to be used. The first and broadest scope considers the allocation of  
88 resources across many healthcare settings and beyond. These analyses consider how  
89 increased spending on one particular area of healthcare might compare in terms of value for  
90 money with the investments made in health services in other areas or even in other sectors of  
91 the economy beyond healthcare. Here, a health-economic approach of societal value is about  
92 understanding those services and outcomes that are important to citizens and allocating  
93 budget to those programmes deemed most important to society, with the aim of maximising  
94 societal welfare [1, 2]. Cost-utility analyses (CUAs) and societal cost-benefit analyses (CBAs)  
95 assess the benefits of a programme in the broad and generic terms of quality-adjusted life  
96 years (QALYs) or monetary units, respectively, so that they allow broad comparisons (e.g.  
97 across disease categories [see **Table 1**]). These analyses inform policy makers about  
98 allocative efficiency: the 'optimal' resource allocation within the constraints of the budget. It is  
99 this resource allocation that needs to be addressed during discussions regarding whether MAR  
100 should be publicly funded at all and, if so, how big its budget should be relative to other  
101 healthcare domains (e.g., oncology/mental health) or other ways of spending public money  
102 (e.g., infrastructure). This type of economic study is rarely executed in MAR.

103 The second scope, which is most commonly reported in the published literature of MAR and  
104 in clinical practice, is narrower and only compares interventions by their ability to achieve a  
105 well-defined set of outcomes. It investigates, within the narrow boundaries of a condition-  
106 specific outcome measure (in this case, a live birth), which intervention delivers this outcome  
107 at the lowest cost or what additional costs are needed to generate one additional unit of that  
108 outcome (i.e., live birth rate). This type of information is provided by a cost-effectiveness



109 analysis (CEA). By the nature of its specific outcome metric, it allows a much narrower scope  
1 of comparisons and, in its traditional use, is most suited to informing the maximisation of a  
2 110 of comparisons and, in its traditional use, is most suited to informing the maximisation of a  
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4 111 desired unit of outcome in the context of an earmarked budget for MAR.  
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7 112 However, it is common to see the results of cost-effectiveness studies of MAR technology  
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9 113 reported as an incremental analysis, whereby an 'incremental cost-effectiveness ratio (ICER)'  
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11 114 is calculated by measuring the incremental cost of a new therapeutic approach and dividing it  
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14 115 by the incremental effects (e.g., additional live births). In situations where this infers an  
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16 116 increase in MAR spending for an increase in live births, this takes us into the realms of  
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18 117 allocative efficiency: depending on the cost of the new therapeutic approach, its adoption may  
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20 118 require further allocation of scarce resources [3].  
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23 119 In short, in the context of MAR, CUAs and societal CBAs address the degree to which  
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25 120 healthcare systems should treat the condition of infertility or increase spending on MAR  
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27 121 treatments compared with other healthcare domains. CEAs help to identify those MAR  
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29 122 programs that optimize live birth rates [4], although an incremental analysis can also be  
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31 123 informative in terms of allocative efficiency when it is understood how valuable one birth is to  
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33 124 society (see also the section **How are economic evaluation results summarized?**).  
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### 40 126 **What is the comparator intervention used in the analysis?**

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43 127 One of the first considerations in conducting or evaluating an economic evaluation is  
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45 128 establishing the comparator: the intervention or standard of care to be replaced following the  
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47 129 introduction of a new medical technology. Alternatively, a new MAR intervention can be  
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50 130 compared to many products when multiple treatment options exist. By choosing the wrong  
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52 131 comparator, or omitting a relevant comparator, an intervention can easily be made to look  
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54 132 more or less cost-effective than it really is. When assessing the applicability of cost-  
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56 133 effectiveness studies to clinical practice, one must scrutinize whether the comparator is  
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58 134 appropriate for their patient population and local setting (see also the section **How can**

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135 **evidence be used in practice?**). For example, when comparing two stimulation protocols, it  
136 is necessary to assess whether the dosing in each arm reflects current standard of care  
137 according to international professional guidelines/recommendations and existing ART  
138 treatment practices in a real-world context. If doses or practices vary dramatically versus a  
139 benchmark/recommended practice, this may substantially limit the interpretation of the  
140 findings when implemented under different settings.

141 When there are many possible interventions to compare, a full incremental analysis should be  
142 performed, whereby all alternatives are compared incrementally in order of increasing costs  
143 and effects. The rules of 'dominance' and 'extended dominance', whereby interventions with  
144 higher costs and lower benefits than one alternative or a combination of alternatives are  
145 eliminated, indicate which of the interventions are the relevant ones to compare using ICERs  
146 [5]. For example, when comparing five alternative *in vitro* fertilisation (IVF) embryo transfer  
147 strategies, van Heesch et al. found that four and five cycles of elective single-embryo transfer  
148 were extendedly dominated by a combination of three cycles of elective single-embryo transfer  
149 and then three cycles according to standard-embryo transfer policy; therefore, the two  
150 dominated strategies were not considered in the incremental analysis [6].

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### 152 **What is the perspective adopted in the analysis?**

153 Depending on the target audience, different costs and outcomes are relevant. The narrowest  
154 perspective would be for the patients themselves, followed by the clinic, private health  
155 insurance provider, public healthcare payer and societal perspectives. Any analysis will  
156 consequently present a comparison of costs and outcomes from the chosen perspective (i.e.,  
157 relative to who pays the costs and who receives the outcome) and excluding costs and effects  
158 that fall on other parties. An intervention can be cost-effective from one perspective but not  
159 from another perspective (e.g., because the former party only carries part of the costs  
160 incurred). For example, influenza or depression cost most in terms of their impact on work  
161 absenteeism (productivity costs) and cost less in terms of actual medical treatment [7, 8].

162 Therefore, prevention of these diseases is much more cost-effective from a societal than from  
1 a healthcare payer perspective, because most of the associated cost savings are societal ones  
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3 that fall outside the budgets of healthcare payers. The cost of multiple pregnancies as a result  
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5 of MAR are paid for by the public health system (society/taxpayers) in most countries, and are  
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7 often not paid directly by the patient [9]. As such, individuals paying out of pocket may prefer  
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9 to transfer two or more embryos, in an effort to increase the probability of live birth per embryo  
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11 transfer, without fully realizing the medical maternal and perinatal risks associated with multiple  
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13 pregnancy and, as such, impose higher costs related to multiple pregnancies/births.  
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170 For many, the societal perspective is considered the most relevant, as this broad perspective  
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19 captures all benefits and costs for society [10]. Alternatively, for others, narrower perspectives  
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21 can be applied to evaluate costs that fall on individuals or organisations being asked to pay for  
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23 a new product or technology. For example, many national health technology assessment  
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25 (HTA) agencies apply a narrow health service perspective that only considers those costs that  
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27 are paid by the public health system, excluding costs like productivity losses or out-of-pocket  
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29 payments for patients. There are limitations to applying a narrow perspective that can lead to  
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31 inefficient allocation of resources and inefficient treatment choices, as this fails to consider the  
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33 implications of certain choices for involved third parties that were not considered in the  
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35 analysis.  
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180 Costs associated with MAR are often shared among public systems, private insurers and  
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42 patients. Whilst all fractions can be important cost components, the perspective defines those  
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44 costs to be included in the analysis. In the strictest sense, once the perspective has been  
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46 determined, only those costs relevant to the perspective applied in the analysis should be  
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48 considered. Similarly, if costs are shared for a particular item of service delivery, for example,  
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50 prescription medicines and patient co-payments, these contributions should be deducted from  
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52 the overall costs of the medicine to ensure only those aligned with the perspective are  
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54 considered. For MAR this can again be complicated, as there is usually a mix of public and  
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56 private funding for different MAR treatment components. Unlike other areas of healthcare,  
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189 there are varied public reimbursement programmes for MAR, but a large proportion of cycles  
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2 190 are delivered in private clinics that incur patient fees [11]. The reliance on patients paying out-  
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4 191 of-pocket for care can cause many issues that disrupt the efficient and timely delivery of care.  
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6 192 Financial barriers can often limit access to treatment for couples, influence treatment practices  
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8 193 and can impact pregnancy outcomes [12-14]. The variance in who funds the treatment  
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10 194 influences how we must think about the cost-effectiveness of treatments. In some instances,  
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12 195 couples may pay for MAR treatment, with the national health system funding the consequences  
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14 196 of adverse outcomes, such as multiple pregnancies, all of which should be considered in  
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16 197 decision-making.  
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### 23 199 **What is the time horizon adopted in the analysis?**

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26 200 Costs and effects can be measured for different time horizons. Usually, in the context of MAR,  
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28 201 a short time perspective is adopted, limiting the analysis to the period of treatment until delivery  
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30 202 of a live birth, often less than one year of follow-up. In some cases, limiting the analysis to the  
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32 203 point of delivery can be short-sighted, as many costs can occur further downstream.  
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34 204 Furthermore, choosing an appropriate analytic time horizon is a particularly important construct  
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36 205 for MAR because a newborn generates a lifetime of benefits (and costs) for parents and  
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38 206 society. Similarly, childlessness, and the physical or mental consequences of childlessness  
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40 207 that can occur, persist over the remaining lifetime. Furthermore, this is also true for multiple  
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42 208 pregnancies, which can give rise to more complications at delivery and perhaps persistent  
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44 209 complications that require ongoing medical care. When a study compares options that confer  
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46 210 different risks of multiple birth or other complications per live birth that affect the health of the  
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48 211 offspring, the chosen time horizon can be influential.  
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53 212 In practice, many published cost-effectiveness studies fail to report the timeline of the analysis,  
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55 213 which can limit their usefulness. A previous review of cost-effectiveness studies observed that  
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57 214 58% of included studies did not clearly define a time horizon, and only 13% considered a time  
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59 215 horizon of more than one year [15]. The general recommendation is that the time horizon  
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216 should be long enough to capture any relevant differences in costs and outcomes between the  
217 different approaches being compared, and this should also consider the use of subsequent  
218 treatments in the context of the clinical care pathways that couples have access to throughout  
219 their MAR journey [16]. However, the question over when to stop counting costs associated  
220 with the outcome of MAR is more ambiguous, and the reality is that most cost-effectiveness  
221 studies based on randomized studies only consider costs to the point of a live birth.

222 Where to draw the line and from which point onwards costs and benefits should not be counted  
223 is arbitrary as, in principle, newborns could be followed for a lifetime. The practice of stopping  
224 cost counting from the delivery onwards may be justified on the grounds that the live birth is  
225 the primary desired outcome of MAR treatment, and future costs and health outcomes for  
226 infants may be considered irrelevant in this context. However, successful MAR leads to future  
227 perinatal and delivery costs, and including these costs in association with a successful  
228 outcome treats live births as an adverse event, which could negatively influence the cost per  
229 live birth comparison. In this context it might be of interest to exclude costs associated with live  
230 births, as interventions increasing live birth rates will be associated with higher costs but may  
231 still be cost-effective if the cost per live birth is considered. Similar arguments may be made in  
232 relation to pregnancy costs, in that the provision of antenatal care is considered a cost-effective  
233 use of healthcare resources for pregnant women, regardless of how the pregnancy was  
234 conceived. However, this becomes less justifiable when comparing MAR treatments that  
235 confer different risks of complications that affect the cost of pregnancy and/or costs and  
236 outcomes for offspring. In the case of multiple births, the excess costs could be considered an  
237 adverse event worth including in the analysis, and a much longer time horizon may be required  
238 to fully capture the impact of this; potentially, the lifetime of the infant. Similarly, if one MAR  
239 treatment led to increases in early pregnancy loss, pre-term births or congenital abnormalities  
240 compared with an alternative MAR treatment, it is important to capture the excess costs  
241 attributable to these. Safety as well as success matters.

243 **Which outcomes are measured to assess effectiveness?**

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3 244 For narrow cost-effectiveness analyses, first live birth (following a fresh or frozen embryo  
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5 245 transfer following ART treatment) or cumulative live birth (all live births following successive  
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7 246 ART treatments) are logical outcome measures, more than pregnancies or deliveries.  
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9 247 Furthermore, (cumulative) live births also integrate the occurrence of early pregnancy loss  
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11 248 related to ectopic pregnancies and miscarriages. For broader analyses (CUAs and CBAs),  
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13 249 however, it becomes more difficult to establish appropriate outcome measures.  
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16 250 What are the likely benefits achieved from treating infertility? For couples, the medical  
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18 251 diagnosis is labelled as infertility, which is 'a disease of the male or female reproductive  
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21 252 system, defined by the failure to achieve pregnancy after 12 months or more of regular  
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23 253 unprotected sexual intercourse' [17]. While infertility is often caused by conditions presenting  
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25 254 with specific symptoms, such as pelvic/menstrual pain (endometriosis, uterine fibroids, chronic  
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27 255 pelvic inflammatory disease, postoperative pelvic adhesions, etc.) or menstrual cycle  
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30 256 abnormalities (anovulation caused by polycystic ovarian syndrome, hyperprolactinemia, etc.),  
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32 257 it can also be largely asymptomatic. The main goal of infertility treatment is to produce a child,  
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34 258 thereby satisfying the desire to have children, while obviously also managing, to whatever  
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36 259 extent possible, the causes of infertility. To inform the allocation of constrained healthcare  
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39 260 budgets, the reference case for economic evaluation tends to focus attention on the individual  
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41 261 medical condition and health benefits acquired by individuals undergoing treatment, although  
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43 262 more attention is paid to spill-over effects in more recent analyses. This approach treats the  
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45 263 resulting live birth as an 'externality'; in other words, it is external to the individuals experiencing  
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47 264 infertility.  
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50 265 Such thinking underpins the cost-effectiveness modelling conducted to inform recent NICE  
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52 266 guidelines on the provision of ART and treatment for endometriosis causing infertility [18, 19],  
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55 267 where the value of treatment was captured through its impact on satisfying the desire of infertile  
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57 268 couples for a child, using QALYs as the unit of outcome. Although it was recognised, in the  
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59 269 interpretation of results, that decision makers may value live births beyond their impact on the  
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270 QALYs of couples seeking treatment. However, the QALYs of the future child are not  
271 considered in the evaluation. Whether or not to count QALYs of future children is a complex  
272 ethical matter in itself, with both exclusion and inclusion leading to counterintuitive results.  
273 QALYs were developed as a generic measure of health benefit to capture improvements in  
274 health among patients. As such, it has been argued that they are not appropriate for capturing  
275 the value of live births achieved through MAR treatment [20]. Nevertheless, focussing only on  
276 the QALYs of couples seeking treatment ignores the fact that the child represents future value  
277 for society and the government, and potentially undervalues MAR treatment. Whilst it is  
278 possible to enumerate the QALYs accruing to children born as result of MAR, it would, in this  
279 situation, be incorrect to apply the same rules of interpretation that are applied to incremental  
280 cost per QALY estimates for other health conditions. How to fully capture the value of live births  
281 to society in the context of economic evaluation remains an area requiring further research and  
282 debate [21]. Monetary valuation in the context of return on investment or CBA offer an  
283 alternative approach in this respect (discussed further below).

284

### 285 **How are economic evaluation results summarized?**

286 The two main cost-effectiveness measures reported in the literature are the average cost-  
287 effectiveness ratio (ACER) and the ICER, both often expressed as a cost per live birth. In  
288 practice, the ACER is a simpler construct to understand, as it is reported as a ratio of average  
289 costs over the average benefits for each alternative intervention considered in the analysis.  
290 The ACER therefore reflects the sum of all costs incurred by an intervention, divided by the  
291 likelihood of achieving a live birth (Equation 1), which can then be compared across all of the  
292 competing MAR interventions to select the most technically efficient option. However, when  
293 comparing interventions with each other, it is customary to report the ICER per live birth (i.e.,  
294 the increase in costs to achieve one additional live birth with one intervention compared with  
295 another [Equation 2]). To interpret this outcome, the ICER per live birth measure does pose a  
296 problem, as no willingness-to-pay threshold has been established for how much society is

297 willing to pay for an additional child. Many cost-effectiveness studies report incremental results,  
 298 but there are no rules that govern whether \$30,000 or \$100,000 per additional live birth is  
 299 considered acceptable. Incremental results would be more applicable if there were better  
 300 understanding of societies' willingness to pay for an additional child in the context of MAR  
 301 provision, where it is often competing for scarce resources with other healthcare interventions.  
 302 This is an area that would benefit from further research to assess the preference of society  
 303 and the willingness to trade-off health benefits for other groups of patients against live births  
 304 delivered through the provision of MAR. In this regard, when making resource allocation  
 305 decisions in relation to MAR treatments, cost-effectiveness analyses might currently be more  
 306 suited to evaluating the average cost per live birth for each product individually, where the one  
 307 with the lowest cost per live birth is likely to be the more technically efficient option.

Equation 1

$$\text{Average cost per live birth of program A} = \frac{\Sigma \text{costs of program A}}{\Sigma \text{live births A}}$$

Equation 2

$$\text{Incremental cost per live birth of program A vs. B} = \frac{(\Sigma \text{costs A} - \Sigma \text{costs B})}{(\Sigma \text{Live Births A} - \Sigma \text{Live Births B})}$$

313 To inform allocative efficiency on the broadest level, CBA is used, whereby all outcomes are  
 314 reported in monetary units (see **Table 1**), in a return-on-investment metric or net-benefit  
 315 estimate. This involves converting live births and other MAR outcomes into monetary values.  
 316 In the strictest sense, MAR is used to treat infertility, which is a medical disease experienced  
 317 by couples, the consequence of which is fewer children are born. As already discussed,  
 318 because the benefits attributed to unborn children are not considered part of the medical  
 319 problem of infertility, successful treatment (i.e., live births) represent an externality to the initial  
 320 medical problem of infertility. However, the resulting child does have value when considered



1  
2 322 from the perspective of parents, families, society and governments. Examples of this were  
3  
4 323 observed in Korea [22] and Japan [23], where the respective governments increased funding  
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6 324 for MAR treatment, not because of the burden on couples caused by infertility, but as a policy  
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8 325 measure to aid falling birth rates [24]. By valuing the externality of children and future economic  
9  
10 326 contributions of the child, these governments elected to fund MAR for infertile couples as a  
11 327 way to secure future economic benefits provided by their children.  
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13  
14 328 The government perspective cost-benefit framework has been applied in several previous  
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16 329 studies, to inform allocative efficiency by capturing the lifetime net tax contributions attributed  
17  
18 330 to MAR-conceived children [25, 26]. The aim of these previous studies was to demonstrate the  
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20 331 future value of MAR-conceived children in relation to treatment costs. Because funding for  
21  
22 332 MAR competes with all other healthcare programmes for funding within tax-financed public  
23  
24 333 systems, to inform on allocative efficiency suggests that we know the relative value of these  
25  
26 334 other programmes, although this is seldom the case. Considering that MAR is the only  
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28 335 intervention that creates human life, when the externalities are valued few medical  
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30 336 technologies are likely to show greater economic value in the future, especially in societies  
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32 337 where the birth rate is below replacement level. The analyses demonstrate that, from the  
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34 338 perspective of government, MAR investment costs yield a significant future fiscal gain over the  
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36 339 lifetimes of these children.  
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41 340 Stated preference techniques offer an alternative approach to placing monetary value on the  
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43 341 provision of MAR and its associated outcomes, by assessing what individuals, patients or  
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45 342 members of the general population are hypothetically willing to pay for it. For example, using  
46  
47 343 such an approach Botha et al. estimated that individuals in a representative sample of the  
48  
49 344 Australian population were willing to pay an additional \$27.43 in annual tax contributions for a  
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51 345 preferred configuration of fertility treatment provision [27]. Furthermore, Spiegel, et al.  
52  
53 346 estimated that the willingness to pay for an IVF cycle in Israel among patients (\$5,482) and the  
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55 347 general public (\$4,398) was more than the actual average cost of IVF treatment (\$3,257),  
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348 highlighting the perceived net benefits of IVF for both patients and society [28]. Such values  
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2 349 provide a means for allowing direct comparison with the costs of providing fertility in a CBA.  
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7 351 **How can evidence be used in practice?**  
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10 352 The extent to which economic evaluation studies inform choices for privately paying patients  
11  
12 353 is not well understood. Patients as customers, who are already paying out-of-pocket for  
13  
14 354 treatment, are clearly governed by different principles, where securing a live birth and time to  
15  
16 355 pregnancy have heightened importance. As privately paying patients may not fully understand  
17  
18 356 the potential risks, in particular for multiple pregnancies, there is potential for adverse treatment  
19  
20 357 selection. In these cases, privately paying couples may expose themselves to the risks of  
21  
22 358 multiple pregnancies, without the consequences of having to pay for many of the additional  
23  
24 359 health service costs associated with multiple deliveries and complications.  
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26

27  
28 360 The results from economic evaluations can be applied at several levels within a healthcare  
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30 361 system to improve efficiency. For instance, national level bodies are often responsible for  
31  
32 362 evaluating the cost-effectiveness of products as part of the national reimbursement process  
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34 363 and are sometimes involved in negotiating prices [29]. Nationally, this mostly occurs when new  
35  
36 364 products enter the market or when developing clinical guidelines, which is performed by  
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38 365 agencies such as NICE in the UK. Additionally, local hospital formulary committees also  
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40 366 consider economic data when making local funding decisions [30]. Many important attributes  
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42 367 are not captured in cost-effectiveness ratios that may need to be considered. For instance, live  
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44 368 births and even QALYs, do not address equity, dignity, autonomy, and patient choice – all  
45  
46 369 factors that have value to individuals and society, but are not captured within CEAs.  
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50  
51 370 The cost-effective delivery of MAR, the area where economic evaluation studies are likely to  
52  
53 371 have the greatest impact, is at the level of the clinic and hospital, where clinical groups are  
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55 372 often charged with procuring products and making decisions regarding products and services  
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57 373 to improve treatment outcomes – often with fixed budgets. There are a few important points  
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374 that clinicians and other decision-makers can take into consideration when adopting economic  
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2 375 evaluation research into their practice. Firstly, the most important consideration is the clinical  
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4 376 evidence on which the model is based. Therefore, evaluating whether the evidence meets  
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6 377 established standards and whether the evidence is based on randomized studies or real-world  
7  
8 378 practices can influence the results. Secondly, it is necessary to consider whether the treatment  
9  
10 379 practices (e.g., diagnostics, clinic visits, oocyte extraction) and associated costs, as reported  
11  
12 380 in the evidence, reflect those applied within the practice. Thirdly, many economic evaluation  
13  
14 381 studies simulate different treatment practices based on sequences of fresh and frozen ART  
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16 382 cycles. Often, these are based on data from RCTs, and only rarely include a long-term, real-  
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18 383 world perspective, which takes into account that many patients are treated during multiple fresh  
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20 384 (up to three and sometimes more) ovarian stimulation cycles for ART and related fresh and  
21  
22 385 frozen ART cycles. Typically, these models mostly reflect a cohort of people treated in a fairly  
23  
24 386 homogenous manner. Therefore, it is important to consider whether the cohort reported in  
25  
26 387 economic evaluation studies reflects those of the local practice, and additionally how one  
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28 388 integrates frozen cycles into treatment practice. The more the practice may vary from the  
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30 389 standard treatment approach reported in the study, the more likely it is that these study findings  
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32 390 have limited applicability.  
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38 391 Another important consideration when looking to apply economic evaluation results is the  
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40 392 transferability of findings across geographic boundaries. By design, cost-effectiveness models  
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42 393 use country-specific cost data. As there is considerable variation in treatment practices and  
43  
44 394 costs, the results from economic evaluation studies are not always transferable to different  
45  
46 395 markets [31]. This limitation suggests the need to localize economic models to each market to  
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48 396 address cost-effectiveness questions. When applying studies in clinical practice, it is important  
49  
50 397 to consider whether the costs are relevant to the practice. The same could be said about  
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52 398 clinical data used in models and whether it is applicable in other settings. Current practice is  
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54 399 often to use the results from international randomized studies for the basis of outcomes data;  
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56 400 however, RCTs are often limited by selected study populations, usually patients with a good  
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1 401 prognosis and excluding patients at risk for poor or exaggerated response to ovarian  
2 402 stimulation, representing only a minority of the wider real-world population treated with ART  
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4 403 [32]. Local real-world data sources may, in this case, provide more accurate estimates of  
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6 404 treatment effects and resource consumption  
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9 405

## 11 406 **Summary**

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14 407 In this educational article, we have bundled a set of key questions that need to be answered,  
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16 408 in order to better understand the value of an economic evaluation in the context of MAR.  
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18 409 These questions, their answers, and also an appropriate understanding of the remaining  
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20 410 issues, can help clinicians understand the strengths and weaknesses of studies and enable  
21  
22 411 them to make best use of the evidence in practice, so that the resources available for MAR  
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24 412 treatment can be used to create maximum value to patients and society. It is important to  
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26 413 recognize that achieving efficiency in budget allocation for MAR treatment is not the same  
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28 414 thing as providing fewer services or lowering costs. In many cases, doing more or spending  
29  
30 415 more is necessary to achieve increased live birth rates. The question to consider is whether  
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32 416 increased expenditure is justifiable in relation to the additional outcomes achieved (i.e.,  
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34 417 whether extra investment is 'worth it', either to patients, clinics, insurance companies,  
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36 418 healthcare systems or societies as a whole). Economic evaluations aim to assist decision  
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38 419 makers in making these judgements. However, they can often appear callous, as they focus  
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40 420 on a single measure of effect and the costs of achieving such measures, often evaluated  
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42 421 over a short time period. Moreover, in the context of MAR there are many methodological  
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44 422 issues that obstruct a clear interpretation of cost-effectiveness analyses.  
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19 432 manuscript.

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23  
24  
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44  
45  
46 443 All authors contributed to the conception and design of the analysis, as well as interpretation  
47  
48 444 of data and critical review of this manuscript. All authors approved the manuscript for  
49  
50 445 submission to the journal.  
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53 446 **Data availability**

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56 447 Any requests for data by qualified scientific and medical researchers for legitimate research  
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58 448 purposes will be subject to Merck KGaA's Data Sharing Policy. All requests should be  
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449 submitted in writing to Merck KGaA's data sharing portal

1  
2 450 <https://www.merckgroup.com/en/research/our-approach-to-research-and->

3  
4 451 [development/healthcare/clinical-trials/commitment-responsible-data-sharing.html](https://www.merckgroup.com/en/research/our-approach-to-research-and-development/healthcare/clinical-trials/commitment-responsible-data-sharing.html). When

5  
6 452 Merck KGaA has a co-research, co-development, or co-marketing or co-promotion

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8 453 agreement, or when the product has been out-licensed, the responsibility for disclosure might

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11 454 be dependent on the agreement between parties. Under these circumstances, Merck KGaA

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13 455 will endeavour to gain agreement to share data in response to requests.

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55  
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57  
58  
59  
60  
61  
62  
63  
64  
65

457 **References**

- 1  
2 458 1. McIntosh E, Luengo-Fernandez R. Economic evaluation. Part 2: frameworks for combining  
3  
4 459 costs and benefits in health care. *J Fam Plann Reprod Health Care* 2006;32(3):176-80.  
5  
6
- 7 460 2. Palmer S, Torgerson DJ. Definitions of efficiency. *BMJ* 1999;318(7191):1136.  
8
- 9 461 3. Donaldson C, Currie G, Mitton C. Cost effectiveness analysis in health care: contraindications.  
10  
11 462 *BMJ* 2002;352:891-4.  
12  
13
- 14 463 4. Drummond M, Sculpher M, Claxton K, Stoddart G, Torrance G. *Methods for the Economic*  
15  
16 464 *Evaluation of Health Care Programmes*. 4th ed. UK: Oxford University Press; 2015.  
17
- 18 465 5. Johannesson M, Weinstein M. On the decision rules of cost-effectiveness analysis. *J Health*  
19  
20 466 *Econ* 1993;12:459-67.  
21  
22
- 23 467 6. van Heesch MM, van Asselt AD, Evers JL, van der Hoeven MA, Dumoulin JC, van Beijsterveldt  
24  
25 468 CE, et al. Cost-effectiveness of embryo transfer strategies: a decision analytic model using long-term  
26  
27 469 costs and consequences of singletons and multiples born as a consequence of IVF. *Hum Reprod*  
28  
29 470 2016;31(11):2527-40.  
30  
31  
32
- 33 471 7. Nichol K. Cost-benefit analysis of a strategy to vaccinate healthy working adults against  
34  
35 472 influenza. *Arch Intern Med* 2001;161:749-59.  
36
- 37 473 8. Luyten J, Knapp M. Economic evaluation of mental health promotion and mental illness  
38  
39 474 prevention. In: Bährer-Kohler S, Carod-Artal FJ, editors. *Global Mental Health: Prevention and*  
40  
41 475 *Promotion*. New York: Springer International Publishing; 2017.  
42  
43
- 44 476 9. Crawford S, Boulet SL, Mneimneh AS, Perkins KM, Jamieson DJ, Zhang Y, et al. Costs of  
45  
46 477 achieving live birth from assisted reproductive technology: a comparison of sequential single and  
47  
48 478 double embryo transfer approaches. *Fertil Steril* 2016;105(2):444-50\*.  
49  
50
- 51 479 10. Jönsson B. Ten arguments for a societal perspective in the economic evaluation of medical  
52  
53 480 innovations. *Eur J Health Econ* 2009;10(4):357-9.  
54  
55
- 56 481 11. Berg Brigham K, Cadier B, Chevreul K. The diversity of regulation and public financing of IVF in  
57  
58 482 Europe and its impact on utilization. *Hum Reprod* 2013;28(3):666-75.  
59  
60  
61  
62  
63  
64  
65

- 483 12. Verbeke E, Luyten J, D'Hooghe T. The Economics of IVF: Evaluating the Necessity and Value of  
1  
2 484 Public Funding. Patient-Centered Assisted Reproduction: How to Integrate Exceptional Care with  
3  
4 485 Cutting-Edge Technology 2020:106\*.  
5  
6  
7 486 13. Boulet SL, Crawford S, Zhang Y, Sunderam S, Cohen B, Bernson D, et al. Embryo transfer  
8  
9 487 practices and perinatal outcomes by insurance mandate status. Fertil Steril 2015;104(2):403-9. e1.  
10  
11 488 14. Reynolds MA, Schieve LA, Jeng G, Peterson HB. Does insurance coverage decrease the risk for  
12  
13 489 multiple births associated with assisted reproductive technology? Fertil Steril 2003;80(1):16-23.  
14  
15  
16 490 15. Moolenaar LM, Cissen M, de Bruin JP, Hompes PG, Repping S, van der Veen F, et al. Cost-  
17  
18 491 effectiveness of assisted conception for male subfertility. Reprod Biomed Online 2015;30(6):659-66.  
19  
20  
21 492 16. Neumann PJ, Johannesson M. The willingness to pay for *in vitro* fertilization: a pilot study using  
22  
23 493 contingent valuation. Med Care 1994;32(7):686-99.  
24  
25  
26 494 17. WHO. International Classification of Diseases, 11th Revision (ICD-11) Geneva, Switzerland:  
27  
28 495 World Health Organization; 2018 [Accessed September]. Available from: [https://www.who.int/news-](https://www.who.int/news-room/fact-sheets/detail/infertility)  
29  
30 496 [room/fact-sheets/detail/infertility](https://www.who.int/news-room/fact-sheets/detail/infertility).  
31  
32  
33 497 18. NICE. Fertility problems: assessment and treatment: Clinical guideline [CG156] London, UK:  
34  
35 498 National Institute for Health and Care Excellence 2013 [Accessed 16 July]. Available from:  
36  
37 499 [https://www.nice.org.uk/guidance/cg156/resources/fertility-problems-assessment-and-treatment-](https://www.nice.org.uk/guidance/cg156/resources/fertility-problems-assessment-and-treatment-pdf-35109634660549*)  
38  
39 500 [pdf-35109634660549\\*](https://www.nice.org.uk/guidance/cg156/resources/fertility-problems-assessment-and-treatment-pdf-35109634660549*).  
40  
41  
42 501 19. NICE. Endometriosis: diagnosis and management: NICE guideline [NG73] London, UK: National  
43  
44 502 Institute for Health and Care Excellence; 2017 [Accessed 16 July]. Available from:  
45  
46 503 <https://www.nice.org.uk/guidance/NG73>.  
47  
48  
49 504 20. Devlin N, Parkin D. Funding fertility: issues in the allocation and distribution of resources to  
50  
51 505 assisted reproduction technologies. Hum Fertil (Camb) 2003;6:S2-6.  
52  
53  
54 506 21. Luyten J, Verbeke E, Schokkaert E. To be or not to be: Future lives in economic evaluation.  
55  
56 507 Health Economics 2021.  
57  
58  
59  
60  
61  
62  
63  
64  
65



- 508 22. Kim S. Reproductive technologies as population control: how pronatalist policies harm  
1  
2 509 reproductive health in South Korea. *Sex Reprod Health Matters* 2019;27:6-12.  
3
- 4 510 23. McCurry J. Japan to help cover IVF costs in attempt to avert demographic crisis. *The Guardian*.  
5  
6 511 2020.  
7
- 8  
9 512 24. Connolly MP, Ledger W, Postma MJ. Economics of assisted reproduction: access to fertility  
10  
11 513 treatments and valuing live births in economic terms. *Hum Fertil (Camb)* 2010;13(1):13-8.  
12  
13
- 14 514 25. Connolly M, Gallo F, Hoorens S, Ledger W. Assessing long-run economic benefits attributed to  
15  
16 515 an IVF-conceived singleton based on projected lifetime net tax contributions in the UK. *Hum Reprod*  
17  
18 516 2009;24(3):626-32.  
19  
20
- 21 517 26. Connolly MP, Postma MJ, Crespi S, Andersen AN, Ziebe S. The long-term fiscal impact of  
22  
23 518 funding cuts to Danish public fertility clinics. *Reprod Biomed Online* 2011;23(7):830-7.  
24  
25
- 26 519 27. Botha W, Donnelly N, Shanahan M, Norman RJ, Chambers GM. Societal preferences for  
27  
28 520 fertility treatment in Australia: a stated preference discrete choice experiment. *J Med Econ*  
29  
30 521 2019;22:95-107.  
31  
32
- 33 522 28. Spiegel U, Gonen LD, Templeman J. Economic implications of *in vitro* fertilization using  
34  
35 523 willingness to pay. *J Public Health* 2013;21:535-57.  
36  
37
- 38 524 29. EUNETHTA. Methods for health economic evaluations - A guideline based on current practices  
39  
40 525 in Europe. 2015.  
41
- 42 526 30. Seigfried RJ, Corbo T, Saltzberg MT, Reitz J, Bennett DA. Deciding which drugs get onto the  
43  
44 527 formulary: a value-based approach. *Value Health* 2013;16(5):901-6.  
45  
46
- 47 528 31. O'Brien B. A tale of two (or more) cities: geographic transferability of pharmacoeconomic data.  
48  
49 529 *Am J Manag Care* 1997;3:S33-9.  
50  
51
- 52 530 32. Hershkop E, Segal L, Fainaru O, Kol S. 'Model' versus 'everyday' patients: can randomized  
53  
54 531 controlled trial data really be applied to the clinic? *Reprod Biomed Online* 2017;34:274-9\*.  
55  
56  
57  
58  
59  
60  
61  
62  
63  
64  
65

- 532 33. van Loendersloot LL, Moolenaar LM, van Wely M, Repping S, Bossuyt PM, Hompes PGA, et al.  
1  
2 533 Cost-effectiveness of single versus double embryo transfer in IVF in relation to female age. Eur J Obstet  
3  
4 534 Gynecol Reprod Biol 2017;214:25-30\*.  
5  
6  
7 535 34. Mennini FS, Marcellusi A, Viti R, Bini C, Carosso A, Revelli A, et al. Probabilistic cost-  
8  
9 536 effectiveness analysis of controlled ovarian stimulation with recombinant FSH plus recombinant LH vs.  
10  
11 537 human menopausal gonadotropin for women undergoing IVF. Reprod Biol Endocrinol 2018;16(1):68.  
12  
13  
14 538 35. Darvishi A, Goudarzi R, Zadeh VH, Barouni M. Cost-benefit Analysis of IUI and IVF based on  
15  
16 539 willingness to pay approach; case study: Iran. PLoS One 2020;15(7):e0231584.  
17  
18  
19 540 36. Scotland G, McLernon D, Kurinczuk J, McNamee P, Harrild K, Lyall H, et al. Minimising twins in  
20  
21 541 *in vitro* fertilisation: a modelling study assessing the costs, consequences and cost–utility of elective  
22  
23 542 single versus double embryo transfer over a 20-year time horizon. BJOG 2011(118):1073-83.  
24  
25

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34  
35  
36  
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38  
39  
40  
41  
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546 **Table 1.** Types of economic evaluation

Type of evaluation	Cost measure	Outcome measure		MAR Example
<b>Cost-effectiveness analysis (CEA)</b>		<b>Physical, natural units</b>	<ul style="list-style-type: none"> <li>• Pregnancy</li> <li>• Delivery</li> <li>• Live birth</li> <li>• Cumulative live birth</li> </ul>	<ul style="list-style-type: none"> <li>• Sequential single embryo transfers, when clinically appropriate, can reduce total ART treatment and pregnancy/infant-associated medical costs by reducing multiple births without lowering live birth rates [9].</li> <li>• Single embryo transfer followed by an additional frozen-thawed single embryo transfer (if available) was dominant: less costly and more effective than double embryo transfer in women under 32 years [33]</li> </ul>
<b>Cost-utility analysis (CUA)</b>	Volume of physical resource times unit cost	'Utilities' that take into account public preferences for different health states	<ul style="list-style-type: none"> <li>• Quality-adjusted life year (QALY)</li> <li>• Disability-adjusted life year (DALY)</li> <li>• Healthy years equivalent (HYE)</li> </ul>	<ul style="list-style-type: none"> <li>• In the short term (1 year), single embryo transfer is more cost-effective than double embryo transfers; however, in the intermediate (5 years) and long terms (18 years), double embryo transfer becomes the most cost-effective strategy, with a ceiling ratio of €20,000 per QALY [6].</li> <li>• The 2:1 combination of recombinant human follicle stimulating hormone and recombinant human luteinising hormone was a cost-effective option for ovarian stimulation (ICER per QALY was below the €20,000 willingness to pay threshold) compared with urinary gonadotrophins in the</li> </ul>

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				Italian National Health System [34].
<b>Cost-benefit analysis (CBA)</b>		<b>Monetary equivalents</b>	<ul style="list-style-type: none"> <li>• Monetary willingness to pay (WTP)</li> <li>• Monetary value of a statistical life (VSL)</li> </ul>	<ul style="list-style-type: none"> <li>• A CBA based on the WTP approach for intra-uterine insemination and assisted reproductive technology highlighted the substantial role of financial constraints on the valuation of advanced fertility treatments in communities in Iran [35].</li> </ul>
<b>Cost-consequence analysis (CCA)</b>		<b>Multiple outcomes used:</b> most appropriate unit for every outcome (information in disaggregated format)	<ul style="list-style-type: none"> <li>• Pregnancies achieved</li> <li>• Live births</li> <li>• QALY</li> <li>• Monetary return on investment (ROI)</li> </ul>	<ul style="list-style-type: none"> <li>• A modelling study compared single and double embryo transfer strategies in terms of costs to the health service, live birth rate, singleton live birth rate, twin pregnancy rate, incidence of disability among infants born, and QALYs accruing to women. The cost-effectiveness of double embryo transfer improves with age, and may be considered cost-effective in some groups of older women. The decision may best be considered on a case-by-case basis for women aged 37–39 years [36].</li> </ul>

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## Practice Points

- Cost-effectiveness analyses help to identify those MAR programmes that optimize live birth rates
- Cost-utility analyses and societal cost-benefit analyses address the degree to which healthcare systems should treat the condition of infertility or increase spending on MAR treatments compared with other healthcare domains
- The comparator used must be appropriate for the patient population and setting
- The time horizon should be reported, and should be long enough to capture any relevant differences in costs/outcomes between different approaches being compared
- For narrow health economic analyses of interventions by their ability to achieve a well-defined set of outcomes, first live birth or cumulative live birth are logical outcome measures.
- Broader health economic analyses that look at the allocation of resources across many healthcare domains and beyond must consider which outcomes will be most appropriate
- To summarize the results, the average cost-effectiveness ratio and the incremental cost effectiveness ratio may be used

## **Research Agenda**

- The preference of society and the willingness to trade-off health benefits for other groups of patients against live births delivered through the provision of MAR
- The extent to which economic evaluation studies inform choices for privately paying patients and for public or private MAR teams
- The concept of willingness-to-pay needs to be further explored from both a societal and a patient perspective, taking into account existing benchmarks where possible.

**Multiple Choice Questions**

**Question 1:** A policy maker who needs to judge whether MAR budgets should be expanded or reduced when compared to other disease budgets should preferably base this decision on the results obtained through:

- A. Cost-effectiveness analyses
- B. Cost-utility analyses
- C. Cost-consequences analyses
- D. Cost-benefit analyses
- E. Cost-of-illness studies

*Correct answer:* D. The policy maker will need to judge the overall return received for the money invested in MAR and will have to compare that result to the one from other programs. When budgets only need to be optimized across disease or health areas, cost-utility analyses will provide the necessary information. When budgets need to be optimized across *all* policy areas, including those outside the health domain, cost-benefit analyses would be necessary.

**Question 2:** When a new MAR program is found to be costlier than the standard of care, but it also leads to more births in the treated population, then, from a health-economic perspective, a policymaker should:

- A. Not recommend the program for funding because it increases costs
- B. Recommend the program for funding because it leads to more live births and reduces the burden of infertility
- C. Recommend the program for funding when the average cost per live birth of the new program is lower than the average cost per live birth of the standard of care
- D. Recommend the program for funding when the incremental costs per live birth of the new program are lower than the societal willingness-to-pay for an additional child
- E. Recommend the program for funding when the incremental costs per live birth of the new program are lower than the societal willingness-to-pay for an additional child plus the infertility problems of its parents

*Correct answer:* E. Judging whether a new MAR program deserves funding cannot be done based on costs or outcomes separately but requires a comparison of both. However, comparing average costs and effects can be misleading as a program can still be cost-effective while having a higher average cost per live birth than its alternative, simply because its additional costs may be justified. To make that judgment one needs to look at the 'incremental' costs and compare these to the incremental effects gained. The latter will consist of both a new child and fertility problems of the parents. The value of these elements will ultimately determine whether the additional costs are 'acceptable'.



**Question 3:** Which cost category is irrelevant when we want to calculate the cost-effectiveness of a MAR program from a healthcare payer perspective?

- A. Costs of making an infertility diagnosis before MAR starts
- B. Costs of maternal follow-up after birth
- C. Costs of pregnancy complications
- D. Costs of work absenteeism during the MAR treatment
- E. Costs of psychological counselling for infertile couples

*Correct answer:* D. From a payer perspective all 'direct' healthcare costs will be relevant.

However, often short time horizons are used in economic evaluation of MAR, excluding postnatal follow up or complications of treatment, but these can differ across treatments and hence should be considered. Also, psychological counselling costs (e.g., incurred in a do nothing scenario) are relevant to the payer perspective. Indirect costs of productivity losses and work absenteeism are usually not born by healthcare payers but by society. These can be excluded in an analysis from a payer perspective, even though they are important from an employer perspective and from an overall societal perspective.