Observations suggest that ART induces vascular dysfunction both in humans and animals

Assisted reproduction: a novel cardiovascular risk factor

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Summary

There is abundant evidence indicating that pathological events during early life predispose to cardiovascular and metabolic diseases later in life. Recent evidence suggests that assisted reproductive technologies (ART) represent a novel important example of this problem. Here we will review recent data in humans and animals demonstrating ART-induced cardiac dysfunction, premature vascular ageing and arterial hypertension that appear to be related, at least of part, to epigenetic mechanisms. Therefore, a better comprehension of mechanisms of ART-induced vascular dysfunction and long-term monitoring of the ART population is of upmost importance in order to prevent or treat the long-term cardiovascular consequences. Given the young age of the ART population, we will have to wait for 20–30 years before we will know in how many premature cardiovascular endpoints these ART-induced alterations will result, ART should be taken into account when counseling putting into place preventive strategies.

Key words: in vitro fertilisation; vascular function; epigenetics; nitric oxide; hypertension

Introduction

Developmental origin of adult diseases

Epidemiological studies have shown an association between pathological events occurring during early life and the development of cardiovascular and metabolic disease in adulthood, leading to the “Barker’s hypothesis of fetal programming of adult disease” [1]. In line with this hypothesis, we have shown that transient perinatal hypoxia induces pulmonary vascular dysfunction that predisposes to exaggerated hypoxic pulmonary hypertension in young apparently healthy adults [2], and preeclampsia induces pulmonary and systemic vascular dysfunction in young apparently healthy offspring [3]. Taken together, these data demonstrate that environmental insults occurring during the perinatal period and the last trimester of gestation have important long-term consequences for the regulation of the cardiovascular system in humans. Studies in a mouse model of preeclampsia indicate that the altered vascular responsiveness in the offspring is related to epigenetic changes [4]. Assisted reproductive technologies (ART) involve the manipulation of early embryos during a period when they may be particularly vulnerable. In line with this concept, recent evidence indicates that ART represents a novel important example of this problem [5–7].

Studies in humans

Vascular dysfunction in children conceived by means of assisted reproductive technologies

The prevalence of infertility has been estimated to be 9% worldwide [8]. ART has allowed millions of infertile couples to have children, and in developed countries ART children now make up for 2% to 5% of births [9]. In a recent study, we found that ART children display marked generalised vascular dysfunction, as evidenced by a roughly 30% higher hypoxic pulmonary artery pressure, a 25% lower flow-mediated dilation (FMD), a markedly faster pulse-wave velocity (PWV) and increased intima-media thickness (IMT) in ART than in control children (fig. 1) [7]. In order to exclude confounding factors that could contribute to ART-induced vascular dysfunction, we studied vascular function in parents of ART children, children conceived after ovarian hyperstimulation alone and naturally conceived siblings of ART children. Vascular function was normal in all these groups (fig. 1), suggesting that ART per se induces vascular dysfunction. In the pulmonary circulation, this defect predisposes to exaggerated hypoxic pulmonary hypertension, already during childhood. In the systemic circulation, endothelial dysfunction, increased vascular stiffness and IMT represent early steps in the development of atherosclerosis and are independent risk factors for premature cardiovascular morbidity [10]. Recent data suggest that vascular dysfunction in ART children may be reversible [11]. In ART mice, vascular dysfunction is related to decreased vascular endothelial nitric oxide (NO) synthase expression and NO synthesis (see below) [5]. We speculated that a similar mechanism underpins vascular dysfunction in ART children. We found that vascular dysfunction in ART children was associated with increased oxidative stress and decreased plasma NO [11]. Antioxidant administration normalised oxidative stress and plasma NO and...
improved endothelial function in the systemic and pulmonary circulation. These data show that in young individuals ART-induced vascular dysfunction is subject to redox-regulation and potentially reversible [11].

Blood pressure in children conceived by means of assisted reproductive technologies
Among the potential long-term consequences of ART-mediated vascular dysfunction, arterial hypertension may represent an important problem [12]. In our study, arterial blood pressure was not different between ART and control children (mean age 11.1 years) [7]. Belva et al. also found no difference in arterial blood pressure between 14-year-old ART and naturally conceived adolescents [13]. However, in a larger study (225 ART children and 225 spontaneously conceived control children) including older subjects (age 8–18 years), Ceelen et al. found that office systolic and diastolic blood pressure were higher in ART children than in controls [14]. Taken together these data suggest that the prevalence of HTA in this population may increase with increasing age. Consistent with this concept, adult ART mice display arterial hypertension (see below).

Additional studies using ambulatory blood pressure monitoring (ABPM) are needed in this population in order to follow the evolution of arterial blood pressure and to determine whether ART-induced vascular dysfunction translates into arterial hypertension.

Cardiac dysfunction in children conceived by means of assisted reproductive technologies
A very recent study showed that children conceived by means of ART manifest cardiac and vascular remodelling in utero that persists in early postnatal life [15]. This study compared cardiac function and morphology in 100 foetuses conceived with ART and 100 control pregnancies. In ART foetuses, the myocardial wall was thicker and the longitudinal heart function was decreased, as shown by a decreased tricuspid ring displacement, impaired relaxation and dilated atria. These signs of cardiovascular remodelling persisted at 6 months of age. However, it is not known how they will evolve later in life. To provide such information, we recently assessed cardiac function in older ART children and adolescents (Von Arx R et al.). We found that, whereas left ventricular function was normal and comparable to that of naturally conceived children, right ventricular end-diastolic area was increased (cardiac remodelling) and right ventricular diastolic function was impaired, particularly under stressful conditions (high altitude exposure). These data suggest that ART-induced cardiac remodelling persists into adulthood and may predispose to cardiac dysfunction later in life.

To assess long-term consequence of ART on cardiovascular function and mortality, and examine potential underlying mechanisms, we studied mice generated by ART.
Studies in mice

Vascular dysfunction in mice generated by assisted reproductive technologies

Consistent with data in humans, we found that ART induces systemic vascular dysfunction also in mice, as evidenced by decreased endothelial-dependent mesenteric artery vasodilation (fig. 2 panel A) and increased carotid artery stiffness (fig. 2 panel B) [5]. Notably, this vascular dysfunction in vitro translated into arterial hypertension in vivo in adult ART mice (fig. 2 panel C). To study the long-term consequences of ART-induced vascular dysfunction, we compared the lifespan of ART mice with that of control mice fed a Western-style high-fat diet. The lifespan was roughly 25% shorter in ART than in control mice, demonstrating the long-term consequences of ART-induced cardiovascular dysfunction (fig. 2 panel D).

Underlying mechanisms

The demonstration of ART-induced cardiovascular dysfunction in normal fertile mice strengthens the concept that ART per se is the main culprit causing this problem. ART implies manipulation of the embryo during a vulnerable period with increased epigenetic activity (fig. 3 panel A) [6]. The term “epigenetics” refers to chromatin-based mechanisms that regulate gene expression without affecting the DNA sequence and can be transmitted to the next generation(s). The best studied epigenetic mechanisms involve DNA methylation (fig. 3 panel B) and histone modifications (acetylation and methylation) [16]. Several observations indicate that epigenetic mechanisms underpin ART-induced cardiovascular alterations. First, male ART mice transmit vascular dysfunction to the next generation. Second, the methylation of the promoter of the endothelial nitric oxide synthase (eNOS) gene in vascular tissue is increased in ART mice. Third, DNA demethylation of the eNOS translated into decreased eNOS expression in the vasculature and decreased vascular NO synthesis in ART compared with control mice. Taken together, these findings indicate that in mice, ART alters the entire chain of events starting...
from eNOS promoter methylation and ending in premature vascular aging and arterial hypertension (fig. 3 panel C) [6]. Finally, epigenetic alterations can be reversed by drugs such as histone deacetylase inhibitors (i.e., butyrate) [4]. We found that butyrate administration normalised eNOS promoter methylation and vascular function in ART mice, and prevented the transmission of this problem to the next generation [5]. Collectively these data indicate that ART induces premature vascular ageing and hypertension in mice by an epigenetic mechanism.

We then started out to test for specific events during embryo culture that may contribute to ART-induced epigenetic and cardiovascular alterations. The time passed by the embryo on the culture medium before its transfer to the recipient mother may play a role. We therefore compared vascular function in mice born after implantation of two cell embryos (24 h in culture media) and blastocysts (86 h in culture media). Vascular dysfunction was comparable in the two groups, suggesting that shortening the culture time does not prevent ART-induced vascular dysfunction. In line with this observation, altered methylation patterns and gene expression are already detectable in two cell stage embryos [17]. Taken together, these data suggest that suboptimal culture conditions contribute to ART-induced epigenetic alterations and vascular dysfunction. In line with this hypothesis, preliminary data in mice suggest that modification of the culture media may attenuate ART-induced epigenetic alterations and vascular dysfunction in mice [18].

**Conclusion**

In conclusion, these observations suggest that ART per se induces vascular dysfunction both in humans and animals. In mice, endothelial dysfunction and arterial hypertension are induced during the embryo development in culture media by an epigenetic mechanism. A similar mechanism may also underpin ART-induced vascular dysfunction in humans. Comparison with young populations with similar vascular dysfunction at a young age and known cardiovascular complications later in life (i.e., preeclampsia), suggest that ART may represent an important novel risk factor predisposing to premature cardiovascular morbidity and mortality (fig. 4) [19]. Given the young age of this population (the first ART
child was born in 1978), the magnitude of the problem is not yet known, since clinically manifest cardiovascular disease has not had time to develop. Now, however, it is already important to avoid traditional cardiovascular risk factors in this population and to detect early cardiovascular alterations with the aim to prevent or at least optimally treat cardiovascular complications.

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**References**