Ovarian function in domestic ruminants: Mechanistic and translational aspects

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Abstract

The purpose of this review is to illustrate the critical value of integration of both in vitro and whole animal approaches in order to make real progress in terms of understanding the underlying mechanisms controlling ovarian follicle development and also in utilising domestic ruminants as models for the human in the translational development of interventions to address infertility. In terms of mechanistic aspects, this review concentrates on interactions between members of the TGFβ super family derived from both somatic cells (AMH) and the oocyte (BMP6, BMP15, GDF9) and illustrates how integration of in vitro and in vivo approaches has allowed us to propose a functional relationship between these factors in regulating the progression of follicles through the gonadotrophin-responsive to dependent phases of follicle development. In this section, a new in situ direct ovarian perfusion system is introduced. In terms of translational aspects, this review details the integration of in vitro and in vivo approaches in order to develop methods in sheep that allow high rates of restoration of ovarian function and fertility following whole ovarian cryopreservation and autotransplantation in adult animals. This methodology should be of value to women at risk of premature ovarian failure. It is concluded that, given the vastly increased cost of ruminant research in developed countries, that meetings such as the RRS can play a critical role in encouraging international collaboration in order to ensure the continued exploitation of monovulatory ruminants as valuable experimental models.

Introduction

It is now well established that ovarian follicle and oocyte development is controlled by a complex array of autocrine and paracrine factors secreted by both the oocyte and follicular somatic cells which interact to initially control follicle development locally during initial gonadotrophin-independent stages

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of follicles development and then increasingly act to modulate the responsiveness of ovarian granulosa and theca cells to the pituitary gonadotrophins through the subsequent gonadotrophin-responsive and -dependent stages of follicle development (Campbell et al., 1995; Webb and Campbell, 2007a; Webb et al., 1999). Advances in the development of physiological, serum-free culture systems for preantral follicles (Picton et al., 2003), somatic cells (Campbell et al., 1996; Campbell et al., 1998) and oocytes (Picton et al., 2003) allied with the enormous power of the “omic” technologies has resulted in the implication of a vast array of local factors from a range of major local regulatory systems (see below) in the control of major development checkpoints during follicle development. The complex nature of these inter-relationships between interacting systems has led to a degree of uncertainty as to the actual identity of the key regulatory systems controlling follicular development and this problem is compounded by contradictory evidence, much of which can be attributed to species differences in the role and structure of the same local factors/system.

This problem is particularly evident in ruminant reproduction research, where despite the economic value of these species and their usefulness as experimental models for the human, there is limited commercial availability of species specific reagents and/or sequence information when compared to polyovular rodents and the human. Further, the current lack of stem cell and specific knock-out technology severely limits our ability to conduct the type of interventionist in vivo studies required to elucidate key development pathways in ruminant species. In this regard it is telling that some of the major advances in ruminant reproduction research in the last decade have come from identification of the naturally occurring mutations in sheep which were known to result in infertility and/or increased prolificacy (McNatty et al., 2005).

Prof George Eric Lamming was an outstanding animal physiologist who understood the need to refer to the whole animal in order to make real progress in understanding reproductive processes that will in turn lead to advances to boost production and/or treat infertility. It is therefore appropriate that the main objective of this review is to emphasise the critical nature of integration of both in vitro and in vivo approaches in ruminant reproduction research, despite the vastly increased cost of this type of research (at least in the UK). Thus after a brief summary of folliculogenesis, we plan to illustrate this point through recent research from our laboratories which will emphasise the value of some classic technology and also introduce some new in vivo models to facilitate both mechanistic and translational aspects of ovarian research. It should be emphasised that this review is not intended to cover the whole of the complex endocrine and local regulatory system controlling follicle development but rather to highlight some aspects of these local regulatory systems in order to illustrate the value of an integrated experimental approach.

**Ovulatory follicle development**

Ovarian folliculogenesis is a lengthy and intricately regulated process marked by dramatic proliferation and precisely orchestrated differentiation of both the somatic and germ cell elements. The ovarian follicle is the fundamental developmental unit of the mammalian ovary, consisting of a germ cell (oocyte) in close association with somatic cells (granulosa cells). Primordial follicles represent the source from which follicles will be recruited for growth throughout life and as the paired ovaries of an individual contain around 100,000-250,000 of these follicles at birth (Cahill, 1981; Turnbull et al., 1977). Once follicles have been initiated to grow, the granulosa cells proliferate to form multilaminar structures (preantral follicles) which subsequently form a fluid filled space (antrum) and a well differentiated theca layer (Young and McNeilly, 2010). Follicular development in humans, sheep and cattle is a lengthy process, taking around 4-6 months (Turnbull et al., 1977; Webb and Campbell, 2007a) with the majority of this time (4 months) being spent in the preantral stages of development. Gap junction-mediated communication between the oocyte and the surrounding somatic cells
Ovarian function in domestic ruminants is essential for the coordinated development of both cell types, making folliculogenesis a highly synchronised process. The factors that initiate this process, the recruitment of primordial follicles, remain unknown although a large number of potential modulators have been identified (Picton et al., 2008; Tingen et al., 2009). Following follicle recruitment, the oocyte enlarges, accumulating maternal transcripts, proteins and organelles that will support embryonic development. At the same time, the granulosa cells change shape and start to proliferate and differentiate, becoming responsive to gonadotrophic signals (Picton et al., 2008). Oocyte-somatic cell communication is maintained throughout follicle growth, during which time somatic cells provide the oocyte with metabolic substrates and meiosis-arresting signals (Picton et al., 2008). When the follicle reaches a diameter of 200-400 µm, the antrum develops and this is followed by widespread atresia (50-70% for follicles over 1 mm) so that the vast majority (>99%) of follicles fail to ovulate. Follicle development from the late preantral stage to the small antral stage (2 mm in sheep, 4-5 mm in cattle and humans) is termed gonadotrophin-responsive because although development of these follicles can be influenced by FSH, the growth of follicles to these stages still occurs in the absence of FSH (Dufour et al., 1979; Webb and Campbell, 2007a). The transition from the gonadotrophin-responsive to the subsequent gonadotrophin-dependent phase is characterised by a dramatic decline in the mitotic index of the ovarian somatic cells (Fig. 1) as differentiation of these cells occurs. Further development of antral follicles, however, is termed gonadotrophin-dependent as FSH must be maintained above a critical threshold concentration (Baird, 1983; Webb et al., 1999) in order to support growth to an

Fig. 1. Schematic diagram illustrating changes in oocyte diameter (A) and relative level of maximal AMH expression (B) in relation to changes in granulosa cell mitotic index during follicle development in sheep (C). Note that majority of oocyte growth occurs prior to rapid increase in mitotic index in somatic cells which is temporarily related to pattern of AMH expression. C redrawn from Turnbull et al., 1977.
ovulatory size, at which point the follicle is highly vascularised, the granulosa cells have high levels of aromatase and LH receptor expression and contains a cytoplasmically mature, developmentally competent oocyte. The mechanisms regulating the recruitment, selection and dominance of the ovulatory follicle in monovulatory species has been the subject of intensive investigation and the occurrence of spontaneous mutations in sheep which result in dysregulation of this process, such as the FecB mutation (see below) are valuable experimental models for elucidating the underlying mechanisms of this selection process.

The role of potential ligands for BMPR1B (Alk6) on ovine follicle development

In the following section we will relate some recent work from our laboratories in order to illustrate the value of integration of both in vitro cell culture and analytical techniques with whole animal studies to ensure significant progress in our understanding of the mechanisms regulating follicular development. The initial impetus for this work came from the identification of the FecB Booroola mutation, which resulted in marked increases in ovulation rate due to precocious ovulatory follicle maturation, as being due to a single point mutation in the intracellular domain of one of the receptors for bone morphogenetic proteins (BMPR1B, also known as Alk6) (Mulsant et al., 2001; Souza et al., 2001; Wilson et al., 2001). We have previously presented evidence to suggest that the FecB mutation exerts its action at the level of the ovary (Campbell et al., 2003a) rather than modulating pituitary gonadotrophin release (McNatty and Henderson, 1987), and more recent data in which depression in FSH levels was shown to negate the effect of the FecB mutation on prolificacy, confirmed this hypothesis by providing unequivocal evidence that the FecB mutation acts through modulation of gonadotrophic signals (AS McNeilly, DT Baird BK Campbell unpublished results). Our resultant studies on potential ligands for Alk6, including bone morphogenetic protein (BMP) 6, anti-mullerian hormone (AMH) and BMP15 (Miyazono et al., 2005; Shi and Massague, 2003), described below, have revealed a much wider range of actions for these factors than those associated with just ovulatory follicle selection. Readers are referred to a number of excellent reviews for detailed descriptions of the BMP system in the ovary (Juengel et al., 2004a; Knight and Glister, 2006; McNatty et al., 2005).

BMP6

Initial cell culture studies in sheep showed that human recombinant BMP2, BMP4 and BMP6 were equally potent enhancers of FSH-induced oestradiol production by sheep granulosa cells whilst in theca cells (TC), similar doses of these BMPs (5-50 ng/ml) inhibited LH-stimulated androstenedione (A4) production whereas lower doses (0.005-0.05 ng/ml) stimulated TC proliferation and hence total androstenedione production (Campbell et al., 2006). Similar findings to these had also been made in the bovine (Glister et al., 2005). In order to resolve these contradictory findings, BMP6, which had been identified as the most highly expressed BMP in sheep GC (Campbell et al., 2006; Juengel et al., 2006), was infused directly into the ovary of sheep with ovarian autotransplants (Campbell et al., 2009). Treated (n = 6) and vehicle treated controls (n = 6) were infused during the early follicular phase, between 12-24 h after luteal regression and ovarian response determined by collection of samples of ovarian venous blood and transdermal ultrasound. In the absence of any change in circulating gonadotrophins or in the antral follicle population, BMP6 infusion resulted in acute but transient increases in ovarian inhibin A, androstenedione and oestradiol secretion during the second half of the infusion period (Fig. 2). Thereafter, treated animals also had an advance in the time of the LH surge by around 10 h and smaller pre-ovulatory follicles and
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corpora lutea, but without an increase in ovulation rate. This study therefore clarified that BMP6 acts as an enhancer of gonadotrophin-stimulated follicular development, with the discrepancy between in vitro and in vivo results in terms of androstenedione secretion being explained by the upregulation in ovarian inhibin secretion which is known to be a potent paracrine stimulator of thecal androgen production (Campbell and Baird, 2001; Hillier and Miro, 1993) through blockade of the inhibitory action of activin (Young and McNeilly, 2012) which is produced early in antral follicular development and also suppresses thecal androgen production (Young et al., 2012). Overall these data provided strong in vivo evidence to support the hypothesis that BMP6 is an important local regulator of ovarian function and that alterations in BMP6 cellular signalling may explain some of the effects of the FecB mutation in inducing precocious maturation of ovulatory follicles.

AMH

Although AMH was first identified in relation to its regulatory role in male sex differentiation, after birth this sexually dimorphic expression pattern is lost and AMH protein is also expressed in granulosa cells of growing follicles in the ovary (Themmen, 2005; Webb et al., 1999). In rodents (Durlinger et al., 2002; Webb et al., 1999), humans (Webb et al., 1999; Weenen et al., 2004) and
sheep (Campbell et al., 2012) AMH protein expression starts in the columnar granulosa cells of primary follicles, is highest in granulosa cells of preantral and small antral follicles and gradually diminishes in the subsequent stages of follicle development so that AMH is no longer expressed during the gonadotropin-dependent terminal stages of follicle development (Fig. 1). In addition, AMH expression disappears when follicles become atretic. Amh null mice have increased rates of primordial follicle recruitment and resulting in premature ovarian failure. Further, AMH has been shown to inhibit gonadotropin-dependent growth of cultured mouse preantral follicles (Durlinger et al., 2002; Themmen, 2005; Webb et al., 1999) and in rat, pig (di Clemente et al., 1994; Webb et al., 1999) and sheep (Campbell et al., 2012) granulosa cell cultures, AMH attenuates the FSH-dependent increase in aromatase activity and LH receptor expression. Further, in sheep we were also able to show that AMH is a potent inhibitor of LH-stimulated thecal androgen synthesis in vitro (Campbell et al., 2012) and we have subsequently confirmed that theca cells do express the specific Type 2 AMH receptor (BK Campbell, P Marsters unpublished results). Thus the available evidence supported a functional role for AMH in modulating both the early and terminal stages of follicle development in polyovular species. However, in relation to early follicle development the evidence was more equivocal for monovulatory species, in that the literature contained two reports which supported a role for AMH in the suppression of primordial follicle initiation in vitro in the human (Carlsson et al., 2006) and bovine (Gigli et al., 2005), whereas another study reported the opposite in humans (Schmidt et al., 2005).

In contrast to BMP6, direct ovarian infusion of AMH in ewes with ovarian autotransplants had no effect on ovarian steroid secretion or the pattern of ovulatory ovarian follicle development (Campbell et al., 2012). The converse experiment however, a knock-down utilising active immunisation in lieu of a naturally occurring mutation, had no effect on primordial follicle number but did lead to a decline in the population of gonadotrophin-responsive preantral and small antral follicles and increases in both the number of gonadotrophin-dependent antral follicles and ovulation rate (Fig. 3; (Campbell et al., 2012). In contrast to the rodent, the results indicate that AMH does not affect the rate of primordial follicle recruitment but appears to regulate the rate at which follicles progress through the gonadotropin-responsive phase, during which time it is maximally expressed. Further, the existence of a functional link between AMH and theca cell function in vivo was confirmed by this experiment as antral fluid levels of androstenedione were greatly elevated in AMH-immunised animals (Campbell et al., 2012) and we have subsequently shown that supplementation of sheep with the weak adrenal androgen DHEA, results in perturbation in AMH protein expression and an increase in the number of gonadotrophin-responsive small antral follicles. The elucidation of a functional relationship between AMH and LH-stimulated thecal androgen production is significant in terms of the aetiology of common forms of anovulatory infertility in women (Franks et al., 2008) and the ability to manipulate the number of recruitable antral follicles may have therapeutic value in women who respond poorly to ovarian stimulation. Overall, this experiment also emphasises the continued value of active immunisation as a means to perturb specific local regulatory systems in domestic ruminants in the absence of natural or induced knock-down models and this conclusion is supported by the findings of Juengel et al. who showed similar ovarian responses to low and high levels of immunisation against BMP15 as observed in ewes that carried one or two copies of the FecX BMP15 mutation (Galloway et al., 2000; Juengel et al., 2004b).

Although the studies described above indicate that AMH represents an important local attenuator of gonadotrophin-induced somatic cell differentiation they also demonstrated that contrary to expectations, AMH cannot be directly implicated in the mechanism of action of the FecB mutation as in a variety of ovarian somatic cell types AMH did not signal via the BMP pathway to phosphorylate SMAD1, SMAD5 and SMAD8 (L Nicol, AS McNeilly, BK Campbell unpublished work).
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Parallel studies utilising a reporter assay for multiple transcriptional responsive elements (Cignal 45-Pathway Reporter Array, SABiosciences, Crawley, U.K.) in human granulosa cell lines confirmed this result and indicated that AMH increases activity of a range of transcription factors including ERE, ARE, SP1 and NFAT (J Hernandez-Medrano, BK Campbell unpublished results) and further work is required to extend these studies. A further area of uncertainty in relation to AMH is the identity of factor(s) that stimulate its expression in granulosa cells during the very early stages of follicle development. A clue to this problem was however observed in our initial immunohistochemical (IHC) studies examining the pattern of AMH protein expression at different stages of follicle development (Campbell et al., 2012) where a marked inverse relationship was observed between AMH and aromatase protein expression. This inverse relationship was most notable in large oestrogenic follicles where the cumulus area surrounding the oocyte

![Graph A](image)

**Fig. 3.** Effect of active immunisation against AMH on the follicle population derived from either histological evaluation of the early follicle population ≤ 1 mm diameter in one ovary (a) or dissection of visible antral follicles > 1 mm diameter from the contra-lateral ovary in each animal (b). Note that immunisation in high titre animals resulted in an increase in the population of both small and large antral follicles which was associated with a decline in the population of small antral follicles less then 1 mm in diameter. It was concluded that AMH, which is maximally expressed in late preantral and small antral follicles regulates the rate at which follicle pass through this gonadotrophin-responsive phase. Ovulation rates in AMH immunised animals were also increased. Mean ± SEM. * P<0.05, **P<0.01 for comparison with controls. Data published in Campbell et al., 2012.
had high levels of AMH expression and low aromatase whereas the membrana granulosa cell layer displayed the inverse pattern of expression (Fig. 4). It appears extremely likely from this observation that maintenance of AMH expression in the undifferentiated cumulus cell layer of these oestrogenic follicles is due to oocyte secreted factors and follow up experiments utilising knock-down approaches in vitro have indicated functional links between granulosa cell AMH expression and BMP15, GDF9 and SOX8 (HM Picton, BK Campbell unpublished results). On the basis of these observations we have proposed the mechanistic model presented in Fig. 4 which proposes that AMH is upregulated in somatic cells by oocyte secreted factors (OSF) and that the down-regulation in AMH expression that occurs as follicles move into the gonadotrophin-dependent phase occurs in response to either a decrease in the concentration of OSF across this morphogen gradient and/or induction of LH-stimulated thecal androgen secretion which acts as a negative modulator of AMH expression.

**Fig. 4.** Immunohistochemistry showing AMH (A) and aromatase (CYP19A1) (B) proteins in consecutive sections from the same oestrogenic large antral follicle showing that AMH is absent from the granulosa cell layer but present in the coronal and cumulus cells surrounding the oocyte. Note that aromatase abundance is inverse to AMH abundance. C proposed model explaining pattern expression of AMH shown in A/B that that AMH is upregulated in somatic cells by OSF and that the down-regulation in AMH expression that occurs as follicles move into the gonadotrophin-dependent phase occurs in response to either a decrease in the concentration of OSF across this morphogen gradient and/or induction of LH-stimulated thecal androgen secretion which acts as a negative modulator of AMH expression. Images A and C previously published in Campbell et al., 2012.
BMP15

The consequences of naturally occurring mutations (Galloway et al., 2000) and/or active immunisation against BMP15 (Juengel et al., 2004b) are well described in the literature and it is clear that BMP15 is major local regulator of ovarian function. There is some evidence to support a role for BMP15 in the mechanism of action of the FecB mutation in that BMP15 expression in oocytes has been shown to be lower in FecB animals (Crawford et al., 2011) and ewes who are heterozygous for the FecX BMP15 mutation show some evidence of the precocious ovulatory follicles phenotype that is so characteristic of the Booroola (McNatty et al., 2009). In our own studies with Booroola ewes we were unable to demonstrate any effect of the FecB mutation in Alk6 on the ability of BMP15 to phosphorylate SMAD1, SMAD5 and SMAD8 (L Nicol, AS McNeilly, BK Campbell unpublished results) and parallel cell line reporter studies have also produced equivocal results (PM Marsters and BK Campbell unpublished results). The biology of BMP15, however, is complex and this is well illustrated by cell culture studies which have shown that low doses of rhBMP15 alone are mildly stimulatory to FSH-stimulated oestradiol production by granulosa cells while higher doses are markedly inhibitory. BMP15 is, however, known to form heterodimers with GDF9 (Mottershead et al., 2013) and exposure of granulosa cells to both rmGDF9 and rhBMP15 induced a 10-fold increase in oestradiol production when compared to the same doses in isolation (A Marsh, J Hernandez-Medrano, BK Campbell unpublished results).

In order to test the in vivo action of ovarian BMP15 exposure on ovulatory follicle development we have developed an alternative in situ infusion model which avoids the surgical complexity.

Fig. 5. Schematic diagram showing an in situ infusion model which involves the cannulation of the side-branch of the ovarian artery which supplies the broad ligament and the placement of another venous cannula in the uterine vein which is advanced so its tip lies on the common utero-ovarian vein. Cannulation of both ovaries in each animal in this manner allows one ovary to act as a vehicle only control for direct ovarian infusion of test factors via the ovarian artery following exteriorisation of all cannulae through the abdominal wall. This model avoids the surgical complexity of the ovarian autotransplant model and also allows collection of ovarian tissue for histological and molecular analysis of effects of infusion. Diagram redrawn from Lee et al. (Lee et al., 2010).
of the ovarian autotransplant model and allows within animal comparisons. This model involves the cannulation of the side-branch of the ovarian artery which supplies the broad ligament and the placement of another venous cannula in the uterine vein which is advanced so its tip lies on the common utero-ovarian vein (Fig 5). Cannulation of both ovaries in each animal in this manner allows one ovary to act as a vehicle only control for direct ovarian infusion of test factors via the ovarian artery following exteriorisation of all cannulae through the abdominal wall. Utilising either this in situ cannulation model or the ovarian autotransplant, contrary to expectations direct ovarian infusion of rhBMP15 resulted in small but statistically significant increases in ovarian androstenedione and oestradiol secretion. This interesting result was confirmed by direct ovarian infusion of antisera raised against specific ovine BMP15 sequences (J Hernandez-Medrano, BK Campbell unpublished results). The biology of BMP15 in ruminant species therefore appears more complex than indicated by the FecX mutants and further work is required, preferably with ovine full size forms of the molecule, to elucidate its actions in vivo in relation to interactions with GDF9 and possible indirect activation of somatic cell factors such as AMH (Fig. 4).

In summary, the integration of in vitro and in vivo approaches has allowed us to propose a functional relationship between members of the transforming growth factor beta super family derived from both somatic cells (AMH) and the oocyte (BMP6, BMP15, GDF9) in regulating the progression of follicles through the gonadotrophin-responsive to dependent phases of follicle development. However, much more work is required to elucidate the complex biology of members of this system e.g. BMP15 and to integrate the actions of the BMP system with other major local regulatory systems such as the IGF system (Webb and Campbell, 2007b).

Ruminant models of fertility restoration in women at risk of premature ovarian failure

The domestic ruminant has been utilised for many years as an appropriate animal model for the human, due to similarities in size (sheep) and physiology, particularly in relation to the dynamics and timing of follicle development and their monovulatory nature (Campbell et al., 2003b). In the following section we will further illustrate the value of integration of both in vitro cell culture and analytical techniques with whole animal studies to ensure significant progress in the development of interventions to address human infertility. Specifically, we have used sheep as models for fertility preservation in women at risk of premature ovarian failure (POF), which occurs in 1-2% of women (Coulam et al., 1986). For a number of reasons such as genetic predisposition (Santoro, 2003) and gonadotoxic treatment (Anderson and Wallace, 2011), the rate of follicle loss is accelerated in some women, leading to POF and early menopause (<40 years old). In addition to primary infertility and amenorrhea, POF can have far-reaching clinical and psychological effects such as failure to develop secondary sexual characteristics, loss of self-confidence, depression, increased risk of osteoporosis, and heart disease (Traub and Santoro, 2010). Due to the continued improvement in childhood cancer patient survival rates, the importance of POF and its impact on human health is expanding (Anderson and Wallace, 2011). In instances when ovarian failure can be predicted, patches of ovarian cortex and the oocytes within it, can be removed and protected from damage or further deterioration in the frozen state and then autografted without vascular reanastomosis to resume development when the restoration of ovarian function is required (Donnez et al., 2013; Gosden and Nagano, 2002). The first demonstration of restoration of fertility following autografting was made in lambs in 1994 (Gosden et al., 1994) and this pioneering paper was followed by a larger scale study examining the endocrine consequences and period of restoration of fertility following cortical autografting (Baird et al., 1999). Some 5 years later the first report of a human birth from a
woman who had received frozen cortical autografts was published (Donnez et al., 2004) and at the present time autografting of ovarian cortical tissue offers the only current proven means of restoring both fertility and ovarian function following POF in women. Although shown to be effective clinically (Donnez et al., 2013) autografting returns only a small proportion of the ovarian reserve and many of these follicles are lost due to post-graft ischaemia. This intervention is therefore less effective in older women, is associated with endocrine disturbance, involves delays of several months before ovarian function is restored and when restored the duration of ovarian function is relatively short term.

In order to try and overcome these limitations of autografting, for a number of years we have been looking at the possibility of whole ovarian cryopreservation and autotransplantation (WOCP&TP) as a superior alternative treatment path. WOCP&TP offers a complete solution for the long-term preservation of ovarian function and fertility in girls and women of all ages at risk of POF. This is because WOCP&TP involves acute preservation of a whole ovary and theoretically should not diminish the ovarian reserve, result in endocrine imbalances or require a significant delay before cyclicity is restored. Further, there should be no age constraints for the long-term restoration of ovarian function.

The primary challenge with organ cryopreservation is the crucial need to preserve an intact vascular system to allow for subsequent organ revascularization and function (Pegg, 2010). All methods of cryopreservation require the replacement of the majority of the tissue water content with a cryoprotective agent (CPA). For organ preservation this has usually been achieved by vascular perfusion. While this approach creates very short diffusion distances for the CPA and water it has the disadvantage of causing microvascular damage (Pegg, 2010). From our understanding of the mechanisms of cellular damage and the mode of action of CPAs it is clear that while intra-cell injury caused by ice formation can usually be avoided, the formation of ice in extracellular locations, particularly in blood vessels, can produce serious injury and prevent organ revascularisation (Pegg, 2010). A number of different cryopreservation approaches for WOCP have been evaluated in sheep by different workers with varying degrees of success, with a large proportion of autotransplants failing after a relatively short period of time mainly due to vascular issues, although a single live birth was achieved from lambs, albeit nearly a year and half after autotransplantation (Imhof et al., 2006).

Research conducted in our laboratories has shown that cryopreservation and thawing of whole ovine ovaries using a slightly modified version of the protocol developed for cortical autografts does not significantly affect follicle viability assessed by in vitro measures (Onions et al., 2008) although some evidence of vascular damage within the ovarian pedicle was observed. Despite this, we went onto show that following slow-freezing, rapid-thawing and heterotopic autotransplantation, the reanastomosed ovarian artery of 7 out of 8 whole ovine ovaries was fully patent 7 months after autotransplantation (Onions et al., 2009). Despite patent ovarian vascular perfusion, however, follicle losses were prohibitively high and evidence of restored ovarian function was only seen in 2 out of 8 ewes. We therefore hypothesised that this follicle loss was either due to cryodamage that acute post-thaw in vitro measures of follicle viability were unable to detect, or that it was due to transient ischaemia in the ovary due to cryodamage to the ovarian vasculature. We therefore set out to both optimise the penetration of CPA through the ovary to maximise follicle survival post thaw and to maintain the integrity of the ovarian vasculature following autotransplantation. Utilising array technology we went on to show that ovarian perfusion, CPA exposure and freeze-thaw all induce apoptotic pathways and alter expression of genes associated with endothelial cell function and this finding was consistent with the observation of blood clots within the ovarian artery 7 days after WOCP&TP (Onions et al., 2013). We therefore hypothesised that deleterious effects to the ovarian follicle
population and surrounding ovarian tissue associated with WOCP&TP could be attributed, at least in part, to post-operative ischaemia (within 7 days) resulting from the induction of clotting cascades within the ovarian vasculature caused by either cryo- or cytotoxic damage to the arterial endothelial cells (Onions et al., 2013). Further *in vitro* studies demonstrated that the period of CPA infusion was linearly related to the infusion time and that shorter periods of infusion were associated with lower levels of lactate dehydrogenase (LDH) release *in vitro*, used as an index of cellular damage. We therefore designed an experiment which examined the effect of cryoprotectant (CPA) perfusion time (10 vs 60 min) and pre- and post-operative treatment with the anti-coagulants aspirin in combination with enoxaparine (Clexane®) or eptifibatide (Integrilin®) on ovarian function and fertility 11-23 months after WOCP&TP. The results of this study were far beyond our expectations as combination of aspirin with additional post-operative anticoagulants resulted in acute restoration of ovarian function in 14/14 ewes within 3 weeks.
of WOCP&TP of which 9/14 became pregnant and 4/14 gave birth to a total of 7 normal lambs (Campbell et al., 2014). There was no difference between anti-coagulants in terms of restoration of reproductive function and fertility. In contrast, the duration of CPA perfusion was highly significant with a 60 min perfusion resulting in an ovary of normal appearance and function with high rates of primordial follicle survival (70%) and an abundant blood supply whereas ovaries perfused for 10 min had either resorbed completely and were vestigial (7/14) or were markedly smaller (P < 0.01). It was concluded that both the degree of CPA penetration and maintenance of post-operative vascular patency are critical determinants of the success of WOCP&TP and it is hoped that this work represents a major advance toward the possible development and clinical application of WOCP&TP as a means to preserve and restore the fertility of girls and women at risk of premature ovarian failure. However, before WOCP&TP can be used clinically, further animal model studies are required to; (i) develop optimised cryopreservation protocols for a range of different sized ovaries similar to those observed in prepubertal girls, adolescents and mature women; (ii) generate more information on the optimal type, dose and duration of post-operative anti-thrombotic therapy and (iii) examine the genetic and epigenetic normality of live offspring derived from WOCP&TP.

Concluding comments

In the preceding review we have sought to relate some recent work in our laboratories in order to illustrate the value of integration of both in vitro cell culture and analytical techniques with whole animal studies to ensure significant progress in our understanding of the mechanisms regulating follicular development and the use of ruminants as translational models for the human. One of the major strengths of Prof Eric Lamming was his ability to understand and interact with industry bodies in order to conduct research which was targeted at real production problems and if whole animal ruminant research is to continue to thrive we need to focus on the translational nature of this work to both animal production industries and clinical medicine. Over the last 10-15 years the cost of ruminant research in the UK has increased 20-fold and similar trends seem evident in other European and North American countries. It is therefore becoming increasingly difficult to obtain funding for these types of studies and while a small proportion of studies will continue to be funded the time is perhaps right to look to the types of international collaborations encouraged by bodies such as this Ruminant Reproduction meeting that Eric Lamming helped to establish, in order to maintain the balance and integration of high quality laboratory research with whole animal studies. Despite the costs of these whole animal approaches, the need for increasing ruminant production while minimising environmental impact has never been greater because of global food security and climate change challenges. Therefore the integration of human and animal research has a double benefit for the health and wealth implications of both developed and developing countries.

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